

2024 Hospital Pay-for-Performance Program (for peer groups 1 through 4)

Hospital CQI Performance Index Scorecards

Hospital CQI Pay-for-Performance Program

Hospitals can earn up to 40% of their P4P points based on performance across Blue Cross-supported CQIs.

The CQI component of the P4P is weighted equally for all hospitals, regardless of the number of CQIs a hospital participates in. Therefore, hospitals participating in fewer CQIs will have a greater portion of their incentive allocated to each initiative, while hospitals participating in a greater number of CQIs will have a smaller portion allocated to each initiative. Hospitals participating in more than 10 CQIs will be scored using only the top 10 individual CQI performance scores.

The following chart provides the weight per CQI based on the number of initiatives a hospital participates in:

Number of CQIs	Overall potential incentive	Potential incentive per CQI
1	40%	40%
2	40%	20%
3	40%	13.33%
4	40%	10%
5	40%	8%
6	40%	6.67%
7	40%	5.71%
8	40%	5%
9	40%	4.44%
10+	40%	4%

CQI performance index scorecards

The CQI performance index scorecards will be made available as a separate addendum to the 2024 Pay-for-Performance program guide in mid- to late-December 2023. In addition, each CQI performance index scorecard will be made available through each coordinating center.

All performance index measures and weights are established by the CQI coordinating centers. The weights and measures of a specific CQI index may be adjusted for newly participating hospitals. The coordinating center for each CQI will evaluate and score each hospital's performance index and submit the final aggregate score to Blue Cross.

The measurement period for each performance index measure is January through December, unless otherwise noted in the scorecard.

Specific questions and comments pertaining to the performance index measures should be directed to the respective CQI coordinating center (refer to the following CQI coordinating center contact table).

CQI Program Manager Contacts

CQI	CQI Clinical Focus Area	Index Scorecard Section (click on page # to go to section)	Coordinating Center Program Manager	Email
ASPIRE	Anesthesiology	3-4	Tory Lacca Kate Buehler	lacca@umich.edu kjbucrek@med.umich.edu
BMC2	Angioplasty & Vascular Surgery	5-17	Annemarie Forrest	avassalo@med.umich.edu
HMS	Hospital Medicine	18-27	Elizabeth McLaughlin	emcnair@umich.edu
MAQI2	Anticoagulation	28-33	Brian Haymart	khaymart@umich.edu
MARCQI	Knee/Hip Arthroplasty	34-63	Tae Kim	taekk@med.umich.edu
MBSC	Bariatric Surgery	64-66	Rachel Ross	rachacoo@umich.edu
MEDIC	Emergency Department	67-73	Andy Scott	afscott@med.umich.edu
MROQC	Radiation Oncology	74-76	Melissa Mietzel	hillmel@umich.edu
MSQC	General Surgery	77-89	Amanda Stricklen	aoreilly@umich.edu
MSSIC	Spine Surgery	90-105	Jamie Myers	Jmyer8@hfhs.org
MSTCVS	Cardiac Surgery	106-107	Patty Theurer	ptheurer@umich.edu
MTQIP	Trauma Surgery	108-112	Judy Mikhail	jmikhail@umich.edu
OBI	Obstetrics	113-115	Helen Costis	hcostis@umich.edu

2024 Anesthesiology Quality Improvement and Reporting Exchange (ASPIRE) Performance Index Scorecard Cohorts 1 – 7 Measurement Period: 01/01/2024 - 12/31/2024

		Measurement Period: 01/01/2024 - 12/31/2024	
Measure #	Weight	Measure Description	Points
1	10%	Collaborative Meeting Participation: ASPIRE Quality Champion and Anesthesiology Clinical Quality Reviewer (ACQR) combined attendance at meetings. Three total meetings with six opportunities for attendance.	
		6 / 6 Meetings	10
		5 / 6 Meetings	5
		4 or Less Meetings	0
2	5%	Attend ASPIRE Quality Committee e-meetings: ASPIRE Quality Champion or ACQR attendance across six meetings	
_	070	5 - 6 / 6 Meetings	5
		4 or less Meetings	0
3	5%	ACQR/ASPIRE Quality Champion perform data validation, case validation and submit data by the 3rd Wednesday of each month for January - November and by the 2nd Wednesday of the month for December. Data must be of high quality upon submission, >90% of diagnostics marked as 'Data Accurately Represented.'	
		10 - 12/12 Months	5
		9 or Less Months	0
4	10%	Site Based Quality Meetings: Sites to hold an onsite in- person or virtual meeting following the three ASPIRE Collaborative meetings to discuss the data and plans for quality improvement at their site	
7	10 70	3 Meetings	10
		2 Meetings	5
		1 Meeting	0
5	25%	Sustainability (SUS 02) Percentage of cases where carbon dioxide equivalents (CO2 eq) normalized by hour for cases receiving halogenated agents and/or nitrous oxide is less than CO2 eq of 2% sevoflurane at 2L FGF = 2.83 kg CO2/hr during the maintenance period of anesthesia OR the Total CO2e is less than 2.83 kg CO2. (cumulative score January 1, 2024 - December 31, 2024)	
		Performance is ≥ 45% or show improvement of 10 percentage points	25
		Performance is ≥ 40%	15
		Performance is ≥ 35%	10
		Performance is < 35%	0

2024 Anesthesiology Quality Improvement and Reporting Exchange (ASPIRE) Performance Index Scorecard Cohorts 1 – 7 Measurement Period: 01/01/2024 - 12/31/2024

		Wedsurement Clod. 01/01/2024 - 12/01/2024	
Measure#	Weight	Measure Description	Points
6	20%	Postoperative Nausea and Vomiting (PONV 05) Percentage of patients who had a procedure requiring general anesthesia or cesarean delivery and administered appropriate prophylaxis for PONV. (cumulative score January 1, 2024 - December 31, 2024)	
	2070	Performance is ≥ 70% or improvement of 15 percentage points	20
		Performance is ≥ 65%	15
		Performance is ≥ 60%	10
		Performance is < 60%	0
		Site Directed Measure: Sites choose a measure they are performing above/below ASPIRE threshold or needs improvement by December 8, 2023 (cumulative score January 1, 2024 through December 31, 2024)	
_		Performance is ≥90%; ≤10%; ≤5% or show ≥15% improvement (absolute)	25
7	25%	Performance is ≥85%; ≤15%; ≤10% or show ≥10% improvement (absolute)	15
		Performance is ≥80%; ≤20%; ≤15% or show ≥5% improvement (absolute)	10
		Performance is <80%; >20%; >15% or show <5% improvement (absolute)	0

2024 Blue Cross Blue Shield of Michigan Cardiovascular Consortium (BMC2) Performance Index Scorecard PCI & Vascular Surgery Sites Measure identified in each measure

N4	\\/a:= a4	Magazina Dagarintian	PCI	VS
Measure #	Weight	Measure Description	points	points
		Meeting Participation - Clinician Lead Measurement Period: 01/01/2024 – 12/31/2024		
1	10	Meetings (attendance at the collaborative-wide meeting earns 1 additional extra credit point)	5	5
		1 Meeting	2.5	2.5
		Did not participate	0	0
		Data Coordinator Expectations Measurement Period: 01/01/2024 – 12/31/2024		
2	10	Meets all expectations	5	5
		Meets most expectations	2.5	2.5
		Does not meet expectations	0	0
3	10	Physicians Complete Cross Site Review of Assigned Cases for Procedural Indications and Technical Quality Measurement Period: 01/01/2024 – 12/31/2024		
		Submitted reviews for 100% of cases	5	5
		Submitted reviews for <100% of cases	0	0
4	10	Vascular Surgery Performance Goal - Documentation of EVAR* imaging performed on the 1-year follow up form Measurement Period: 01/01/2024 – 09/30/2024		
·		≥80%	NA	10
		70% - <80%	NA	5
		<70%	NA	0
5		Vascular Surgery Performance Goal - Duplex ultrasound completed prior to asymptomatic carotid endarterectomy Measurement Period: 01/01/2024 – 09/30/2024		
υ	10	≥90%	NA	10
		80% - <90%	NA	5
		<80%	NA	0
6	10	Vascular Surgery Performance Goal - Vein mapping completed before elective lower extremity open bypass Measurement Period: 01/01/2024 – 09/30/2024		
		≥50%	NA	10
		40% - <50%	NA	5
		<40%	NA	0
	Sites	select two measures for scoring from measures 4, 5	, ნ	

2024 Blue Cross Blue Shield of Michigan Cardiovascular Consortium (BMC2) Performance Index Scorecard PCI & Vascular Surgery Sites Measure identified in each measure

Magazina #	\\/aialat	Magazira Dagarintian	PCI	VS
Measure #	Weight	Measure Description	points	points
7	15	Vascular Surgery Performance Goal - Smokers receive smoking cessation treatment prior to discharge Measurement Period: 01/01/2024 – 09/30/2024		
		≥25%	NA	15
		20% - <25%	NA	10
		<20%	NA	0
		PCI Performance Goal - Use of IVUS/OCT^ for stent optimization Measurement Period: 01/01/2024 – 09/30/2024		
8	10	≥45% in EITHER all cases OR ≥45% in cases involving the left main coronary artery, in-stent restenosis, or stent thrombosis	10	NA
		≥10 percentage points absolute increase in all cases* from Q4 YTD 2023	5	NA
		<10 percentage points absolute increase in all cases from Q4 YTD 2023 PCI Performance Goal - Outcomes and	0	NA
		mortality, risk-adjusted AKI, risk-adjusted major bleeding, guideline medications prescription at discharge (aspirin, statin, P2Y12), and referral to cardiac rehab. Measurement Period: 01/01/2024 – 09/30/2024		
		Risk-adjusted mortality		
		A/P <1	5	NA
		A/P > 1, < 1.5	3	NA
		A/P >1.5	0	NA
9	25	Risk-adjusted acute kidney injury A/P <1		NA
3	20	A/P >1, <1.5	5 3	NA NA
		A/P >1.5	0	NA NA
		Risk-adjusted major bleeding		1471
		A/P <1	5	NA
		A/P >1, <1.5	3	NA
		A/P >1.5	0	NA
		Guideline medications prescription at discharge		
		<u>≥</u> 95%	5	NA
		90% - <95%	3	NA
		<90%	0	NA

2024 Blue Cross Blue Shield of Michigan Cardiovascular Consortium (BMC2) Performance Index Scorecard PCI & Vascular Surgery Sites Measure identified in each measure

Measure#	Weight	Measure Description	PCI	VS points
	J	· ·	points	points
		Referral to cardiac rehabilitation		
		<u>></u> 95%	5	NA
		90% - <95%	3	NA
		<90%	0	NA
		PCI Performance Goal - Cardiac rehabilitation utilization within 90 days after PCI discharge†		
		Site performance ≥40% or absolute increase of ≥5 points from baseline site performance. Baseline 1/1/2022 - 9/30/2022. Scored in 2025.	10	NA
10	0	Site performance ≥37% - <40% or absolute increase of ≥3 points from baseline site performance. Scored in 2025.	5	NA
		Site performance <37% and absolute increase of <3 points from baseline site performance. Scored in 2025.	0	NA
11	n/a	Extra credit: 1 point per approved activity (maximum of 5 points) Examples include: Physician attendance at the collaborative-wide meeting, presentation at a meeting, engagement in a work group/task force, referral of an engaged patient advisor, special initiatives, TBD	1-5	

^{*}EVAR = endovascular aneurysm repair

[^]IVUS = intravascular ultrasound; OCT = optical coherence tomography

^{*}Does not apply to the LM, ISR, IST measure

[†]Cardiac rehabilitation utilization will be measured in 2024, but due to data lag, will be scored in 2025

2024 Blue Cross Blue Shield of Michigan Cardiovascular Consortium (BMC2) Performance Index Scorecard PCI only sites Measurement Period identified in the measure

Measure#	Weight	Measure Description	PCI points
		Meeting Participation - Clinician Lead Measurement Period: 01/01/2024 – 12/31/2024	ронно
1	10	2 Meetings (attendance at the collaborative-wide meeting earns 1 additional extra credit point)	10
		1 Meeting	5
		Did not participate	0
		Data Coordinator Expectations Measurement Period: 01/01/2024 – 12/31/2024	
2	10	Meets all expectations	10
		Meets most expectations	5
		Does not meet expectations	0
3	10	Physicians Complete Cross Site Review of Assigned Cases for Procedural Indications and Technical Quality Measurement Period: 01/01/2024 – 12/31/2024	
		Submitted reviews for 100% of cases	10
		Submitted reviews for <100% of cases	0
		PCI Performance Goal - Use of IVUS/OCT^ for stent optimization Measurement Period: 01/01/2024 – 09/30/2024	
4	10	≥45% in EITHER all cases OR ≥45% in cases involving the left main coronary artery, in-stent restenosis, or stent thrombosis	10
		≥10 percentage points absolute increase in all cases ^x from Q4 YTD 2023	5
		<10 percentage points absolute increase in all cases from Q4 YTD 2023	0
5	00	PCI Performance Goal - Outcomes and Process Composite, inclusive of risk-adjusted mortality, risk- adjusted AKI, risk-adjusted major bleeding, guideline medications prescription at discharge (aspirin, statin, P2Y12), and referral to cardiac rehab. Measurement Period: 01/01/2024 – 09/30/2024 Risk-adjusted mortality	
	60	A/P <1	12
		A/P >1, <1.5	8
		A/P >1.5	0
		Risk-adjusted acute kidney injury	
		A/P <1	12
		A/P >1, <1.5	8

2024 Blue Cross Blue Shield of Michigan Cardiovascular Consortium (BMC2) Performance Index Scorecard PCI only sites

Measurement Period identified in the measure

Measure#	Weight	Measure Description	PCI points
		A/P >1.5	0
		Risk-adjusted major bleeding	
		A/P <1	12
		A/P >1, <1.5	8
		A/P >1.5	0
		Guideline medications prescription at discharge	
		<u>></u> 95%	12
Part of		90% - <95%	8
Measure 5		<90%	0
		Referral to cardiac rehabilitation	
		<u>></u> 95%	12
		90% - <95%	8
		<90%	0
	0	PCI Performance Goal - Cardiac rehabilitation utilization within 90 days after PCI discharge† Measurement Period: 01/01/2024 – 09/30/2024	
6		Site performance >40% or absolute increase of >5 points from baseline site performance. Baseline 1/1/2022 - 9/30/2022. Scored in 2025.	10
		Site performance ≥37% - <40% or absolute increase of ≥3 points from baseline site performance. Scored in 2025.	5
		Site performance <37% and absolute increase of <3 points from baseline site performance. Scored in 2025.	0
7	0	Extra credit: 1 point per approved activity (maximum of 5 points) Examples include: Physician attendance at the collaborative-wide meeting, presentation at a meeting, engagement in a work group/task force, referral of an engaged patient advisor, special initiatives, TBD	1-5

[^]IVUS = intravascular ultrasound; OCT = optical coherence tomography

^{*}Does not apply to the LM, ISR, IST measure

[†]Cardiac rehabilitation utilization will be measured in 2024, but due to data lag, will be scored in 2025

2024 Blue Cross Blue Shield of Michigan Cardiovascular Consortium (BMC2) Performance Index Scorecard Vascular Surgery only sites Measurement Period identified in each measure

Measure#	Weight	Measure Description	VS	
ivicasule #	vveigni	·	points	
		Meeting Participation - Clinician Lead Measurement Period: 01/01/2024 - 12/31/2024		
1	10	2 Meetings (attendance at the collaborative-wide meeting earns 1 additional extra credit point)	10	
		1 Meeting	5	
		Did not participate	0	
		Data Coordinator Expectations Measurement Period: 01/01/2024 – 12/31/2024		
2	10	Meets all expectations	10	
		Meets most expectations	7.5	
		Does not meet expectations	0	
3	10	Physicians Complete Cross Site Review of Assigned Cases for Procedural Indications and Technical Quality Measurement Period: 01/01/2024 – 12/31/2024		
		Submitted reviews for 100% of cases	10	
		Submitted reviews for <100% of cases	0	
4	25	Vascular Surgery Performance Goal - Documentation of EVAR* imaging performed on the 1-year follow up form Measurement Period: 01/01/2024 – 09/30/2024		
	20	≥80%	25	
			70% - <80%	15
		<70%	0	
5	25	Vascular Surgery Performance Goal - Duplex ultrasound completed prior to asymptomatic carotid endarterectomy Measurement Period: 01/01/2024 – 09/30/2024		
	20	≥90%	25	
		80% - <90%	15	
		<80%	0	
		Vascular Surgery Performance Goal - Vein mapping completed before elective lower extremity open bypass Measurement Period: 01/01/2024 – 09/30/2024		
6	25	≥50%	25	
		40% - <50%	15	
		<40%	0	
	Sites	select two measures for scoring from measures 4, 5, 6		
7	20	Vascular Surgery Performance Goal - Smokers receive smoking cessation treatment prior to discharge		

2024 Blue Cross Blue Shield of Michigan Cardiovascular Consortium (BMC2) Performance Index Scorecard Vascular Surgery only sites Measurement Period identified in each measure

Measure#	Weight	Measure Description	VS points
		Measurement Period: 01/01/2024 – 09/30/2024	
	≥25		20
		20% - <25%	15
		<20%	0
11	0	Extra credit: 1 point per approved activity (maximum of 5 points) Examples include: Physician attendance at the collaborative-wide meeting, presentation at a meeting, engagement in a work group/task force, referral of an engaged patient advisor, special initiatives, TBD	1-5

^{*}EVAR = endovascular aneurysm repair

2024 Blue Cross Blue Shield of Michigan Cardiovascular Consortium (BMC2) Performance Index Scorecard PCI & YEAR 2 Vascular Surgery Sites Measure identified in each measure

Measure#	Weight	Measure Description	PCI points	VS points
1		Meeting Participation - Clinician Lead Measurement Period: 01/01/2024 – 12/31/2024	рошко	pomie
	15	2 Meetings (attendance at the collaborative-wide meeting earns 1 additional extra credit point)	5	10
		1 Meeting	2.5	5
		Did not participate	0	0
		Data Coordinator Expectations Measurement Period: 01/01/2024 – 12/31/2024		
2	20	Meets all expectations	5	15
		Meets most expectations	2.5	10
		Does not meet expectations	0	0
3	15	Physicians Complete Cross Site Review of Assigned Cases for Procedural Indications and Technical Quality Measurement Period: 01/01/2024 – 12/31/2024		
		Submitted reviews for 100% of cases	5	10
		Submitted reviews for <100% of cases	0	0
4	5	Vascular Surgery Performance Goal - Documentation of EVAR* imaging performed on the 1-year follow up form Measurement Period: 01/01/2024 – 09/30/2024		
		≥80%	NA	5
		70% - <80%	NA	2.5
		<70%	NA	0
5	5	Vascular Surgery Performance Goal - Duplex ultrasound completed prior to asymptomatic carotid endarterectomy Measurement Period: 01/01/2024 – 09/30/2024		
Ü		≥90%	NA	5
		80% - <90%	NA	2.5
		<80%	NA	0
6	5	Vascular Surgery Performance Goal - Vein mapping completed before elective lower extremity open bypass Measurement Period: 01/01/2024 – 09/30/2024		
		≥50%	NA	5
		40% - <50%	NA	2.5
		<40%	NA	0
	Sites	select two measures for scoring from measures 4, 5	, 6	
7	5	Vascular Surgery Performance Goal - Smokers receive smoking cessation treatment prior to discharge		

2024 Blue Cross Blue Shield of Michigan Cardiovascular Consortium (BMC2) Performance Index Scorecard PCI & YEAR 2 Vascular Surgery Sites Measure identified in each measure

Massamall	\A/a:ada4	Manage Description	PCI	VS
Measure #	Weight	Measure Description	points	points
		Measurement Period: 01/01/2024 – 09/30/2024		
		≥25%	NA	5
		20% - <25%	NA	2.5
		<20%	NA	0
		PCI Performance Goal - Use of IVUS/OCT^ for stent optimization Measurement Period: 01/01/2024 – 09/30/2024		
8	10	≥45% in EITHER all cases OR ≥45% in cases involving the left main coronary artery, in-stent restenosis, or stent thrombosis	10	NA
		≥10 percentage points absolute increase in all cases* from Q4 YTD 2023	5 NA	NA
		<10 percentage points absolute increase in all cases from Q4 YTD 2023	0	NA
		mortality, risk-adjusted AKI, risk-adjusted major bleeding, guideline medications prescription at discharge (aspirin, statin, P2Y12), and referral to cardiac rehab. Measurement Period: 01/01/2024 – 09/30/2024		
		Risk-adjusted mortality A/P <1	5	NΙΛ
		A/P >1, <1.5	3	5 2.5 0 NA NA
		A/P > 1, < 1.5 A/P > 1.5	0	
		Risk-adjusted acute kidney injury		INA
•	0.5	A/P <1	5	NA
9	25	A/P >1, <1.5	3	
		A/P >1.5	0	
		Risk-adjusted major bleeding		
		A/P <1	5	NA
		A/P >1, <1.5	3	NA
		A/P >1.5	0	NA
		Guideline medications prescription at discharge		
	1	>95%	5	NA
		90% - <95%	3	
		-	3 0	NA

2024 Blue Cross Blue Shield of Michigan Cardiovascular Consortium (BMC2) Performance Index Scorecard PCI & YEAR 2 Vascular Surgery Sites Measure identified in each measure

Measure#	Weight	Maggura Description	PCI	VS
ivicasure #	vveignt	Measure Description	points	points
		<u>≥</u> 95%	5	NA
		90% - <95%	3	NA
		<90%	0	NA
		PCI Performance Goal - Cardiac rehabilitation utilization within 90 days after PCI discharge† Measurement Period: 01/01/2024 – 09/30/2024		
10	0	Site performance ≥40% or absolute increase of ≥5 points from baseline site performance. Baseline 1/1/2022 - 9/30/2022. Scored in 2025.	10	NA
10	0	Site performance >37% - <40% or absolute increase of >3 points from baseline site performance. Scored in 2025.	5	NA
		Site performance <37% and absolute increase of <3 points from baseline site performance. Scored in 2025.	0	NA
11	0	Extra credit: 1 point per approved activity (maximum of 5 points) Examples include: Physician attendance at the collaborative-wide meeting, presentation at a meeting, engagement in a work group/task force, referral of an engaged patient advisor, special initiatives, TBD	1-5	

^{*}EVAR = endovascular aneurysm repair

[^]IVUS = intravascular ultrasound; OCT = optical coherence tomography

^{*}Does not apply to the LM, ISR, IST measure

[†]Cardiac rehabilitation utilization will be measured in 2024, but due to data lag, will be scored in 2025

2024 Blue Cross Blue Shield of Michigan Cardiovascular Consortium (BMC2) Performance Index Scorecard VS only sites in Year 2 of consortium participation Measure identified in each measure

Measure#	Weight	Measure Description	VS	
Weasure #	vveignt	·	points	
		Meeting Participation - Clinician Lead Measurement Period: 01/01/2024 – 12/31/2024		
1	25	2 Meetings (attendance at the collaborative-wide meeting earns 1 additional extra credit point)	25	
		1 Meeting	15	
		Did not participate	0	
		Data Coordinator Expectations Measurement Period: 01/01/2024 – 12/31/2024	15 0 25 15 0 20 0	
2	25	Meets all expectations	25	
		Meets most expectations	15	
		Does not meet expectations	0	
3	20	Physicians Complete Cross Site Review of Assigned Cases for Procedural Indications and Technical Quality Measurement Period: 01/01/2024 – 12/31/2024		
		Submitted reviews for 100% of cases	20	
		Submitted reviews for <100% of cases	0	
4	10	Vascular Surgery Performance Goal - Documentation of EVAR* imaging performed on the 1-year follow up form Measurement Period: 01/01/2024 – 09/30/2024		
		≥80%		
		70% - <80%		
5	10	<70% Vascular Surgery Performance Goal - Duplex ultrasound completed prior to asymptomatic carotid endarterectomy Measurement Period: 01/01/2024 – 09/30/2024		
	10	≥90%	10	
		80% - <90%	5	
		<80%	0	
6	40	Vascular Surgery Performance Goal - Vein mapping completed before elective lower extremity open bypass Measurement Period: 01/01/2024 – 09/30/2024		
	10	≥50%	10	
		40% - <50%	5	
		<40%	0	
	Sites	select two measures for scoring from measures 4, 5, 6		
7	10	Vascular Surgery Performance Goal - Smokers receive smoking cessation treatment prior to discharge Measurement Period: 01/01/2024 – 09/30/2024		

2024 Blue Cross Blue Shield of Michigan Cardiovascular Consortium (BMC2) Performance Index Scorecard VS only sites in Year 2 of consortium participation Measure identified in each measure

Measure#	Weight	Measure Description	VS points
		≥25%	10
		20% - <25%	5
		<20%	0
8	0	Extra credit: 1 point per approved activity (maximum of 5 points) Examples include: Physician attendance at the collaborative-wide meeting, presentation at a meeting, engagement in a work group/task force, referral of an engaged patient advisor, special initiatives, TBD	1-5

^{*}EVAR = endovascular aneurysm repair

2024 Blue Cross Blue Shield of Michigan Cardiovascular Consortium (BMC2) Performance Index Scorecard VS only sites in Year 1 of consortium participation Measurement Period: 01/01/2024 – 12/31/2024

Measure#	Weight	Measure Description	VS Points
		Meeting Participation - Clinician Lead	
		2 Meetings	35
1	35	1 Meeting	20
		Did not participate	0
		Data Coordinator Expectations	
		Meets all expectations	35
2	35	Meets most expectations	20
		Does not meet expectations	0
3	20	Physicians Complete Cross Site Review of Assigned Cases for Procedural Indications and Technical Quality	
	30	Submitted reviews for 100% of cases	30
		Submitted reviews for <100% of cases	0

2024 Michigan Hospital Medicine Safety (HMS) Collaborative Quality Initiative Performance Index Scorecard PICC Insertions/Hospital DischargesHospitals Enrolled Prior to 2020

Hospitals Ellioned Filor to 2020
Measurement Period: 08/01/2024-11/06/2024

Measure#	Weight	Measure Description	Points
		Timeliness of HMS Data at Mid-Year and End of Year ¹	
1	5	On time > 95% at Mid-Year AND End of Year	5
'	5	On time ≥ 95% at Mid-Year OR End of Year	3
		On time < 95% at Mid-Year AND End of Year	0
		Completeness ¹ and Accuracy ^{2,3} of HMS Data	
		≥ 95% of registry data complete & accurate, semi-annual	
2	5	QI activity surveys completed, AND audit case corrections	5
2	3	completed by due date < 95% of registry data complete & accurate, semi-annual	
		QI activity survey not completed OR audit case corrections	0
		not completed by due date	
		Consortium-wide Meeting Participation ⁴ – clinician lead or designee	
	40	3 meetings	1
3	10		0
		2 meetings	
		1 meeting	
		No meetings	0
		Consortium-wide Meeting Participation⁴ – data abstractor, QI staff, or other	
		3 meetings	1
4	10		0
		2 meetings	5
		1 meeting	0
		No meetings	0
		Increase Use of 5 Days of Antibiotic Treatment ⁵ in Uncomplicated CAP (Community Acquired Pneumonia) Cases (i.e., reduce excess durations) ^{6,7}	
		≥ 70% uncomplicated CAP cases receive 5 days⁵ of	
		antibiotics OR	3 0 5 0 1 0 5 0 0
		≥ 70% relative increase in the number of uncomplicated	
		CAP cases that receive 5 days ⁵ of antibiotics during the current performance year ⁸	U
_	40	55-69% uncomplicated CAP cases receive 5 days ⁵ of	
5	10	antibiotics OR	
		50-69% relative increase in the number of uncomplicated	5
		CAP cases that receive 5 days ⁵ of antibiotics during the current performance year ⁸	
		< 55% uncomplicated CAP cases receive 5 days ⁵ of	
		antibiotics AND	
		< 50% relative increase in the number of uncomplicated	0
		CAP cases that receive 5 days of antibiotics during the	
		current performance year ⁸	

2024 Michigan Hospital Medicine Safety (HMS) Collaborative Quality Initiative Performance Index Scorecard PICC Insertions/Hospital DischargesHospitals Enrolled Prior to 2020

		Measurement Period: 08/01/2024-11/06/2024	
Measure#	Weight	Measure Description	Points
		Reduce Use of Antibiotics in Patients with ASB (Asymptomatic Bacteriuria) ⁹ and Questionable Pneumonia ^{6,7,10}	
6	10	≤ 13% of positive urine culture cases treated with an antibiotic are ASB cases ⁹ AND ≤ 11% of pneumonia cases treated with an antibiotic are questionable pneumonia ¹⁰	1 0
	10	≤ 13% of positive urine culture cases treated with an antibiotic are ASB cases ⁹ OR ≤ 11% of pneumonia cases treated with an antibiotic are questionable pneumonia ¹⁰	5
		> 13% of positive urine culture cases treated with an antibiotic are ASB cases ⁹ AND > 11% of pneumonia cases treated with an antibiotic are questionable pneumonia ¹⁰	0
		Increase Antibiotics Delivered within 3 hours of Arrival for Septic Shock Patients ^{11,21}	
7	15	≥ 67% septic shock cases ¹¹ receive antibiotics within 3 hours of arrival	1 5
		55 – 66% septic shock cases ¹¹ receive antibiotics within 3 hours of arrival	1 0
		< 55% septic shock cases ¹¹ receive antibiotics within 3 hours of arrival	0
		Increase Discharge/Post-Discharge Care Coordination for Sepsis Patients Discharged to Home-like Setting ^{12,13,21}	
8	15	≥ 65% sepsis cases discharged to home-like setting 12 received at least 1 of 3 discharge/post- discharge coordination of care measures 13	1 5
	10	45 – 64% sepsis cases discharged to home-like setting ¹² received at least 1 of 3 discharge/post-discharge coordination of care measures ¹³	1 0
		< 45% sepsis cases discharged to home-like setting 12 received at least 1 of 3 discharge/post-discharge coordination of care measures 13	0
		Reduce Inappropriate PICC Placements in Special Populations – Active Malignancy ¹⁴ or Critical Care ¹⁵	
		Active Malignancy ¹⁴	
9	15	≤ 25% of PICCs placed in active malignancy cases ¹⁴ are triple lumens and in for < 5 Days AND participation in special population workgroup ^{16,17,18}	1 5
		≤ 25% of PICCs placed in active malignancy cases ¹⁴ are triple lumens and in for < 5 Days OR participation in special population workgroup ^{16,17,18}	1 0
		> 25% of PICCs placed in active malignancy cases ¹⁴ are triple lumens and in for < 5 Days AND No participation in special population workgroup ^{16,17,18}	0

2024 Michigan Hospital Medicine Safety (HMS) Collaborative Quality Initiative Performance Index Scorecard PICC Insertions/Hospital DischargesHospitals Enrolled Prior to 2020 Measurement Period: 08/01/2024-11/06/2024

Measure#	Weight	Measure Description	Points
		OR	
		Critical Care (ICU) ¹⁵	
		≤ 30% of PICCs placed in critical care cases ¹⁵ are triple	
		lumens AND Participation in special population workgroup ^{16,17,18}	15
		≤ 30% of PICCs placed in critical care cases ¹⁵ are triple	40
		lumens OR Participation in special population workgroup ^{16,17,18}	10
		> 30% of PICCs placed in critical care cases ¹⁵ are triple	0
		lumens AND No Participation in special population workgroup ^{16,17,18}	0
		Reduce Use of Inappropriate Empiric Broad-Spectrum	
	5	Antibiotics ¹⁹ for Patients with Uncomplicated CAP (Community Acquired Pneumonia) ²⁰	
_		<u>< 10% collaborative-wide average</u> ²⁰ of uncomplicated	
С		CAP cases receive an inappropriate broad-spectrum antibiotic empirically ¹⁹	5
		> 10% collaborative-wide average ²⁰ of uncomplicated	
		CAP cases receive an inappropriate broad-spectrum antibiotic empirically ¹⁹	0
		Optional Bonus	
		Specialist ¹⁸ attendance at 3 of the virtual special population	
		workgroup meetings ¹⁷ during the calendar year in the	
Optional	5	hospital's pre-determined workgroup area OR	5
Optional		Specialist ¹⁸ attendance at 1 or more of the special	0
		population workgroup meetings ¹⁷ that is not the hospital's	
		pre-determined workgroup area during the calendar year	
	_	Emergency Medicine Physician attendance at the 2 in-	_
Optional	5	person collaborative wide meetings convened during the calendar year (July & November)	5

HMS Supporting Documentation

Hospitals Enrolled Prior to 2020

- ¹ Registry data for all initiatives (Antimicrobial, PICC/Midline and Sepsis) assessed during mid-year performance evaluation review and at year end based on data submitted during calendar year 2024. All required cases must be completed by the mid-year performance evaluation review AND by year end. Mid-year due date and final due date will be announced by Coordinating Center.
- ² Assessed based on scores received for site audits conducted during performance year 2024. Scores are averaged if multiple audits take place during the year. Both semi-annual QI activity surveys must be completed by due dates announced by Coordinating Center.
- ³ For audits conducted during the performance year, audit case corrections must be completed, or discrepancies addressed within 3 months of audit summary receipt (due date for case corrections provided in audit summary) or end of performance year deadline whichever comes first.
- ⁴ Based on all meetings scheduled during calendar year 2024. Clinician lead or designee must be a physician as outlined in Hospital Expectations.
- ⁵ Duration is considered appropriate if 6 or fewer days of total antibiotic treatment (inpatient and outpatient) is administered.
- ⁶ Assessed at year end based on final quarter of data entered (per the data abstraction calendar) in the data registry during the performance year 2024. To determine the final score, an adjusted statistical model will be utilized. The method for obtaining each hospital's adjusted performance measurement utilizes all available data from the most recent 4 quarters. The collaborative wide average and collaborative wide improvement or decline, as well as the average rate change over time of each individual hospital are incorporated into the final adjusted rate. Each hospital's adjusted rate reflects both change in performance over time and overall performance relative to the collaborative averages. The adjusted performance is a more stable and reliable estimate of each hospital's current performance, their performance relative to collaborative as a whole, and reflects the improvement work each hospital is doing over a given performance year.
- ⁷ The adjusted model for this measure includes all cohorts.
- ⁸ Rate of change is based on the adjusted method between Q1 2024 and Q4 2024 and may not reflect raw rates from quarter to quarter.
- ⁹ Antibiotic treatment for ASB is assessed based on treatment on day 2 or later of the entire hospital encounter. This portion of the measure is assessed out of all positive urine culture cases abstracted during the performance year.
- ¹⁰ Questionable pneumonia is defined as cases abstracted into the pneumonia registry who do not meet the clinical and radiographic criteria to be classified as Community Acquired Pneumonia. Antibiotic treatment for these cases is assessed on day 3 or later of the hospital encounter. This portion of the measure is assessed out of all pneumonia cases abstracted during the performance year.
- ¹¹ Cases with septic shock are defined as those with hypotension (vasopressors initiated within two hours of arrival OR systolic blood pressure < 90 mmHg within two hours of arrival OR calculated MAP < 65 within two hours of arrival). Patients excluded from review in this measure include those with < 2 SIRS, normal WBC, no elevated lactate, and no symptoms of infection.
- ¹² Home-like Setting = home (with or without home services), assisted living, custodial nursing, temporary shelter.
- ¹³ Discharge/post- discharge coordination of care measures:
 - Hospital contact information provided at discharge (in discharge summary)

- Scheduled visit with Primary Care Physician (PCP) within 2 weeks (at time of discharge)
- Post-discharge telephone call or PCP visit/home health services within 3 calendar days of hospital discharge
- ¹⁴ PICC placements where the medical record reflects a cancer diagnosis **AND** the PICC was placed for a cancer-related admission.
- ¹⁵ PICC placements where the patient was in the ICU at the time of PICC insertion.
- ¹⁶ Participation in Special Population Workgroup
 - At least 3 individuals representing the following roles must attend 3 of the 3 tri-annual initiative specific work group meetings:
 - 1 Quality Professional
 - 1 Physician (the physician in attendance for at least 1 of the 3 meetings per year must be a specialist in the special population area)
 - o 1 Vascular Access Team Member or Interventional Radiologist Representative
- ¹⁷ Initiative specific workgroups will take place virtually during the calendar year 2024.
- ¹⁸ Specialist is considered Critical Care, Oncology or Hematology Physician
 - Oncology workgroup area = Oncology or Hematology Physician
 - Critical Care (ICU) workgroup area = Critical Care Physician

If hospital does not have a specialist either employed at the hospital or contracted by the hospital, the HMS Physician Champion is acceptable, however, must be approved by the Coordinating Center.

- ¹⁹ Empiric antibiotic therapy is assessed on day 2 of the hospital encounter for Uncomplicated CAP cases that are eligible for 5 days of antibiotic therapy.
- ²⁰ Assessed at year end based on the collaborative-wide average for the final quarter of data entered (per the data collection calendar) in the data registry during the performance year 2024. This is different than the other performance measures in the index, which are applied to each individual hospital.
- ²¹ Assessed at year end based on the raw rate for the individual site for the final quarter of data entered (per the data collection calendar) in the data registry during the performance year 2024. This is different than the other site-only performance measures, which are based on the adjusted rate for the site.

2024 Michigan Hospital Medicine Safety (HMS) Collaborative Quality Initiative Performance Index Scorecard (PICC Insertions/Hospital Discharges)- Hospitals Enrolled in 2020, 2021, 2022 Measurement Period: 08/01/2024-11/06/2024

Measure#	Weight	Measure Description	Points
		Timeliness of HMS Data at Mid-Year and End of Year ¹	
1	5	On time <u>></u> 95% at Mid-Year AND End of Year	5
'	3	On time ≥ 95% at Mid-Year OR End of Year	3
		On time < 95% at Mid-Year AND End of Year	0
		Completeness ¹ and Accuracy ^{2,3} of HMS Data	
		≥ 95% of registry data complete & accurate, semi-	
2	5	annual QI activity surveys completed, AND audit case	5
2	5	corrections completed by due date	
		< 95% of registry data complete & accurate, semi- annual QI activity survey not completed OR audit case	0
		corrections not completed by due date	U
		Consortium-wide Meeting Participation⁴ – clinician lead	
		or designee	
3	10	3 meetings	10
3	10	2 meetings	5
		1 meeting	0
		No meetings	0
		Consortium-wide Meeting Participation⁴ – data abstractor, QI staff, or other	
4	10	3 meetings	10
4	10	2 meetings	0 10 5 0
		1 meeting	
		No meetings	0
		Increase Use of 5 Days of Antibiotic Treatment ⁵ in Uncomplicated CAP (Community Acquired Pneumonia) Cases (i.e., reduce excess durations) ^{6,7}	
		≥ 70% uncomplicated CAP cases receive 5 days⁵ of	
		antibiotics OR	
		≥ 70% relative increase in the number of uncomplicated	10
		CAP cases that receive 5 days ⁵ of antibiotics during the current performance year ⁸	
		55-69% uncomplicated CAP cases receive 5 days ⁵	
5	10	of antibiotics OR	
		50-69% relative increase in the number of uncomplicated	5
		CAP cases that receive 5 days ⁵ of antibiotics during the	
		current performance year ⁸ < 55% uncomplicated CAP cases receive 5 days ⁵ of	
		antibiotics AND	
		< 50% relative increase in the number of uncomplicated	0
		CAP cases that receive 5 days ⁵ of antibiotics during the	•
		current performance year ⁸	
6	10	Reduce Use of Antibiotics in Patients with ASB (Asymptomatic Bacteriuria) ⁹ and Questionable Pneumonia ^{6,7,10}	

2024 Michigan Hospital Medicine Safety (HMS) Collaborative Quality Initiative Performance Index Scorecard (PICC Insertions/Hospital Discharges)- Hospitals Enrolled in 2020, 2021, 2022 Measurement Period: 08/01/2024-11/06/2024

Measure#	Weight	Measure Description	Points
		≤ 13% of positive urine culture cases treated with an antibiotic are ASB cases ⁹ AND ≤ 11% of pneumonia cases treated with an antibiotic are questionable pneumonia ¹⁰	10
		≤ 13% of positive urine culture cases treated with an antibiotic are ASB cases OR < 11% of pneumonia cases treated with an antibiotic are questionable pneumonia 10	5
		> 13% of positive urine culture cases treated with an antibiotic are ASB cases AND > 11% of pneumonia cases treated with an antibiotic are questionable pneumonia 10	0
		Increase Antibiotics Delivered within 3 hours of Arrival for Septic Shock Patients ^{11,19}	
7	15	≥ 67% septic shock cases ¹¹ receive antibiotics within 3 hours of arrival	15
		55 – 66% septic shock cases ¹¹ receive antibiotics within 3 hours of arrival	10
		< 55% septic shock cases ¹¹ receive antibiotics within 3 hours of arrival	0
		Increase Discharge/Post-Discharge Care Coordination for Sepsis Patients Discharged to Home-like Setting ^{12,13,19}	
0	45	≥ 65% sepsis cases discharged to home-like setting ¹² received at least 1 of 3 discharge/post- discharge coordination of care measures ¹³	15
8	15	45 – 64% sepsis cases discharged to home-like setting ¹² received at least 1 of 3 discharge/post-discharge coordination of care measures ¹³	10
		< 45% sepsis cases discharged to home-like setting ¹² received at least 1 of 3 discharge/post-discharge coordination of care measures ¹³	5 0 15 10 0
		Reduce PICCs (Peripherally-Inserted Central Catheters) in for ≤ 5 Days (excluding deaths) ^{6,14}	
9	5	≤ 10% of PICC cases in for ≤ 5 Days	0 15 10 0 15 10 0 5 3 0
		1115% of PICC cases in for ≤ 5 Days	3
		> 15% of PICC cases in for ≤ 5 Days	0
		Increase Use of Single Lumen PICCs in Non-ICU (Intensive Care Unit) Cases ^{6,14}	
10	5	≥ 85% of non-ICU PICC cases have a single lumen	
		70-84% of non-ICU PICC cases have a single lumen	
		< 70% of non-ICU PICC cases have a single lumen	0
11	5	PICCs in Patients with eGFR (estimated glomerular filtration rate) <45 (without Nephrology approval) ^{6,14}	

2024 Michigan Hospital Medicine Safety (HMS) Collaborative Quality Initiative Performance Index Scorecard (PICC Insertions/Hospital Discharges)- Hospitals Enrolled in 2020, 2021, 2022 Measurement Period: 08/01/2024-11/06/2024

Measure #	Weight	Measure Description	Points
		≤ 5% of cases with PICC have eGFR <45 without	5
		Nephrology approval	3
		6 -12% of cases with PICC have eGFR <45 without	3
		Nephrology approval	3
		>12% of cases with PICC have eGFR <45 without	0
		Nephrology approval	0
		Reduce Use of Inappropriate Empiric Broad-Spectrum	
		Antibiotics ¹⁵ for Patients with Uncomplicated CAP	
		(Community Acquired Pneumonia) ¹⁶	
		< 10% collaborative-wide average 16 of uncomplicated	
С	5	CAP cases receive an inappropriate broad-spectrum	5
		antibiotic empirically ¹⁵	
		> 10% collaborative-wide average 16 of uncomplicated	
		CAP cases receive an inappropriate broad-spectrum	0
		antibiotic empirically ¹⁵	
		Optional Bonus	
0 "	_	Specialist ¹⁷ attendance at 1 or more of the special	_
Optional	5	population workgroup meetings ¹⁸ during the calendar year	5
		Emergency Medicine Physician attendance at the 2 in-	
Optional	5	person collaborative wide meetings convened during the	5
Ориона	3	calendar year (July & November)	J
		Calendar year (Sury & November)	

HMS Supporting Documentation

Hospitals Enrolled in 2020, 2021, 2022

- ¹ Registry data for all initiatives (Antimicrobial, PICC/Midline and Sepsis) assessed during mid-year performance evaluation review and at year end based on data submitted during calendar year 2024. All required cases must be completed by the mid-year performance evaluation review AND by year end. Mid-year due date and final due date will be announced by Coordinating Center.
- ² Assessed based on scores received for site audits conducted during performance year 2024. Scores are averaged if multiple audits take place during the year. Both semi-annual QI activity surveys must be completed by due dates announced by Coordinating Center.
- ³ For audits conducted during the performance year, audit case corrections must be completed, or discrepancies addressed within 3 months of audit summary receipt (due date for case corrections provided in audit summary) or end of performance year deadline whichever comes first.
- ⁴ Based on all meetings scheduled during calendar year 2024. Clinician lead or designee must be a physician as outlined in Hospital Expectations.
- ⁵ Duration is considered appropriate if 6 or fewer days of total antibiotic treatment (inpatient and outpatient) is administered.
- ⁶ Assessed at year end based on final quarter of data entered (per the data abstraction calendar) in the data registry during the performance year 2024. To determine the final score, an adjusted statistical model will be utilized. The method for obtaining each hospital's adjusted performance measurement utilizes all available data from the most recent 4 quarters. The collaborative wide average and collaborative wide improvement or decline, as well as the average rate change over time of each individual hospital are incorporated into the final adjusted rate. Each hospital's adjusted rate reflects both change in performance over time and overall performance relative to the collaborative averages. The adjusted performance is a more stable and reliable estimate of each hospital's current performance, their performance relative to collaborative as a whole, and reflects the improvement work each hospital is doing over a given performance year.
- ⁷ The adjusted model for this measure includes all cohorts.
- ⁸ Rate of change is based on the adjusted method between Q1 2024 and Q4 2024 and may not reflect raw rates from quarter to quarter.
- ⁹ Antibiotic treatment for ASB is assessed based on treatment on day 2 or later of the entire hospital encounter. This portion of the measure is assessed out of all positive urine culture cases abstracted during the performance year.
- ¹⁰ Questionable pneumonia is defined as cases abstracted into the pneumonia registry who do not meet the clinical and radiographic criteria to be classified as Community Acquired Pneumonia. Antibiotic treatment for these cases is assessed on day 3 or later of the hospital encounter. This portion of the measure is assessed out of all pneumonia cases abstracted during the performance year.
- ¹¹ Cases with septic shock are defined as those with hypotension (vasopressors initiated within two hours of arrival OR systolic blood pressure < 90 mmHg within two hours of arrival OR calculated MAP < 65 within two hours of arrival). Patients excluded from review in this measure include those with < 2 SIRS, normal WBC, no elevated lactate, and no symptoms of infection.
- ¹² Home-like Setting = home (with or without home services), assisted living, custodial nursing, temporary shelter.
- ¹³ Discharge/post- discharge coordination of care measures:
 - Hospital contact information provided at discharge (in discharge summary)

- Scheduled visit with Primary Care Physician (PCP) within 2 weeks (at time of discharge)
- Post-discharge telephone call or PCP visit/home health services within 3 calendar days of hospital discharge
- ¹⁴ The adjusted model for this measure includes cohorts 2020, 2021 and 2022 only.
- ¹⁵ Empiric antibiotic therapy is assessed on day 2 of the hospital encounter for Uncomplicated CAP cases that are eligible for 5 days of antibiotic therapy.
- ¹⁶ Assessed at year end based on the collaborative-wide average for the final quarter of data entered (per the data collection calendar) in the data registry during the performance year 2024. This is different than the other performance measures in the index, which are applied to each individual hospital.
- ¹⁷ Specialist is considered Critical Care, Oncology or Hematology Physician
 - Oncology workgroup area = Oncology or Hematology Physician
 - Critical Care (ICU) workgroup area = Critical Care Physician

If hospital does not have a specialist either employed at the hospital or contracted by the hospital, the HMS Physician Champion is acceptable, however, must be approved by the Coordinating Center.

- ¹⁸ Initiative specific workgroups will take place virtually during the calendar year 2024.
- ¹⁹ Assessed at year end based on the raw rate for the individual site for the final quarter of data entered (per the data collection calendar) in the data registry during the performance year 2024. This is different than the other site-only performance measures, which are based on the adjusted rate for the site.

2024 Michigan Anticoagulation Improvement Initiative (MAQI2) Collaborative Quality Initiative Performance Index Scorecard Ascension St. John and Trinity Ann Arbor - Year 2 Measurement period: 01/01/2024 – 12/31/2024

		Measurement period: 01/01/2024 – 12/31/2024	
Measure #	Weight	Measure Description	Points
		Implementation of DOAC dashboard* Ascension St. John and Trinity Ann Arbor Year 2 1/1-12/31	
		Fully implemented DOAC dashboard	10
1	10	Site IT staff fully engaged and programming process underway	8
'	10	Organizational approval received and project added to IT project list	6
		Site team making demonstrable effort to get organizational approval	4
		No site engagement regarding DOAC dashboard implementation	0
		Inappropriate aspirin in patients on DOACs*	
2	10	≤5% or a relative reduction of ≥10%	10
		6-10% or a relative reduction of 6-9%	5
		11-15% or a relative reduction of 2-5%	3
		>15% or a relative reduction of <2%	0
	10	Collaborative-wide meeting participation – Clinical Champion	
3		attended all 3 meetings	10
		attended 2 out of 3 meetings	7
		attended 1 out of 3 meetings	3
		did not attend any meetings	0
	10	Collaborative-wide meeting participation – Coordinator/Lead Abstractor	
4		attended all 3 meetings	10
		attended 2 out of 3 meetings	7
		attended 1 out of 3 meetings	3
		did not attend any meetings	0
5	30	Completeness and Accuracy of data	
		Critical data elements are complete/accurate in ≥90% of cases	10
		Critical data elements are complete/accurate in 70-89% of cases	5
		Critical data elements are complete/accurate in <70% of cases	0

2024 Michigan Anticoagulation Improvement Initiative (MAQI2) Collaborative Quality Initiative Performance Index Scorecard Ascension St. John and Trinity Ann Arbor - Year 2 Measurement period: 01/01/2024 – 12/31/2024

Measure#	Weight	Measure Description	Points
		Volume of data abstraction	
		≥ 90% of expected volume entered by site (per 0.5 FTE: enter 500 new pts or maintain follow-ups in 300 active registry patients)	30
		70-89% of expected volume entered by site (per 0.5 FTE: enter 500 new pts or maintain follow-ups in 300 active registry patients)	20
6	30	50-69% of expected volume entered by site (per 0.5 FTE: enter 500 new pts or maintain follow-ups in 300 active registry patients)	10
		<50% of expected volume entered by site (per 0.5 FTE: enter 500 new pts or maintain follow-ups in 300 active registry patients)	0

^{*}Sites can chose to do one or both of these two initiatives. If one is chosen, it will be worth 20 maximum points.

2024 Michigan Anticoagulation Improvement Initiative (MAQI2) Collaborative Quality Initiative Performance Index Scorecard Epic Sites Measurement period: 01/01/2024 – 12/31/2024

Measure#	Weight	Measure Description	Points
1	10	Smoking status assessment in newly enrolled patients	
		≥90% of newly enrolled patients in 2022 will have smoking status assessed and documented per site protocol	10
		70-89% of newly enrolled patients in 2022 will have smoking status assessed and documented per site protocol	5
		50-69% of newly enrolled patients in 2022 will have smoking status assessed and documented per site protocol	3
		<50% of newly enrolled patients in 2022 will have smoking status assessed and documented per site protocol	0
		DOAC Dashboard Utilization	
		≥750 known/possible critical alerts addressed^ in 2024	10
2	10	500-749 known/possible critical alerts addressed in 2024	5
		250-499 known/possible critical alerts addressed in 2024	3
		<250 known/possible critical alerts addressed in 2024	0
		Inappropriate aspirin use in warfarin patients (updated 2024 criteria)	
3	20	≤12% or relative decrease of ≥15%	20
3	20	13-19% or relative decrease of 10-14%	10
		20-24% or relative decrease of 5-9%	5
		>24% or relative decrease of <5%	0
		Extended International Normalized Ratio (INR) testing interval	
4	10	≥80% of eligible patients received extended intervals	10
4		60-79% of eligible patients received extended intervals	5
		40-59% of eligible patients received extended intervals	3
		<40% of eligible patients received extended intervals	0
		Gastroprotection* in warfarin patients at high-risk for upper GI bleeding (site level)	
5	10	≥55% or relative increase of ≥10%	10
5		40-54% or relative increase of 6-9%	5
		25-39% or relative increase of 2-5%	3
		<25% or relative increase of <2%	0
	10	Gastroprotection* in warfarin patients at high-risk for upper GI bleeding (consortium level)	
6		≥55% or relative increase of ≥10%	10
6		40-54% or relative increase of 6-9%	5
		25-39% or relative increase of 2-5%	3
		<25% or relative increase of <2%	0

2024 Michigan Anticoagulation Improvement Initiative (MAQI2) Collaborative Quality Initiative Performance Index Scorecard **Epic Sites**

Measurement period: 01/01/2024 - 12/31/2024

Measure#	Weight	Measure Description	Points
		Collaborative-wide meeting participation -Clinical Champion	
_		attended all 3 meetings	10
7	10	attended 2 out of 3 meetings	7
		attended 1 out of 3 meetings	3
		did not attend any meetings	0
	10	Collaborative-wide meeting participation – Coordinator/Lead Abstractor	
•		attended all 3 meetings	10
8		attended 2 out of 3 meetings	7
		attended 1 out of 3 meetings	4
		did not attend any meetings	0
	10	Completeness and Accuracy of data	
		Critical data elements are complete/accurate in ≥90% of cases	10
9		Critical data elements are complete/accurate in 70-89% of cases	5
		Critical data elements are complete/accurate in <70% of cases	0

^{*}PPI prescribed in patients on aspirin; PPI or H2 receptor blocker prescribed in patients on P2Y12 inhibitor

[^]Alerts addressed means the dashboard monitor staff (pharmacist/RN) notified provider or took other action (i.e., order labs, completing missing information).

2024 Michigan Anticoagulation Improvement Initiative (MAQI2) Collaborative Quality Initiative Performance Index Scorecard Memorial

Measurement Period: 01/01/2024 - 12/31/2024

Measure#	Weight	Measure Description	Points
1	10	Smoking status assessment in newly enrolled patients	
		≥90% of newly enrolled patients in 2022 will have smoking status assessed and documented per site protocol	10
		70-89% of newly enrolled patients in 2022 will have smoking status assessed and documented per site protocol	5
		50-69% of newly enrolled patients in 2022 will have smoking status assessed and documented per site protocol	3
		<50% of newly enrolled patients in 2022 will have smoking status assessed and documented per site protocol	0
		Inappropriate aspirin use in warfarin patients (updated 2024 criteria)	
	00	≤12% or relative decrease of ≥15%	30
2	30	13-19% or relative decrease of 10-14%	20
		20-24% or relative decrease of 5-9%	10
		>24% or relative decrease of <5%	0
		Extended International Normalized Ratio (INR) testing interval	
	10	≥80% of eligible patients received extended intervals	10
3		60-79% of eligible patients received extended intervals	5
		40-59% of eligible patients received extended intervals	3
		<40% of eligible patients received extended intervals	0
	10	Gastroprotection* in warfarin patients at high-risk for upper GI bleeding (site level)	
		≥55% or relative increase of ≥10%	10
4		40-54% or relative increase of 6-9%	5
		25-39% or relative increase of 2-5%	3
		<25% or relative increase of <2%	0
	10	Gastroprotection* in warfarin patients at high-risk for upper GI bleeding (consortium level)	
_		≥55% or relative increase of ≥10%	10
5		40-54% or relative increase of 6-9%	5
l		25-39% or relative increase of 2-5%	3
		<25% or relative increase of <2%	0
	10	Collaborative-wide meeting participation -Clinical Champion	
		attended all 3 meetings	10
6		attended 2 out of 3 meetings	7
		attended 1 out of 3 meetings	3
		did not attend any meetings	0

2024 Michigan Anticoagulation Improvement Initiative (MAQI2) Collaborative Quality Initiative Performance Index Scorecard Memorial

Measurement Period: 01/01/2024 - 12/31/2024

Measure#	Weight	Measure Description	Points
	10	Collaborative-wide meeting participation –	
		Coordinator/Lead Abstractor	
_		attended all 3 meetings	10
7		attended 2 out of 3 meetings	7
		attended 1 out of 3 meetings	4
		did not attend any meetings	0
	10	Completeness and Accuracy of data	
		Critical data elements are complete/accurate in ≥90%	10
8		of cases	10
		Critical data elements are complete/accurate in 70-89%	5
		of cases	ပ
		Critical data elements are complete/accurate in <70%	0
		of cases	J

^{*}PPI prescribed in patients on aspirin; PPI or H2 receptor blocker prescribed in patients on P2Y12 inhibitor

2024 Michigan Arthroplasty /Registry Collaborative Quality Initiative (MARCQI) Performance Index Scorecard

HOSPITAL - Year 1

Participation Measurement Period: 01/01/2024 - 11/30/2024 QI Measurement Period: 01/01/2024 - 06/30/2024

Measure#	Weight	Measure Description	Points
1	20	Collaborative Meeting Participation*-Clinical Champions (01.01.2024-11.30.2024) *Attendance at both the Medical Advisory Committee and Collaborative- wide meeting on February 2, 2024 (Virtual); June 7, 2024 (Grand Rapids, MI); and September 27, 2024 (Ann Arbor, MI) for 80% of the meeting time	
		3 out of 3 meetings attended	20
		2 out of 3 meetings attended	10
		<3 meetings attended	0
2	20	Collaborative Meeting Participation*-Clinical Data Abstractors (01.01.2024-11.30.2024) *Attendance at both the CDA Breakout and Collaborative-wide meeting on February 2, 2024 (Virtual); June 7, 2024 (Grand Rapids, MI); and September 27, 2024 (Ann Arbor, MI) for 80% of the meeting time	
		3 out of 3 meetings attended	20
		2 out of 3 meetings attended	2
		<3 meetings attended	0
4	5	New site kickoff: Completion of all necessary pre-meeting modules prior to attendance at 80% of the live/interactive New site kickoff meeting time on Wednesday, January 17, 2024 for: 1. Clinical Champion 2. Quality Administrator 3. Clinical Data Abstractor (if identified)	5
5	5	New site kickoff: Completion of all necessary pre-meeting modules prior to attendance at 80% of the live/interactive New site kickoff meeting time on Wednesday, January 17, 2024 for: 1. Site IT Support	5
6	20	Accuracy and Completeness of Data Submission (audits 01.01.2024-06.30.2024) - 5 metrics 1. Complete data entry (e.g. Data quality assurance and inclusion review scores) > 97% - 100% of the time 2. On-time data entry (e.g. Data abstraction completed 91-150 days post-op) > 97% - 100% of the time 3. Data abstraction started by May 31, 2024. 4. All cases performed or before May 4, 2024 abstracted by October 1, 2024 5. Documentation of utilization of all MARCQI FTEs awarded towards MARCQI activities or documentation of request to lower MARCQI FTE award to site submitted to MARCQI coordinating center by 11.30.2024 5 of 5 metrics met	20

2024 Michigan Arthroplasty /Registry Collaborative Quality Initiative (MARCQI) Performance Index Scorecard

HOSPITAL - Year 1

Participation Measurement Period: 01/01/2024 - 11/30/2024 QI Measurement Period: 01/01/2024 - 06/30/2024

Measure#	Weight	Measure Description	Points
		4 of 5 metrics met	16
		3 of 5 metrics met	12
		2 of 5 metrics met	8
		1 of 5 metrics met	4
		0 of 5 metrics met	0
		Access to Surgeon's Office Records (90 day events): (Surgery dates 01.01.2024-08.31.2024)	
7	15	90% + patient data captured	15
		75% - 89% patient data captured	7.5
		Less than 75% data captured	0
8	15	Site based Quality Meetings:(02.03.2024-11.15.2024) The site is awarded points for holding 3 meetings or more a year (following the MARCQI Collaborative meetings) to discuss surgeon, site based and collaborative outcomes with the orthopedic surgeons. The minimum requirement is 1 site based QI meeting after each MARCQI collaborative-wide meeting for a minimum total of 3 site based QI meetings. The CDA and Clinical Champion participate in the discussion and development of Quality Improvement plans. The Clinical Champion must be present to lead the discussion for the meeting to qualify as a site-based QI meeting following a collaborative-wide meeting. The site will complete the 'Site Based QI Meeting' sign-in form and send minutes/agenda to the Coordinating Center before the next Collaborative meeting.	15
Extra Credit	0	87.5% of MARCQI surgeons at the site attend 3 of 3 site-based QI meetings (02.03.2024-11.15.2024)meaning the MARCQI surgeons must attend 1 site based QI meeting after each MARCQI collaborative-wide meeting for a minimum total of 3 site based QI meetings. *Maximum total of 0.5 extra credit points per site	0.5
Extra credit	0	MODB data submission: Provide site's written planned process for site's submission of all MARCQI eligible cases data to Michigan Health and Hospital Association's Michigan Outpatient Database (MODB) by August 31, 2024 *Maximum total of 0.5 extra credit points per site	0.5
Extra Credit	0	Site based Quality Improvement: Alignment with MARCQI's 2024 demographic variables (e.g., AMA's recommendations for gender, race, ethnicity, etc.) If a site can document in their electronic health record that options for race, ethnicity, and gender are inclusive to MARCQI's 2024 definition (AMA recommendations) are available for MARCQI patients by June 30, 2024, extra credit points will be rewarded *Maximum total of 0.5 extra credit points per site	0.5

HOSPITAL - Year 2

QI Measurement Period: 07/01/2023 - 06/30/2024			
Measure#	Max. Weight	Measure Description	Points
1 20		Collaborative Meeting Participation*-Clinical Champions (01.01.2024-11.30.2024) *Attendance at both the Medical Advisory Committee and Collaborative- wide meeting on February 2, 2024 (Virtual); June 7, 2024 (Grand Rapids, MI); and September 27, 2024 (Ann Arbor, MI) for 80% of the meeting time 3 out of 3 meetings attended 2 out of 3 meetings attended <3 meetings attended	20 10 0
		Clinical Champion active engagement and participation in	U
Extra Credit	0	Quality Improvement working groups (e.g., PROS, ASC, Infection, Hip Fractures, Pain control, Device committee, Outlier work group, Quality Metrics work group, Executive Committee, Academic Quality Team) *Maximum total 0.5 point extra credit per site	0.5
2	20	Collaborative Meeting Participation*-Clinical Data Abstractors (01.01.2024-11.30.2024) *Attendance at both the CDA Breakout and Collaborative- wide meeting on February 2, 2024 (Virtual); June 7, 2024 (Grand Rapids, MI); and September 27, 2024 (Ann Arbor, MI) for 80% of the meeting time	20
		3 out of 3 meetings attended	20
		2 out of 3 meetings attended	10
Extra Credit	0	<3 meetings attended CDA active engagement and participation in CDA Committees or working groups (e.g., CDA Committee, Specifications Manual work group, PROS, ASC, Infection, Patient education, etc.) *Maximum total 0.5 point extra credit per site	0.5
Extra Credit	0	Site's with CDAs in their role for 9 months or greater as of January 31, 2024 who have at least one CDA attend the inperson CDA refresher training course. *Maximum total 0.5 point extra credit per site	
3	Accuracy and Completeness of Data Submission (audits 07.01.2023-06.30.2024) - 5 metrics 1. Complete data entry (e.g. Data quality assurance and inclusion review scores) > 97% - 100% of the time 2. On-time data entry (e.g. Data abstraction completed 91-150 days post-op) > 97% - 100% of the time 3. All 2023 cases abstracted completely by 05.31.2024 4. All cases performed or before May 4, 2024 abstracted by October 1, 2024 5. Documentation of utilization of all MARCQI FTEs awarded towards MARCQI activities or documentation of		

request to lower MARCQI FTE award to site submitted to MARCQI coordinating center by 11.30.2024 2 of 5 metrics met 16 3 of 5 metrics met 12 2 of 5 metrics met 12 2 of 5 metrics met 12 3 of 5 metrics met 4 0 of 5 metrics met 4 0 of 5 metrics met 4 0 of 5 metrics met 0 Site level: On-time data entry score is greater than or equal to 99% during the measurement period (07.01.2023 - 06.30.2024) AND complete data entry score is greater than or equal to 99% (QA & CI review score average) **Maximum total 0.5 point extra credit per site Site based Quality Meetings: (02.03.2024-11.15.2024) The site is awarded points for holding 3 meetings or more a year (following the MARCQI Collaborative meetings) to discuss surgeon, site based and collaborative outcomes with the orthopedic surgeons. The minimum requirement is 1 site based QI meeting after each MARCQI collaborative-wide meeting for a minimum total of 3 site based QI meeting for a minimum total of 3 site based QI meeting following a collaborative-wide meeting. The site will complete the 'Site Based QI weeting sign-in form and send minutes/agenda to the Coordinating Center before the next Collaborative meeting. 87.5% of MARCQI surgeons at the site attend 3 of 3 site-based QI meetings (02.03.2024-11.15.2024)—meaning the MARCQI collaborative-wide meeting for a minimum total of 3 extra credit points per site Extra Credit 8	Measure#	Max. Weight	Measure Description	
Extra Credit Site level: On-time data entry score is greater than or equal to 99% during the measurement period (07.01.2023 - 06.30.2024) AND complete data entry score is greater than or equal to 99% during the measurement period (07.01.2023 - 06.30.2024) AND complete data entry score is greater than or equal to 99% (QA & CI review score average) Maximum total 0.5 point extra credit per site				
Extra Credit Site level: On-time data entry score is greater than or equal to 99% during the measurement period (07.01.2023 - 06.30.2024) AND complete data entry score is greater than or equal to 99% during the measurement period (07.01.2023 - 06.30.2024) AND complete data entry score is greater than or equal to 99% (QA & CI review score average) "Maximum total 0.5 point extra credit per site Site based Quality Meetings: (02.03.2024-11.15.2024) The site is awarded points for holding 3 meetings or more a year (following the MARCQI Collaborative meetings) to discuss surgeon, site based and collaborative meetings) to discuss surgeon, site based and collaborative outcomes with the orthopedic surgeons. The minimum requirement is 1 site based QI meeting after each MARCQI collaborative-wide meeting for a minimum total of 3 site based QI meetings following a collaborative-wide meeting. The CDA and Clinical Champion participate in the discussion for the meeting to qualify as a site-based QI meeting following a collaborative-wide meeting. The site will complete the 'Site Based QI Meeting' sign-in form and send minutes/agenda to the Coordinating Center before the next Collaborative meeting. S7.5% of MARCQI surgeons at the site attend 3 of 3 site-based QI meetings (02.03.2024-11.15.2024)—meaning the MARCQI surgeons must attend 1 site based QI meeting after each MARCQI collaborative-wide meeting for a minimum total of 3 site based QI meetings. "Maximum total of 3 extra credit points per site Site level PROS Collection: Completed primary Pre-op and 2-16 week post-op HOOS JR or KOOS-JR + PROMIS10 (Overall average as of surgeries on or before 06.30.2024) When the difference between the PROS submission and completion rate at the site is >5%, the PROS completion rate will be used for this metric. The site is awarded full points for collection rates of 70%+			5 of 5 metrics met	20
Extra Credit Site level: On-time data entry score is greater than or equal to 99% during the measurement period (07.01.2023 - 06.30.2024) AND complete data entry score is greater than or equal to 99% (QA & CI review score average) *Maximum total 0.5 point extra credit per site			4 of 5 metrics met	16
Extra Credit Site level: On-time data entry score is greater than or equal to 99% during the measurement period (07.01.2023 - 06.30.2024) AND complete data entry score is greater than or equal to 99% (QA & CI review score average) *Maximum total 0.5 point extra credit per site Site based Quality Meetings: (02.03.2024-11.15.2024) The site is awarded points for holding 3 meetings or more a year (following the MARCQI Collaborative meetings) to discuss surgeon, site based and collaborative outcomes with the orthopedic surgeons. The minimum requirement is 1 site based QI meeting after each MARCQI collaborative-wide meeting for a minimum total of 3 site based QI meetings. The CDA and Clinical Champion participate in the discussion for the meeting to qualify as a site-based QI meeting following a collaborative-wide meeting. The site will complete the 'Site Based QI Meeting' sign-in form and send minutes/agenda to the Coordinating Center before the next Collaborative meeting. S7.5% of MARCQI surgeons at the site attend 3 of 3 site-based QI meetings (02.03.2024-11.15.2024)—meaning the MARCQI collaborative-wide meeting for a minimum total of 3 site based QI meetings after each MARCQI collaborative-wide meeting for a minimum total of 3 site based QI meetings. Site level PROS Collection: Completed primary Pre-op and 2-16 week post-op HOOS -JR or KOOS-JR + PROMIS10 (Overall average as of surgeries on or before 66.30.2024) When the difference between the PROS submission and completion rate at the site is 55%, the PROS completion rate will be used for this metric. The site is awarded full points for collection rates of 70%+			3 of 5 metrics met	12
Extra Credit Site level: On-time data entry score is greater than or equal to 99% during the measurement period (07.01.2023 - 06.30.2024) AND complete data entry score is greater than or equal to 99% (QA & CI review score average) *Maximum total 0.5 point extra credit per site			2 of 5 metrics met	8
Extra Credit Site level: On-time data entry score is greater than or equal to 99% during the measurement period (07.01.2023 - 06.30.2024) AND complete data entry score is greater than or equal to 99% (QA & CI review score average) *Maximum total 0.5 point extra credit per site			1 of 5 metrics met	4
to 99% during the measurement period (07.01.2023 - 06.30.2024) AND complete data entry score is greater than or equal to 99% (QA & CI review score average) *Maximum total 0.5 point extra credit per site Site based Quality Meetings:(02.03.2024-11.15.2024) The site is awarded points for holding 3 meetings or more a year (following the MARCQI Collaborative meetings) to discuss surgeon, site based and collaborative outcomes with the orthopedic surgeons. The minimum requirement is 1 site based QI meeting after each MARCQI collaborative-wide meetings. The CDA and Clinical Champion participate in the discussion and development of Quality Improvement plans. The Clinical Champion must be present to lead the discussion for the meeting to qualify as a site-based QI meeting following a collaborative-wide meeting. The site will complete the 'Site Based QI Meeting' sign-in form and send minutes/agenda to the Coordinating Center before the next Collaborative meeting. 87.5% of MARCQI surgeons at the site attend 3 of 3 site-based QI meetings (02.03.2024-11.15.2024)meaning the MARCQI collaborative-wide meeting for a minimum total of 3 site based QI meeting after each MARCQI collaborative-wide meeting for a minimum total of 3 site based QI meetings. *Maximum total of 3 extra credit points per site Site level PROS Collection: Completed primary Pre-op and 2-16 week post-op HOOS -JR or KOOS-JR + PROMIS10 (Overall average as of surgeries on or before 06.30.2024) When the difference between the PROS submission and completion rate at the site is >5%, the PROS completion rate will be used for this metric. The site is awarded full points for collection rates of 70%+			0 of 5 metrics met	0
Credit O	Extra		to 99% during the measurement period (07.01.2023 -	
*Maximum total 0.5 point extra credit per site Site based Quality Meetings:(02.03.2024-11.15.2024) The site is awarded points for holding 3 meetings or more a year (following the MARCQI Collaborative meetings) to discuss surgeon, site based and collaborative outcomes with the orthopedic surgeons. The minimum requirement is 1 site based QI meeting after each MARCQI collaborative-wide meeting for a minimum total of 3 site based QI meetings. The CDA and Clinical Champion participate in the discussion and development of Quality Improvement plans. The Clinical Champion must be present to lead the discussion for the meeting to qualify as a site-based QI meeting following a collaborative-wide meeting. The site will complete the 'Site Based QI Meeting' sign-in form and send minutes/agenda to the Coordinating Center before the next Collaborative meeting. 87.5% of MARCQI surgeons at the site attend 3 of 3 site-based QI meetings (02.03.2024-11.15.2024)meaning the MARCQI surgeons must attend 1 site based QI meeting after each MARCQI collaborative-wide meeting for a minimum total of 3 site based QI meetings. *Maximum total of 3 extra credit points per site Site level PROS Collection: Completed primary Pre-op and 2-16 week post-op HOOS -JR or KOOS-JR + PROMIS10 (Overall average as of surgeries on or before 06.30.2024) When the difference between the PROS submission and completion rate at the site is >5%, the PROS completion rate will be used for this metric. The site is awarded full points for collection rates of 70%+		0	, <u> </u>	0.5
The site is awarded points for holding 3 meetings or more a year (following the MARCQI Collaborative meetings) to discuss surgeon, site based and collaborative outcomes with the orthopedic surgeons. The minimum requirement is 1 site based QI meeting after each MARCQI collaborative-wide meeting for a minimum total of 3 site based QI meetings. The CDA and Clinical Champion participate in the discussion and development of Quality Improvement plans. The Clinical Champion must be present to lead the discussion for the meeting to qualify as a site-based QI meeting following a collaborative-wide meeting. The site will complete the 'Site Based QI Meeting' sign-in form and send minutes/agenda to the Coordinating Center before the next Collaborative meeting. 87.5% of MARCQI surgeons at the site attend 3 of 3 site-based QI meetings (02.03.2024-11.15.2024)meaning the MARCQI surgeons must attend 1 site based QI meeting after each MARCQI collaborative-wide meeting for a minimum total of 3 site based QI meetings. *Maximum total of 3 extra credit points per site Site level PROS Collection: Completed primary Pre-op and 2-16 week post-op HOOS -JR or KOOS-JR + PROMIS10 (Overall average as of surgeries on or before 06.30.2024) When the difference between the PROS submission and completion rate at the site is >5%, the PROS completion rate will be used for this metric. The site is awarded full points for collection rates of 70%+			*Maximum total 0.5 point extra credit per site	
Extra Credit Dassed QI meetings (02.03.2024-11.15.2024)meaning the MARCQI surgeons must attend 1 site based QI meeting after each MARCQI collaborative-wide meeting for a minimum total of 3 site based QI meetings. *Maximum total of 3 extra credit points per site Site level PROS Collection: Completed primary Pre-op and 2-16 week post-op HOOS -JR or KOOS-JR + PROMIS10 (Overall average as of surgeries on or before 06.30.2024) When the difference between the PROS submission and completion rate at the site is >5%, the PROS completion rate will be used for this metric. The site is awarded full points for collection rates of 70%+ 5	4	20	The site is awarded points for holding 3 meetings or more a year (following the MARCQI Collaborative meetings) to discuss surgeon, site based and collaborative outcomes with the orthopedic surgeons. The minimum requirement is 1 site based QI meeting after each MARCQI collaborative-wide meeting for a minimum total of 3 site based QI meetings. The CDA and Clinical Champion participate in the discussion and development of Quality Improvement plans. The Clinical Champion must be present to lead the discussion for the meeting to qualify as a site-based QI meeting following a collaborative-wide meeting. The site will complete the 'Site Based QI Meeting' sign-in form and send minutes/agenda to the Coordinating Center before the next Collaborative meeting.	20
and 2-16 week post-op HOOS -JR or KOOS-JR + PROMIS10 (Overall average as of surgeries on or before 06.30.2024) When the difference between the PROS submission and completion rate at the site is >5%, the PROS completion rate will be used for this metric. The site is awarded full points for collection rates of 70%+ 5		0	based QI meetings (02.03.2024-11.15.2024)meaning the MARCQI surgeons must attend 1 site based QI meeting after each MARCQI collaborative-wide meeting for a minimum total of 3 site based QI meetings.	3
60% 3	5	5	and 2-16 week post-op HOOS -JR or KOOS-JR + PROMIS10 (Overall average as of surgeries on or before 06.30.2024) When the difference between the PROS submission and completion rate at the site is >5%, the PROS completion rate will be used for this metric. The site is awarded full points for collection rates of	5
50% 1			50%	

Measure#	Max. Weight	Measure Description		
	vvoigin	The site is not awarded points if collection is less than 50%	0	
6	2.5	Site level PROS Collection: Completed primary Pre-op HOOS -JR or KOOS-JR + PROMIS10 (Surgeries on from July 1, 2023 to June 30, 2024) When the difference between the PROS submission and completion rate at the site is >5%, the PROS completion rate will be used for this metric.		
0	2.5	The site is awarded full points for collection rates of 90%+	2.5	
		75 - 89%	1.5	
		The site is not awarded points if collection is less than 60%	0.5	
		% of Opioid naive THA patients at the SITE meet the MARCQI Pain control pathway guidelines (<240 OME)		
7	2.5	85% or greater of THA patients meet the guidelines of 240 OME or less	2.5	
		60-84% of THA patients prescribed <240 OME	1.5	
		Less than 60% of patients meet the prescribing criteria	0	
		% of Opioid naive TKA patients at the SITE meet the MARCQI Pain control pathway guidelines (<320 OME)		
8	2.5	90% or greater of TKA patients meet the guidelines of 320 OME or less	2.5	
		70-89% of TKA patients prescribed <320 OME	1.5	
		Less than 70% of patients meet the prescribing criteria	0	
9		90-Day Hip fracture: Reduce COLLABORATIVE rate of 90-Day Hip fracture for all primary HIP procedures (excluding conversions) by 10% (Baseline FY2023 = 1.17%; 10% reduction goal = 1.05%)		
	2.5	1.05% or less of all primary HIPS experience a 90- Day hip fracture	2.5	
		Greater than 1.05% but less than 1.17% of all primary HIPS experience a 90-Day hip fracture	1.5	
		The site is not awarded points if there is no collaborative-wide improvement (>=1.17%) in all primary HIPS with 90-Day hip fractures	0	

Measure#	Max. Weight	Measure Description	Points
10	5	90-Day Hip fracture: Reduce baseline FY2023 rate by 10%, achieve, or maintain SITE level rate of 90-Day Hip fracture for all primary HIP procedures (excluding conversions) to 1.05%Reduce by 10%: If the site's April 2023 Quarterly report shows a 90-Day hip fracture rate for all primary HIP procedures (excluding conversion) that is greater than 1.05%, the site-level goal will be to reduce this rate by 10% or meet the 1.05% rate at their siteMaintain site-level: If the site's April 2023 Quarterly report shows a 90-Day hip fracture rate for all primary HIP procedures (excluding conversion) to be less than or equal to 1.17%, the site-level goal will be to maintain or improve upon their baseline rateFinal determination will be made by the January 2025 Quarterly Report When April 2023 baseline is: >1.05%, the site meets a site-level reduction of 10% or meets the 1.05% rate of 90-day hip fractures ≤1.05%, the site meets a site-level reduction of ≥ 5% site-level rate of 90-day hip fractures ≤1.05%, the site sees a site-level increase of ≤ 2% site- level rate of 90-day hip fractures When April 2023 baseline is: >1.05%, the site sees a site-level reduction is <5% or increases the site-level rate of 90-day hip fractures ≤1.05%, the site attains a site-level reduction is <5% or increases the site-level rate of 90-day hip fractures ≤1.05%, the site sees a site-level reduction is <5% or increases the site-level rate of 90-day hip fractures	5 2.5
Extra Credit 0		level rate of 90-day hip fractures Site based Quality Improvement: Alignment with MARCQI's 2024 demographic variables (e.g., AMA's recommendations for gender, race, ethnicity, etc.) If a site can document in their electronic health record that options for race, ethnicity, and gender are inclusive to MARCQI's 2024 definition (AMA recommendations) are available for MARCQI patients by December 31, 2023, extra credit points will be rewarded *Maximum total of 2 extra credit points per site	2

HOSPITAL - Year 3+

Measure#	Weight	Measure Description	
1	10	Collaborative Meeting Participation*-Clinical Champions (01.01.2024-11.30.2024) *Attendance at both the Medical Advisory Committee and Collaborative- wide meeting on February 2, 2024 (Virtual); June 7, 2024 (Grand Rapids, MI); and September 27, 2024 (Ann Arbor, MI) for 80% of the meeting time	
		3 out of 3 meetings attended	10
		2 out of 3 meetings attended	5
		<3 meetings attended	0
Extra Credit	0	Clinical Champion active engagement and participation in Quality Improvement working groups (e.g., PROS, ASC, Infection, Hip Fractures, Pain control, Device committee, Outlier work group, Quality Metrics work group, Executive Committee, Academic Quality Team) *Maximum total 0.5 point extra credit per site	0.5
2 5		Collaborative Meeting Participation*-Clinical Data Abstractors (01.01.2024-11.30.2024) *Attendance at both the CDA Breakout and Collaborative-wide meeting on February 2, 2024 (Virtual); June 7, 2024 (Grand Rapids, MI); and September 27, 2024 (Ann Arbor, MI) for 80% of the meeting time	
		3 out of 3 meetings attended	5
		2 out of 3 meetings attended	2
		<3 meetings attended	0
Extra Credit	0	CDA active engagement and participation in CDA Committees or working groups (e.g., CDA Committee, Specifications Manual work group, PROS, ASC, Infection, Patient education, etc.) *Maximum total 0.5 point extra credit per site	0.5
Extra Credit	0	Site's with CDAs in their role for 9 months or greater as of January 31, 2024 who have at least one CDA attend the inperson CDA refresher training course. *Maximum total 0.5 point extra credit per site	
3	10	Accuracy and Completeness of Data Submission (audits 07.01.2023-06.30.2024) - 5 metrics 1. Complete data entry (e.g. Data quality assurance and inclusion review scores) > 97% - 100% of the time 2. On-time data entry (e.g. Data abstraction completed 91-150 days post-op) > 97% - 100% of the time 3. All 2023 cases abstracted completely by 05.31.2024 4. All cases performed or before May 4, 2024 abstracted by October 1, 2024 5. Documentation of utilization of all MARCQI FTEs awarded towards MARCQI activities or documentation of request to lower MARCQI FTE award to site submitted to MARCQI coordinating center by 11.30.2024	

HOSPITAL - Year 3+

Measure#	Weight	Measure Description		
		4 of 5 metrics met	8	
		3 of 5 metrics met	6	
		2 of 5 metrics met	4	
		1 of 5 metrics met	2	
		0 of 5 metrics met	0	
Extra Credit	0	Site level: On-time data entry score is greater than or equal to 99% during the measurement period (07.01.2023 - 06.30.2024) AND complete data entry score is greater than or equal to 99% (QA & CI review score average) *Maximum total 0.5 point extra credit per site	0.5	
		Site based Quality Meetings:(02.03.2024-11.15.2024)		
4	5	The site is awarded points for holding 3 meetings or more a year (following the MARCQI Collaborative meetings) to discuss surgeon, site based and collaborative outcomes with the orthopedic surgeons. The minimum requirement is 1 site based QI meeting after each MARCQI collaborative-wide meeting for a minimum total of 3 site based QI meetings. The CDA and Clinical Champion participate in the discussion and development of Quality Improvement plans. The Clinical Champion must be present to lead the discussion for the meeting to qualify as a site-based QI meeting following a collaborative-wide meeting. The site will complete the 'Site Based QI Meeting' sign-in form and send minutes/agenda to the Coordinating Center before the next Collaborative meeting.		
Extra Credit	0	87.5% of MARCQI surgeons at the site attend 3 of 3 site-based QI meetings (02.03.2024-11.15.2024)meaning the MARCQI surgeons must attend 1 site based QI meeting after each MARCQI collaborative-wide meeting for a minimum total of 3 site based QI meetings. *Maximum total of 3 extra credit points per site	3	
5	10	Site level PROS Collection: Completed primary Pre-op and 2-16 week post-op HOOS -JR or KOOS-JR + PROMIS10 (Overall average as of surgeries on or before 06.30.2024) When the difference between the PROS submission and completion rate at the site is >5%, the PROS completion rate will be used for this metric. The site is awarded full points for collection rates of 70%+ 60% The site is not awarded points if collection is less	10 5 2.5	
		than 50%	0	

HOSPITAL - Year 3+

Measure#	Weight	Measure Description		
Extra 0		Site level PROS Collection: Completed primary Pre-op and 1-year post-op HOOS -JR or KOOS-JR + PROMIS10 (Overall average as of surgeries on 06.01.2022 to 11.03.2023 and 1 year post op300-425 days post-op accepted.) When the difference between the PROS submission and completion rate at the site is >5%, the PROS completion rate will be used for this metric.		
Credit		The site is awarded full points for collection rates of 50%+	5	
		40%	3	
		30%	1	
		<30%	0	
	Site level PROS Collection: Completed primary Pre-op HOOS -JR or KOOS-JR + PROMIS10 (Surgeries on from July 1, 2023 to June 30, 2024) When the difference between the PROS submission and completion rate at the site is >5%, the PROS completion rate will be used for this metric.			
6	5	The site is awarded full points for collection rates of 90%+	5	
		75 - 89%	3	
		60 - 74%	1	
		The site is not awarded points if collection is less than 60%	0	
		% of Opioid naive THA patients at the SITE meet the MARCQI Pain control pathway guidelines (<240 OME)		
7	5	85% or greater of THA patients meet the guidelines of 240 OME or less	5	
		60-84% of THA patients prescribed <240 OME	2	
		Less than 60% of patients meet the prescribing criteria	0	
		% of Opioid naive TKA patients at the SITE meet the MARCQI Pain control pathway guidelines (<320 OME)		
8	90% or greater of TKA patie	90% or greater of TKA patients meet the guidelines of 320 OME or less	5	
		70-89% of TKA patients prescribed <320 OME	2	
		Less than 70% of patients meet the prescribing criteria	0	
9	15	90-Day Hip fracture: Reduce COLLABORATIVE rate of 90-Day Hip fracture for all primary HIP procedures (excluding conversions) by 10% (Baseline FY2023 = 1.17%; 10% reduction goal = 1.05%) 1.05% or less of all primary HIPS experience a 90-		
		Day hip fracture	15	

2024 Michigan Arthroplasty Registry Collaborative Quality Initiative (MARCQI)

Performance Index Scorecard

HOSPITAL - Year 3+

Measure#	Weight	·	
		Greater than 1.05% but less than 1.17% of all	7
		primary HIPS experience a 90-Day hip fracture	'
		The site is not awarded points if there is no	
		collaborative-wide improvement (>=1.17%) in all primary	0
		HIPS with 90-Day hip fractures	
		90-Day Hip fracture: Reduce baseline FY2023 rate by 10%, achieve, or maintain SITE level rate of 90-Day Hip fracture for all primary HIP procedures (excluding conversions) to 1.05%Reduce by 10%: If the site's April 2023 Quarterly report shows a 90-Day hip fracture rate for all primary HIP procedures (excluding conversion) that is greater than 1.05%, the site-level goal will be to reduce this rate by 10% or meet the 1.05% rate at their siteMaintain site-level: If the site's April 2023 Quarterly report shows a 90-Day hip fracture rate for all primary HIP procedures (excluding conversion) to be less than or equal to 1.17%, the site-level goal will be to maintain or improve upon their baseline rateFinal determination will be made by the January 2025	
10	10	Quarterly Report When April 2023 baseline is: >1.05%, the site meets a site-level reduction of 10% or meets the 1.05% rate of 90-day hip fractures site-level-reduction of 10% or meets the 1.05% rate of 90-day hip fractures site-level-reduction of 10% or meets the 1.05%, the site maintains baseline outcome or reduces hip fracture rate	10
		When April 2023 baseline is: >1.05%, the site meets a site-level reduction of ≥ 5% site-level rate of 90-day hip fractures ≤1.05%, the site sees a site-level increase of ≤ 2% site-level rate of 90-day hip fractures	5
		When April 2023 baseline is: >1.05%, the site attains a site-level reduction is <5% or increases the site-level rate of 90-day hip fractures < 1.05% , the site sees a site-level increase of >2% site-level rate of 90-day hip fractures	0
11	20	Implementation of one site specific quality initiative (linked to a MARCQI quality initiative). If red on scorecard of April, 2023, you must strongly consider choosing this as the project. If you have worked on this metric previously or have difficultly developing a goal, a yellow target area may be chosen with the Program Manager's approval. If you would like to continue with your FY2023 site based QI project topic, please discuss with the Program Manager. If no red, you will choose a 'yellow'. Progress Reports are due in May 2024 & January 2025.	

HOSPITAL - Year 3+

Measure#	Weight	Measure Description	
		Final results are based on quarterly reports of <u>January</u> , <u>2025</u>	
		Plan submitted and approved, reporting requirements & timelines met, A3 submitted, and goal met* *Clinical Champion and Clinical Data Abstractor must be available to present at June 2025 Collaborative-wide session if asked *Not presenting if asked will yield -20 points on the FY2025 P4P scorecard	20
		Plan submitted and approved Reporting requirements are met & timelines, but the target identified is not met. A3 submitted with final report and presentation by clinical champion and clinical data abstractor given at June 2025 MARCQI Collaborative-wide sessions* *Clinical Champion and Clinical Data Abstractor must be available to present at June 2025 Collaborative-wide session if asked *Not presenting if asked will yield -20 points on the FY2025 P4P scorecard	10
		Plan submitted and approved All reporting requirements are met, but the identified goal and timelines are not met.	5
		Plan is not developed, reports not done.	0
Extra Credit	I II I ontions for race athnicity, and dender are inclusive to		2

HOSPITAL - Extra Credit

Measure#	Max.		Points	Eligible P4P
Extra Credit	0	Clinical Champion active engagement and participation in Quality Improvement working groups (e.g., PROS, ASC, Infection, Hip Fractures, Pain control, Device committee, Outlier work group, Quality Metrics work group, Executive Committee, Academic Quality Team) *Maximum total 0.5 point extra credit per site	0.5	2, 3+
Extra Credit	0	CDA active engagement and participation in CDA Committees or working groups (e.g., CDA Committee, Specifications Manual work group, PROS, ASC, Infection, Patient education, etc.) *Maximum total 0.5 point extra credit per site	0.5	2, 3+
Extra Credit	0	Site's with CDAs in their role for 9 months or greater as of January 31, 2024 who have at least one CDA attend the in-person CDA refresher training course. *Maximum total 0.5 point extra credit per site	0.5	2, 3+
Extra Credit	0	Site level: On-time data entry score is greater than or equal to 99% during the measurement period (07.01.2023 - 06.30.2024) AND complete data entry score is greater than or equal to 99% (QA & CI review score average) *Maximum total 0.5 point extra credit per site	0.5	2, 3+
Extra Credit	0	87.5% of MARCQI surgeons at the site attend 3 of 3 site-based QI meetings (02.03.2024-11.15.2024)meaning the MARCQI surgeons must attend 1 site based QI meeting after each MARCQI collaborative-wide meeting for a minimum total of 3 site based QI meetings. *Maximum total of 3 extra credit points per site	3	2, 3+

HOSPITAL - Extra Credit

Measure#	Max.		Points	Eligible P4P
Extra Credit	0	87.5% of MARCQI surgeons at the site attend 3 of 3 site-based QI meetings (02.03.2024-11.15.2024)meaning the MARCQI surgeons must attend 1 site based QI meeting after each MARCQI collaborative-wide meeting for a minimum total of 3 site based QI meetings. *Maximum total of 0.5 extra credit points per site	0.5	1
Extra Credit	0	Site level PROS Collection: Completed primary Pre-op and 1-year post-op HOOS - JR or KOOS-JR + PROMIS10 (Overall average as of surgeries on 06.01.2022 to 11.03.2023 and 1 year post op300-425 days post-op accepted.) When the difference between the PROS submission and completion rate at the site is >5%, the PROS completion rate will be used for this metric. The site is awarded full points for collection rates of 50%+ 40% 30%	5 3 1 0	3+
Extra Credit	0	Site based Quality Improvement: Alignment with MARCQI's 2024 demographic variables (e.g., AMA's recommendations for gender, race, ethnicity, etc.) If a site can document in their electronic health record that options for race, ethnicity, and gender are inclusive to MARCQI's 2024 definition (AMA recommendations) are available for MARCQI patients by December 31, 2023, extra credit points will be rewarded *Maximum total of 2 extra credit points per site	2	2, 3+

HOSPITAL - Extra Credit

Measure#	Max.		Points	Eligible P4P
Extra Credit	0	Site based Quality Improvement: Alignment with MARCQI's 2024 demographic variables (e.g., AMA's recommendations for gender, race, ethnicity, etc.) If a site can document in their electronic health record that options for race, ethnicity, and gender are inclusive to MARCQI's 2024 definition (AMA recommendations) are available for MARCQI patients by June 30, 2024, extra credit points will be rewarded *Maximum total of 0.5 extra credit points per site	0.5	1

Period: 01/01/2024 - 06/30/2024

Measure#	Max.	1 enod. 01/01/2024 - 00/30/2024	Dointo
ivieasure #	Weight	Measure Description	Points
1	20	COLLABORATIVE MEETING PARTICIPATION*- CLINICAL CHAMPIONS (01.01.2024-11.30.2024) *ATTENDANCE AT BOTH THE MEDICAL ADVISORY COMMITTEE AND COLLABORATIVE-WIDE MEETING ON FEBRUARY 2, 2024 (VIRTUAL); JUNE 7, 2024 (GRAND RAPIDS, MI); AND SEPTEMBER 27, 2024 (ANN ARBOR, MI) FOR 80% OF THE MEETING TIME 3 out of 3 meetings attended 2 out of 3 meetings attended <3 meetings attended	20 10 0
2	20	Collaborative Meeting Participation*-Clinical Data Abstractors (01.01.2024-11.30.2024) *Attendance at both the CDA Breakout and Collaborative-wide meeting on February 2, 2024 (Virtual); June 7, 2024 (Grand Rapids, MI); and September 27, 2024 (Ann Arbor, MI) for 80% of the meeting time	-
		3 out of 3 meetings attended	20
		2 out of 3 meetings attended	2
		<3 meetings attended	0
3	5	Collaborative Meeting Participation*-Quality Administrator (01.01.2024-11.30.2024) *Attendance at the Collaborative-wide meeting in February 2024; June 2024 and September 2024 for 80% of the meeting time	
		3 out of 3 meetings attended	5
		2 out of 3 meetings attended	2
		<3 meetings attended	0
4	5	New site kickoff: Completion of all necessary pre-meeting modules prior to attendance at 80% of the live/interactive New site kickoff meeting time on Wednesday, January 17, 2024 for: 1. Clinical Champion 2. Quality Administrator 3. Clinical Data Abstractor (if identified)	5
5	5	New site kickoff: Completion of all necessary pre-meeting modules prior to attendance at 80% of the live/interactive New site kickoff meeting time on Wednesday, January 17, 2024 for: 1. Site IT Support	5

Period: 01/01/2024 - 06/30/2024

	Mov	Period: 01/01/2024 - 06/30/2024	
Measure #	Max. Weight	Measure Description	Points
6	20	Accuracy and Completeness of Data Submission (audits 01.01.2024-06.30.2024) - 5 metrics 1. Complete data entry (e.g. Data quality assurance and inclusion review scores) > 97% - 100% of the time 2. On-time data entry (e.g. Data abstraction completed 91-150 days post-op) > 97% - 100% of the time 3. Data abstraction started by May 31, 2024. 4. All cases performed or before May 4, 2024 abstracted by October 1, 2024 5. Attestation of BCBSM ASF CQI participation funding utilization to support dedicated data abstraction support for effective and complete MARCQI data abstraction and quality improvement work to the MARCQI Coordinating Center by 11.30.2024	
		5 of 5 metrics met	20
		4 of 5 metrics met	16
		3 of 5 metrics met	12
		2 of 5 metrics met	8
		1 of 5 metrics met	4
		0 of 5 metrics met	0
	12.5	Access to Surgeon's Office Records (90 day events): (Surgery dates 01.01.2024-08.31.2024)	
7		90% + patient data captured	12.5
		75% - 89% patient data captured	6
		Less than 75% data captured	0
8	12.5	Site based Quality Meetings:(02.03.2024-11.15.2024) The site is awarded points for holding 3 meetings or more a year (following the MARCQI Collaborative meetings) to discuss surgeon, site based and collaborative outcomes with the orthopedic surgeons. The minimum requirement is 1 site-based QI meeting after each MARCQI collaborative-wide meeting for a minimum total of 3 site based QI meetings. The CDA and Clinical Champion participate in the discussion and development of Quality Improvement plans. The Clinical Champion must be present to lead the discussion for the meeting to qualify as a site-based QI meeting following a collaborative-wide meeting. The site will complete the 'Site Based QI Meeting' sign-in form and send minutes/agenda to the Coordinating Center before the next Collaborative meeting.	12.5

2024 Michigan Arthroplasty Registry Collaborative Quality Initiative (MARCQI)

Performance Incentive Scorecard

For Ambulatory Surgery Facility/Center (ASF/ASC) - Year 1

Participation Measurement Period: 01/01/2024 - 11/30/2024 QI Measurement

Period: 01/01/2024 - 06/30/2024

Feliod: 01/01/2024 - 00/30/2024			
Measure#	Max. Weight	Measure Description	Points
EXTRA CREDIT	0	MODB data submission: Provide site's written planned process for site's submission of all MARCQI eligible cases data to Michigan Health and Hospital Association's Michigan Outpatient Database (MODB) by August 31, 2024 *Maximum total 0.5 point extra credit per site	0.5
EXTRA CREDIT	0	Site based Quality Improvement: Alignment with MARCQI's 2024 demographic variables (e.g., AMA's recommendations for gender, race, ethnicity, etc.) If a site can document in their electronic health record that options for race, ethnicity, and gender are inclusive to MARCQI's 2024 definition (AMA recommendations) are available for MARCQI patients by June 30, 2024, extra credit points will be rewarded *Maximum total 0.5 point extra credit per site	0.5

ASC - Year 2

Measure#	Max. Weight	Measure Description	Points
1	20	Collaborative Meeting Participation*-Clinical Champions (01.01.2024-11.30.2024) *Attendance at both the Medical Advisory Committee and Collaborative- wide meeting on February 2, 2024 (Virtual); June 7, 2024 (Grand Rapids, MI); and September 27, 2024 (Ann Arbor, MI) for 80% of the meeting time	
		3 out of 3 meetings attended	20
		2 out of 3 meetings attended	10
Extra Credit	0	<3 meetings attended Clinical Champion active engagement and participation in Quality Improvement working groups (e.g., PROS, ASC, Infection, Hip Fractures, Pain control, Device committee, Outlier work group, Quality Metrics work group, Executive Committee, Academic Quality Team) *Maximum total 0.5 point extra credit per site	0.5
2	20	Collaborative Meeting Participation*-Clinical Data Abstractors (01.01.2024-11.30.2024) *Attendance at both the CDA Breakout and Collaborative-wide meeting on February 2, 2024 (Virtual); June 7, 2024 (Grand Rapids, MI); and September 27, 2024 (Ann Arbor, MI) for 80% of the meeting time 3 out of 3 meetings attended	20
		2 out of 3 meetings attended	10
Extra Credit	0	<3 meetings attended CDA active engagement and participation in CDA Committees or working groups (e.g., CDA Committee, Specifications Manual work group, PROS, ASC, Infection, Patient education, etc.) *Maximum total 0.5 point extra credit per site	0.5
Extra Credit	0	Site's with CDAs in their role for 9 months or greater as of January 31, 2024 who have at least one CDA attend the inperson CDA refresher training course. *Maximum total 0.5 point extra credit per site	0.5
Extra Credit	0	Collaborative Meeting Participation*- Site Administrator (01.01.2024-11.30.2024) *Attendance at 3 of 3 Collaborative-wide meeting in February 2024; June 2024; and September 2024 Individual must be identified and confirmed to fill this role by 1st 2024 MARCQI Collaborative-wide meeting *Maximum total 0.5 point extra credit per site	0.5

ASC - Year 2

Measure#	Max. Weight	Measure Description	Points
3	20	Accuracy and Completeness of Data Submission (audits 07.01.2023-06.30.2024) - 5 metrics 1. Complete data entry (e.g. Data quality assurance and inclusion review scores) > 97% - 100% of the time 2. On-time data entry (e.g. Data abstraction completed 91-150 days post-op) > 97% - 100% of the time 3. All 2023 cases abstracted completely by 05.31.2024 4. All cases performed or before May 4, 2024 abstracted by October 1, 2024 5. Attestation of BCBSM ASF CQI participation funding utilization to support dedicated data abstraction support for effective and complete MARCQI data abstraction and quality improvement work to the MARCQI Coordinating Center by 11.30.2024	
		5 of 5 metrics met	20
		4 of 5 metrics met	16
		3 of 5 metrics met	12
		2 of 5 metrics met	8
		1 of 5 metrics met	4
		0 of 5 metrics met	0
Extra Credit	0	Site level: On-time data entry score is greater than or equal to 99% during the measurement period (07.01.2023 - 06.30.2024) AND complete data entry score is greater than or equal to 99% (QA & CI review score average) *Maximum total 0.5 point extra credit per site	0.5
4	20	Site based Quality Meetings:(02.03.2024-11.15.2024) The site is awarded points for holding 3 meetings or more a year (following the MARCQI Collaborative meetings) to discuss surgeon, site based and collaborative outcomes with the orthopedic surgeons. The minimum requirement is 1 site based QI meeting after each MARCQI collaborative-wide meeting for a minimum total of 3 site based QI meetings. The CDA and Clinical Champion participate in the discussion and development of Quality Improvement plans. The Clinical Champion must be present to lead the discussion for the meeting to qualify as a site-based QI meeting following a collaborative-wide meeting. The site will complete the 'Site Based QI Meeting' sign-in form and send minutes/agenda to the Coordinating Center before the next Collaborative meeting.	20
Extra Credit	0	Access to Surgeon's Office Records (90 day events): (Surgery dates 07.01.2023 - 06.30.2024): 90% + patient data captured *Maximum total 1 point extra credit per site	1

ASC - Year 2

Measure #	Max. Weight	Measure Description	Points
		Site level PROS Collection: Completed primary Pre-op and 2-16 week post-op HOOS -JR or KOOS-JR + PROMIS10 (Overall average as of surgeries on or before 06.30.2024) When the difference between the PROS submission and completion rate at the site is >5%, the PROS completion rate will be used for this metric.	
5	5	The site is awarded full points for collection rates of 70%+	5
		60%	3
		50%	1
		The site is not awarded points if collection is less than 50%	0
		Site level PROS Collection: Completed primary Pre-op HOOS -JR or KOOS-JR + PROMIS10 (Surgeries on from July 1, 2023 to June 30, 2024) When the difference between the PROS submission and completion rate at the site is >5%, the PROS completion rate will be used for this metric.	
6	2.5	The site is awarded full points for collection rates of 90%+	2.5
		75 - 89%	1.5
		60 - 74%	0.5
		The site is not awarded points if collection is less than 60%	0
		% of Opioid naive THA patients at the SITE meet the MARCQI Pain control pathway guidelines (<240 OME) 85% or greater of THA patients meet the guidelines	
7	2.5	of 240 OME or less	2.5
		60-84% of THA patients prescribed <240 OME	1.5
		Less than 60% of patients meet the prescribing criteria	0
		% of Opioid naive TKA patients at the SITE meet the MARCQI Pain control pathway guidelines (<320 OME)	
8	2.5	90% or greater of TKA patients meet the guidelines of 320 OME or less	2.5
		70-89% of TKA patients prescribed <320 OME	1.5
		Less than 70% of patients meet the prescribing criteria	0
9	2.5	90-Day Hip fracture: Reduce COLLABORATIVE rate of 90-Day Hip fracture for all primary HIP procedures (excluding conversions) by 10% (Baseline FY2023 = 1.17%; 10% reduction goal = 1.05%)	

ASC - Year 2

Measure #	Max. Weight	Measure Description	Points
		1.05% or less of all primary HIPS experience a 90- Day hip fracture	2.5
		Greater than 1.05% but less than 1.17% of all primary HIPS experience a 90-Day hip fracture	1.5
		The site is not awarded points if there is no collaborative-wide improvement (>=1.17%) in all primary HIPS with 90-Day hip fractures	0
10	5	90-Day Hip fracture: Reduce baseline FY2023 rate by 10%, achieve, or maintain SITE level rate of 90-Day Hip fracture for all primary HIP procedures (excluding conversions) to 1.05%Reduce by 10%: If the site's April 2023 Quarterly report shows a 90-Day hip fracture rate for all primary HIP procedures (excluding conversion) that is greater than 1.05%, the site-level goal will be to reduce this rate by 10% or meet the 1.05% rate at their siteMaintain site-level: If the site's April 2023 Quarterly report shows a 90-Day hip fracture rate for all primary HIP procedures (excluding conversion) to be less than or equal to 1.17%, the site-level goal will be to maintain or improve upon their baseline rateFinal determination will be made by the January 2025 Quarterly Report	
		When April 2023 baseline is: >1.05%, the site meets a site-level reduction of 10% or meets the 1.05% rate of 90-day hip fractures ≤1.05%, the site maintains baseline outcome or reduces hip fracture rate	5
		When April 2023 baseline is: >1.05%, the site meets a site-level reduction of ≥ 5% site-level rate of 90-day hip fractures ≤1.05%, the site sees a site-level increase of ≤ 2% site- level rate of 90-day hip fractures	2.5
		When April 2023 baseline is: >1.05%, the site attains a site-level reduction is <5% or increases the site-level rate of 90-day hip fractures <1.05%, the site sees a site-level increase of >2% site-level rate of 90-day hip fractures	0

ASC - Year 2

Measure #	Max. Weight	Measure Description	Points
Extra Credit	0	Site based Quality Improvement: Alignment with MARCQI's 2024 demographic variables (e.g., AMA's recommendations for gender, race, ethnicity, etc.) If a site can document in their electronic health record that options for race, ethnicity, and gender are inclusive to MARCQI's 2024 definition (AMA recommendations) are available for MARCQI patients by December 31, 2023, extra credit points will be rewarded *Maximum total 2 point extra credit per site	2

QI Measurement Period: 07/01/2023 - 06/30/2024

Measure#	Max. Weight	Measure Description	Points
1	10	Collaborative Meeting Participation*-Clinical Champions (01.01.2024-11.30.2024) *Attendance at both the Medical Advisory Committee and Collaborative-wide meeting on February 2, 2024 (Virtual); June 7, 2024 (Grand Rapids, MI); and September 27, 2024 (Ann Arbor, MI) for 80% of the meeting time	
		3 out of 3 meetings attended	10
		2 out of 3 meetings attended	5
		<3 meetings attended	0
Extra Credit	0	Clinical Champion active engagement and participation in Quality Improvement working groups (e.g., PROS, ASC, Infection, Hip Fractures, Pain control, Device committee, Outlier work group, Quality Metrics work group, Executive Committee, Academic Quality Team) *Maximum total 0.5 point extra credit per site	0.5
2	5	Collaborative Meeting Participation*-Clinical Data Abstractors (01.01.2024-11.30.2024) *Attendance at both the CDA Breakout and Collaborative-wide meeting on February 2, 2024 (Virtual); June 7, 2024 (Grand Rapids, MI); and September 27, 2024 (Ann Arbor, MI) for 80% of the meeting time 3 out of 3 meetings attended	5
		2 out of 3 meetings attended	2
		<3 meetings attended	0
Extra Credit	0	CDA active engagement and participation in CDA Committees or working groups (e.g., CDA Committee, Specifications Manual work group, PROS, ASC, Infection, Patient education, etc.) *Maximum total 0.5 point extra credit per site	0.5
Extra Credit	0	Site's with CDAs in their role for 9 months or greater as of January 31, 2024 who have at least one CDA attend the inperson CDA refresher training course. *Maximum total 0.5 point extra credit per site	0.5
Extra Credit	0	Collaborative Meeting Participation*- Site Administrator (01.01.2024-11.30.2024) *Attendance at 3 of 3 Collaborative-wide meeting in February 2, 2024 (Virtual); June 7, 2024 (Grand Rapids, MI); and September 27, 2024 (Ann Arbor, MI) Individual must be identified and confirmed to fill this role by 1st 2024 MARCQI Collaborative-wide meeting *Maximum total 0.5 point extra credit per site	0.5

QI Measurement Period: 07/01/2023 - 06/30/2024

Measure#	Max. Weight	Measure Description	Points
3	10	Accuracy and Completeness of Data Submission (audits 07.01.2023-06.30.2024) - 5 metrics 1. Complete data entry (e.g. Data quality assurance and inclusion review scores) > 97% - 100% of the time 2. On-time data entry (e.g. Data abstraction completed 91-150 days post-op) > 97% - 100% of the time 3. All 2023 cases abstracted completely by 05.31.2024 4. All cases performed or before May 4, 2024 abstracted by October 1, 2024 5. Attestation of BCBSM ASF CQI participation funding utilization to support dedicated data abstraction support for effective and complete MARCQI data abstraction and quality improvement work to the MARCQI Coordinating Center by 11.30.2024	
		5 of 5 metrics met	10
		4 of 5 metrics met	8
		3 of 5 metrics met	6
		2 of 5 metrics met	4
		1 of 5 metrics met 0 of 5 metrics met	0
		Site level: On-time data entry score is greater than or equal	U
Extra Credit	0	to 99% during the measurement period (07.01.2023 - 06.30.2024) AND complete data entry score is greater than or equal to 99% (QA & CI review score average) *Maximum total 0.5 point extra credit per site	0.5
4	5	Site based Quality Meetings:(02.03.2024-11.15.2024) The site is awarded points for holding 3 meetings or more a year (following the MARCQI Collaborative meetings) to discuss surgeon, site based and collaborative outcomes with the orthopedic surgeons. The minimum requirement is 1 site based QI meeting after each MARCQI collaborative-wide meeting for a minimum total of 3 site based QI meetings. The CDA and Clinical Champion participate in the discussion and development of Quality Improvement plans. The Clinical Champion must be present to lead the discussion for the meeting to qualify as a site-based QI meeting following a collaborative-wide meeting. The site will complete the 'Site Based QI Meeting' sign-in form and send minutes/agenda to the Coordinating Center before the next Collaborative meeting.	5

Ql Measurement Period: 07/01/2023 - 06/30/2024

Measure#	Max. Weight	Measure Description	Points
Extra Credit	0	Access to Surgeon's Office Records (90 day events): (Surgery dates 07.01.2023 - 06.30.2024): 90% + patient data captured *Maximum total 1 point extra credit per site	1
5	15	Site level PROS Collection: Completed primary Pre-op and 2-16 week post-op HOOS -JR or KOOS-JR + PROMIS10 (Overall average as of surgeries on or before 06.30.2024) When the difference between the PROS submission and completion rate at the site is >5%, the PROS completion rate will be used for this metric.	
	10	The site is awarded full points for collection rates of 70%+	15
		60%	10
		50%	5
		The site is not awarded points if collection is less than 50%	0
Extra Credit	0	Site level PROS Collection: Completed primary Pre-op and 1-year post-op HOOS -JR or KOOS-JR + PROMIS10 (Overall average as of surgeries on 06.01.2022 to 11.03.2023 and 1 year post op300-425 days post-op accepted.) When the difference between the PROS submission and completion rate at the site is >5%, the PROS completion rate will be used for this metric. The site is awarded full points for collection rates of 50%+	5 3
		30%	1
		<30%	0
6	10	Site level PROS Collection: Completed primary Pre-op HOOS -JR or KOOS-JR + PROMIS10 (Surgeries on from July 1, 2023 to June 30, 2024) When the difference between the PROS submission and completion rate at the site is >5%, the PROS completion rate will be used for this metric. The site is awarded full points for collection rates of	
		90%+	10
		75 - 89%	6
		60 - 74%	4
		The site is not awarded points if collection is less than 60%	0
7	15	% of Opioid naive THA patients at the SITE meet the MARCQI Pain control pathway guidelines (<240 OME)	

Ql Measurement Period: 07/01/2023 - 06/30/2024

Measure#	Max. Weight	Measure Description	Points
		85% or greater of THA patients meet the guidelines of 240 OME or less	15
		60-84% of THA patients prescribed <240 OME	7
		Less than 60% of patients meet the prescribing criteria	0
		% of Opioid naive TKA patients at the SITE meet the MARCQI Pain control pathway guidelines (<320 OME)	
8	15	90% or greater of TKA patients meet the guidelines of 320 OME or less	15
		70-89% of TKA patients prescribed <320 OME	7
		Less than 70% of patients meet the prescribing criteria	0
		90-Day Hip fracture: Reduce COLLABORATIVE rate of 90-Day Hip fracture for all primary HIP procedures (excluding conversions) by 10% (Baseline FY2023 = 1.17%; 10% reduction goal = 1.05%)	
9	5	1.05% or less of all primary HIPS experience a 90- Day hip fracture	5
		Greater than 1.05% but less than 1.17% of all primary HIPS experience a 90-Day hip fracture	2
		The site is not awarded points if there is no collaborative-wide improvement (>=1.17%) in all primary HIPS with 90-Day hip fractures	0
10	10	Implementation of one site specific quality initiative (linked to a MARCQI quality initiative). Using the April 2023 Quarterly reports, sites not performing equal to or better than their peers in PROS data collection, OME prescribing patterns, hip fractures, or 30-Day ED visits must choose one of those areas of focus as their project. If a site is performing better than their peers in PROS data collection, OME prescribing patterns, hip fractures, and 30-Day ED visits, they may choose a site specific quality initiative utilizing the April 2023 MARCQI Quarterly Report. Progress Reports are due in May 2024 & January 2025. Final results are based on quarterly reports of January, 2025 Plan submitted and approved, reporting requirements & timelines met, A3 submitted, and goal met* *Clinical Champion and Clinical Data Abstractor must be available to present at June 2025 Collaborative-wide session if asked *Not presenting if asked will yield -20 points on the FY2025 Performance Incentive scorecard	10

QI Measurement Period: 07/01/2023 - 06/30/2024

Measure#	Max. Weight	Measure Description	Points
		Plan submitted and approved Reporting requirements are met & timelines, but the target identified is not met. A3 submitted with final report and presentation by clinical champion and clinical data abstractor given at June 2025 MARCQI Collaborative-wide sessions* *Clinical Champion and Clinical Data Abstractor must be available to present at June 2025 Collaborative-wide session if asked *Not presenting if asked will yield -20 points on the FY2025	6
		Performance Incentive scorecard	
		Plan submitted and approved All reporting requirements are met, but the identified goal and timelines are not met.	3
		Plan is not developed, reports not done.	0
Extra Credit	0	Site based Quality Improvement: Alignment with MARCQI's 2024 demographic variables (e.g., AMA's recommendations for gender, race, ethnicity, etc.) If a site can document in their electronic health record that options for race, ethnicity, and gender are inclusive to MARCQI's 2024 definition (AMA recommendations) are available for MARCQI patients by December 31, 2023, extra credit points will be rewarded *Maximum total 2 point extra credit per site	2

2024 Michigan Arthroplasty Registry Collaborative Quality Initiative (MARCQI)

Performance Index Scorecard Extra Credit for Ambulatory Surgery Facility/Center (ASF/ASC) Participation Measurement Period: 01/01/2024 - 11/30/2024 QI Measurement Period: 07/01/2023-06/30/2024

Measure#	Max. Weight	Measure Description	Points	Eligible P4P Years
Extra Credit	0	Clinical Champion active engagement and participation in Quality Improvement working groups (e.g., PROS, ASC, Infection, Hip Fractures, Pain control, Device committee, Outlier work group, Quality Metrics work group, Executive Committee, Academic Quality Team) *Maximum total 0.5 point extra credit per site	0.5	2, 3+
Extra Credit	0	CDA active engagement and participation in CDA Committees or working groups (e.g., CDA Committee, Specifications Manual work group, PROS, ASC, Infection, Patient education, etc.) *Maximum total 0.5 point extra credit per site	0.5	2, 3+
Extra Credit	0	Site's with CDAs in their role for 9 months or greater as of January 31, 2024 who have at least one CDA attend the in-person CDA refresher training course. *Maximum total 0.5 point extra credit per site	0.5	2, 3+
Extra Credit	0	Collaborative Meeting Participation*- Site Administrator (01.01.2024- 11.30.2024) *Attendance at 3 of 3 Collaborative-wide meeting in February 2, 2024 (Virtual); June 7, 2024 (Grand Rapids, MI); and September 27, 2024 (Ann Arbor, MI) Individual must be identified and confirmed to fill this role by 1st 2024 MARCQI Collaborative-wide meeting *Maximum total 0.5 point extra credit per site	0.5	2, 3+
Extra Credit	0	Site level: On-time data entry score is greater than or equal to 99% during the measurement period (07.01.2023 - 06.30.2024) AND complete data entry score is greater than or equal to 99% (QA & CI review score average) *Maximum total 0.5 point extra credit per site	0.5	2, 3+

2024 Michigan Arthroplasty Registry Collaborative Quality Initiative (MARCQI)

Performance Index Scorecard Extra Credit for Ambulatory Surgery Facility/Center (ASF/ASC) Participation Measurement Period: 01/01/2024 - 11/30/2024 QI Measurement Period: 07/01/2023-06/30/2024

Measure#	Max. Weight	Measure Description	Points	Eligible P4P Years
Extra Credit	0	Access to Surgeon's Office Records (90 day events): (Surgery dates 07.01.2023 - 06.30.2024): 90% + patient data captured *Maximum total 1 point extra credit per site	1	2, 3+
Extra Credit	0	Site level PROS Collection: Completed primary Pre-op and 1-year post-op HOOS -JR or KOOS-JR + PROMIS10 (Overall average as of surgeries on 06.01.2022 to 11.03.2023 and 1 year post op300-425 days post-op accepted.) When the difference between the PROS submission and completion rate at the site is >5%, the PROS completion rate will be used for this metric.		3+
		The site is awarded full points for collection rates of 50%+ 40% 30%	5 3 1 0	
Extra Credit	0	Site based Quality Improvement: Alignment with MARCQI's 2024 demographic variables (e.g., AMA's recommendations for gender, race, ethnicity, etc.) If a site can document in their electronic health record that options for race, ethnicity, and gender are inclusive to MARCQI's 2024 definition (AMA recommendations) are available for MARCQI patients by December 31, 2023, extra credit points will be rewarded *Maximum total of 2 extra credit points per site	2	2, 3+
Extra Credit	0	Site based Quality Improvement: Alignment with MARCQI's 2024 demographic variables (e.g., AMA's recommendations for gender, race, ethnicity, etc.) If a site can document in their electronic health record that options for race, ethnicity, and gender are inclusive to	0.5	1

Extra Credit for Ambulatory Surgery Facility/Center (ASF/ASC) Participation Measurement Period: 01/01/2024 - 11/30/2024 QI Measurement Period: 07/01/2023-06/30/2024

Measure#	Max.	Measure Description	Points	Eligible P4P
inicadare ii	Weight	MARCQI's 2024 definition (AMA recommendations) are available for MARCQI patients by June 30, 2024, extra credit points will be rewarded *Maximum total of 0.5 extra credit points per site		Years
Extra credit	0	MODB data submission: Provide site's written planned process for site's submission of all MARCQI eligible cases data to Michigan Health and Hospital Association's Michigan Outpatient Database (MODB) by August 31, 2024 *Maximum total 0.5 point extra credit per site	0.5	1

2024 Michigan Bariatric Surgery Collaborative Quality Initiative (MBSC) Performance Index Scorecard Measurement period identified in each measure Measure# Weight Measure Description **Points** Improvement/Excellence In Grade 1 Complication Rate (Adjusted) **Measurement Periods:** Improvement: OR dates 10/1/2021 to 9/30/2024 Excellence: OR Dates 10/1/2023 to 9/30/2024 Major improvement (z-score less than -1 or Grade 1 10 1 complication rate ≤4%) 10 Moderate improvement/maintained complication rate (zscore between 0 and -1) 5 No improvement/rates of grade 1 complications increased 0 (z-score ≥0) Improvement/Excellence in Serious Complication Rate (Adjusted) **Measurement Periods:** Improvement: OR dates 10/1/2021 to 9/30/2024 Excellence: OR Dates 10/1/2023 to 9/30/2024 2 20 Major improvement (z-score less than -1 or serious complication rate ≤2.4%) 20 Moderate improvement/maintained complication rate (zscore between 0 and -1) 10 No improvement/rates of serious complications increased 0 (z-score ≥0) 1-Year Follow-up Rates (Adjusted) ≥67% OR > 5% **relative** improvement from previous year (10/1/2021-9/30/2022) 10 Maintained 1-year follow-up rate/ >0 to <5% relative 3 10 improvement from previous year (10/1/2021-9/30/2022) 5 1-year follow-up rate decreased/No improvement in 1-year follow-up rate (10/1/2021-9/30/2022) 0 Compliance with VTE prophylaxis - Pre-operatively and Post-operatively (Unadjusted) Measurement Period: 01/01/2024 - 12/31/2024 5 4 ≥94% compliance with guidelines 5 O 0 to 93% compliance with guidelines Compliance with VTE prophylaxis - Post Discharge (Unadjusted) Measurement Period: 01/01/2024 - 12/31/2024 5 5 ≥70% compliance with guidelines or a >2.5% relative improvement from the previous year (1/1/2023 to 12/31/2023) 5 0 0 to 69% compliance with guidelines Opioid Use - Opioid prescriptions within 30 days (measured in MMEs) (Unadjusted) Measurement Period: 10/01/2023 - 09/30/2024 6 5 <42 MME or > 10% relative reduction in opioid use 5 2.5 5-9% relative reduction in opioid use < 5% relative reduction 0

2024 Michigan Bariatric Surgery Collaborative Quality Initiative (MBSC) Performance Index Scorecard Measurement period identified in each measure Measure# Weight Measure Description **Points** Opioid Use - Opioid prescriptions within 30 days (measured in MMEs) (Unadjusted) Measurement Period: 10/01/2023 - 09/30/2024 7 10 <42 MME or > 10% relative reduction in opioid use 10 5-9% relative reduction in opioid use 5 0 < 5% relative reduction ED Visits (not resulting in a readmission, "avoidable") (Unadjusted) 8 5 Measurement Period: 10/01/2023 - 09/30/2024 5 < 6% Avoidable ED visits > 6% Avoidable ED visits 0 Meeting Attendance - Surgeon: (Calendar Year 2024) Attended 3 out of 3 meetings 5 9 5 3 Attended 2 out of 3 meetings 0 Attended fewer than 2 meetings Meeting Attendance - Abstractor/Coordinator: (Calendar Year 2024) Attended 3 out of 3 meetings 5 10 5 Attended 2 out of 3 meetings 3 0 Attended fewer than 2 meetings Timely Monthly Data Submissions (30-day information & registry paperwork) 01/01/2024 - 12/31/2024: On time 11-12 months 5 11 5 3 On time 10 months 0 On time 9 months or less Consent Rate (10/01/2023 - 09/30/2024) (Unadjusted): ≥90% consented patients 5 12 5 3 80-89% consented patients 0 <80% consented patients 10 Physician Engagement (01/01/2024 – 12/31/2024): Following items count as 1 activity point: ** Note: For each site, a surgeon or surgeons must participate in at least 2 of the engagement activities listed below in order to receive the 10 points available for this measure.** ***In order for a surgeon to earn points for a hospital, they must complete 10 bariatric surgery cases at that hospital for the dates of 1/1/2024 to 12/31/2024 13 10 Committee participation MBSC survey response Participate in a qualitative interview Coauthor a paper Participate in quality improvement initiatives (MPIRRE/FUTURE/MSHEILD/etc.)

2	2024 Michigan Bariatric Surgery Collaborative Quality Initiative (MBSC) Performance Index Scorecard Measurement period identified in each measure					
Measure#	Weight	Measure Description	Points			
		Attend or present at a pre-meeting session (IH committee/surgeon skill/etc.) on the day of the MBSC triannual meeting				
		Present MBSC data at a MBSC tri-annual meeting				
		Attend quality site visit as a guest surgeon				
		Following items count as 2 activity points:				
		Host quality site visit				
		Present MBSC data at a national meeting				
		Lead author on an MBSC publication				
		No participation	0			

2024 Michigan Emergency Department Improvement Collaborative (MEDIC) Quality Initiative Performance Index Scorecard Measurement Period: 11/1/2023 – 10/31/2024 Year 1 Sites

Data Delivery: Timeliness	Measure#	Weight	Measure Description	Points	
1	1		Data Delivery: Timeliness		
9-10 months of data transfers on time 4			All 12 months of data transfers on time	10	
Successful submission within 1 month of deadline Successful submission within 1 month of deadline Submission more than 1 month after deadline Submission hathin 1 month of deadline Submission deadline Submissi		10	11 months of data transfers on time	8	
Data Delivery: Adherence & Accuracy			9-10 months of data transfers on time	4	
All 12 months of data transfers adhere to MEDIC data dictionary and are accurate			< 9 months of data transfers on time	0	
10					
10				10	
10					
9-10 months of data transfers adhere to MEDIC data dictionary and are accurate 4	2	10		8	
Successful submission by deadline Successful submission by deadline Successful submission by deadline Successful submission within 1 month of deadline Submission more than 1 month after deadline Outline	_		•	4	
Blectronic Data Dictionary Update				4	
Blectronic Data Dictionary Update				0	
Successful submission by deadline 6			·	_	
Successful submission within 1 month of deadline Submission more than 1 month after deadline O			<u> </u>	- 6	
Submission more than 1 month after deadline 0	3	6	·		
Abstraction: Timeliness					
All cases abstracted by quarterly deadline 13 1 deadline missed 8 2+ deadlines missed 0				0	
13		13			12
2+ deadlines missed 0	4				
Meeting Attendance: Clinical Champion					
12				U	
Miss 1 Meeting 6			·	12	
Meeting Attendance: Data Abstractor	5	12			
Meeting Attendance: Data Abstractor					
Attend All Meetings 12 Miss 1 Meeting 6 Miss >1 Meeting 0 Time from Agreement being signed to hiring date of data abstractor 7 14 < 90 days 14 91-120 days 8 >120 days 14 Time from Agreements signed to successful submission of electronic production data 8 14 < 90 days 14 91-120 days 8			,		
Miss 1 Meeting 6 Miss > 1 Meeting 0				12	
Time from Agreement being signed to hiring date of data abstractor 7 14	6	12			
Time from Agreement being signed to hiring date of data abstractor 7 14 			· · · · · · · · · · · · · · · · · · ·		
7 14			_		
91-120 days 8 >120 days 0 Time from Agreements signed to successful submission of electronic production data 8 14 <90 days 14 91-120 days 8					
Time from Agreements signed to successful submission of electronic production data 14 <90 days 14 91-120 days 8	7	14	•	14	
Time from Agreements signed to successful submission of electronic production data 14 <90 days 14 91-120 days 8			·	8	
8 14 submission of electronic production data <90 days 14 91-120 days 8			>120 days	0	
8 14 < 90 days 14 91-120 days 8					
91-120 days 8	8	14	·	14	
			>120 days	0	

2024 Michigan Emergency Department Improvement Collaborative (MEDIC) Quality Initiative Performance Index Scorecard Measurement Period: 11/1/2023 – 10/31/2024 Year 1 Sites

Measure#	Weight	Measure Description	Points
9	9	Intervention Planning for Year 2 (Intervention Templates, etc.)	
		All Year 2 materials complete and submitted on time	9
		Year 2 materials incomplete and/or submitted late	0

2024 Michigan Emergency Department Improvement Collaborative (MEDIC) Quality Initiative Performance Index Scorecard Measurement Period: 11/1/2023 – 10/31/2024 Year 2 Sites

Measure#	Weight	Measure Description	Points
		Data Delivery: Timeliness	
		All 12 months of data transfers on time	12
1	12	11 months of data transfers on time	9
		9-10 months of data transfers on time	6
		< 9 months of data transfers on time	0
		Data Delivery: Adherence & Accuracy	
		All 12 months of data transfers adhere to MEDIC data dictionary and are accurate	12
2	12	11 months of data transfers adhere to MEDIC data dictionary and are accurate	9
		9-10 months of data transfers adhere to MEDIC data dictionary and are accurate	6
		< 9 months of data transfers adhere to MEDIC data dictionary and are accurate	0
		Electronic Data Dictionary Update	
3	6	Successful submission by deadline	6
		Successful submission within 1 month of deadline	3
		Submission more than 1 month after deadline	0
		Abstraction: Timeliness	
4	10	All cases abstracted by quarterly deadline	10
		1 deadline missed	5
		2+ deadlines missed	0
		Meeting Attendance: Clinical Champion	
5	10	Attend All Meetings	10
		Miss 1 Meeting	5
		Miss >1 Meeting	0
		Meeting Attendance: Data Abstractor	
6	10	Attend All Meetings	10
		Miss 1 Meeting	5
		Miss >1 Meeting	0
7	10	Annual Abstraction Audit: SNAP (<u>S</u> haring K <u>n</u> owledge <u>A</u> nd <u>P</u> erspectives) Review	
7	10	≥ 90% of case cohort decisions are correct	4
		≥ 75% of case cohort decisions are correct	2

2024 Michigan Emergency Department Improvement Collaborative (MEDIC) Quality Initiative Performance Index Scorecard Measurement Period: 11/1/2023 – 10/31/2024 Year 2 Sites

Measure#	Weight	Measure Description	Points
		< 75% of case cohort decisions are correct	0
		≥ 97% of abstracted registry data accurate	6
		95%-97% of abstracted registry data accurate	3
		<95% of abstracted registry data accurate	0
		Site Specific - Timely Administration of Steroids in Pediatric Asthma *Measures and targets identified in Appendix	
0-	40	QI Project developed and implemented and site met or exceeded target	10
8a	10	QI Project developed and implemented and site made improvement toward, but did not meet, the target	7
		QI Project developed and implemented but there was no improvement to the target	4
		QI Project not developed or implemented	0
	10	Site Specific - Adult Low Risk Chest Pain Safe Discharge Initiative *Measures and targets identified in Appendix	
OI.		QI Project developed and implemented and site met or exceeded target	10
8b		QI Project developed and implemented and site made improvement toward, but did not meet, the target	7
		QI Project developed and implemented but there was no improvement to the target	4
		QI Project not developed or implemented	0
	40	Collaborative-Wide Measure: Naloxone Distribution for Opioid Use Harm Reduction	
9	10	Met Naloxone Distribution Target	10
		Did meet target	0
10	_	Collaborative-Wide Measure: Peds Chest Xray Utilization	
10	5	Met CXR Utilization Target	5
		Did meet target	0
4.4	5	Collaborative-Wide Measure: Adult Suspected PE CT Diagnostic Yield	
11) 	Met Diagnostic Yield Target	5
		Did meet target	0

2024 Michigan Emergency Department Improvement Collaborative (MEDIC) Quality Initiative Performance Index Scorecard Measurement Period: 11/1/2023 – 10/31/2024 Year 3+ Sites

Measure#	Weight	Measure Description	Points
4		Data Delivery: Timeliness	
	4	All 12 months of data transfers on time	4
1	4	11 months of data transfers on time	3
		9-10 months of data transfers on time	2
		< 9 months of data transfers on time	0
		Data Delivery: Adherence & Accuracy	
		All 12 months of data transfers adhere to MEDIC data dictionary and are accurate	4
2	4	11 months of data transfers adhere to MEDIC data dictionary and are accurate	3
		9-10 months of data transfers adhere to MEDIC data dictionary and are accurate	2
		< 9 months of data transfers adhere to MEDIC data dictionary and are accurate	0
		Electronic Data Dictionary Update	
3	2	Successful submission by deadline	2
		Successful submission within 1 month of deadline	1
		Submission more than 1 month after deadline	0
		Abstraction: Timeliness	
4	5	All cases abstracted by quarterly deadline	5
		1 deadline missed	3
		2+ deadlines missed	0
		Meeting Attendance: Clinical Champion	
5	5	Attend All Meetings	5
		Miss 1 Meeting	3
		Miss >1 Meeting	0
		Meeting Attendance: Data Abstractor	
6	5	Attend All Meetings	5
		Miss 1 Meeting	3
		Miss >1 Meeting	0
7		Annual Abstraction Audit: SNAP (<u>S</u> haring K <u>n</u> owledge <u>A</u> nd <u>P</u> erspectives) Review	
	5	≥ 90% of case cohort decisions are correct	2
'	5	≥ 75% of case cohort decisions are correct	1
		< 75% of case cohort decisions are correct	0
		≥ 97% of abstracted registry data accurate	3

2024 Michigan Emergency Department Improvement Collaborative (MEDIC) Quality Initiative Performance Index Scorecard Measurement Period: 11/1/2023 – 10/31/2024 Year 3+ Sites

Measure#	Weight	Measure Description	Points
		95%-97% of abstracted registry data accurate	2
		<95% of abstracted registry data accurate	0
	25	Site Specific - Timely Administration of Steroids in Pediatric Asthma *Measures and targets identified in Appendix	
8a		QI Project developed and implemented and site met or exceeded target	25
. Ja		QI Project developed and implemented and site made improvement toward, but did not meet, the target	15
		QI Project developed and implemented but there was no improvement to the target	10
		QI Project not developed or implemented	0
		Site Specific - Adult Low Risk Chest Pain Safe Discharge Initiative *Measures and targets identified in Appendix	
Oh	25	QI Project developed and implemented and site met or exceeded target	25
8b		QI Project developed and implemented and site made improvement toward, but did not meet, the target	15
		QI Project developed and implemented but there was no improvement to the target	10
		QI Project not developed or implemented	0
9	10	Collaborative-Wide Measure: Naloxone Distribution for Opioid Use Harm Reduction	
9		Met Naloxone Distribution Target	10
		Did meet target	0
10	5	Collaborative-Wide Measure: Peds Chest Xray Utilization	
10	5	Met CXR Utilization Target	5
		Did meet target	0
11	5	Collaborative-Wide Measure: Adult Suspected PE CT Diagnostic Yield	
		Met Diagnostic Yield Target	5
		Did meet target	0
12a	25	Site Specific - Quality Improvement Initiative: Adult Head Injury CT Appropriateness	
		QI Project developed and implemented and site met or exceeded target	25
		QI Project developed and implemented and site made improvement toward, but did not meet, the target	15
		QI Project developed and implemented but there was no improvement to the target	10

2024 Michigan Emergency Department Improvement Collaborative (MEDIC) Quality Initiative Performance Index Scorecard Measurement Period: 11/1/2023 – 10/31/2024 Year 3+ Sites

Measure#	Weight	Measure Description	Points
		QI Project not developed or implemented	0
12b		Site Specific - Quality Improvement Initiative: Pediatric Intermediate Risk Head Injury CT Utilization	
		QI Project developed and implemented and site met or exceeded target	25
	25	QI Project developed and implemented and site made improvement toward, but did not meet, the target	15
		QI Project developed and implemented but there was no improvement to the target	10
		QI Project not developed or implemented	0

2024 Michigan Radiation Oncology Quality Consortium (MROQC) Quality Initiative Performance Index Scorecard Measurement Period: 01/01/2024-09/30/2024

Contours and dose reported in 50-69% of patients Contours and dose reported in <50% of patients For lung cancer patients treated with conventional fractionation, the mean esophageal dose is <29 Gy AND the esophageal max dose (D2cc) is <61 Gy. 10 ≥65% of lung patients met both constraints 50-64% of lung patients met both constraints <50% of lung patients met both constraints For SBRT treatment of lung cancer with a single PTV, the Paddick Conformity Index is ≥0.85. ≥80% of patients treated for lung cancer with a single PTV with SBRT met the criterion 60-79% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion (Collaborative-Wide Goal Metric A ONLY) Use of shorter course radiotherapy for bone	Measure#	Weight	Measure Description	Points
Three Metrics Met Two Metrics Met One Metric Met None Met Timely Submission of High-Quality Physics & Dosimetry Data Three Metrics Met Two Metrics Met Two Metrics Met Three Metrics Met One Metric Met None Met In node-positive breast cancer patients, the irradiated nodal group(s) is(are) contoured and named per TG- 263 naming convention AND the dose to the SCV, ICV (Axillary Level 3), Axilla (Level 1 & 2), and/or IMN is reported. Contours and dose reported in ≥70% of patients Contours and dose reported in 50% of patients Contours and dose reported in <50% of patients For lung cancer patients treated with conventional fractionation, the mean esophageal dose is <29 Gy AND the esophageal max dose (D2cc) is <61 Gy. ≥65% of lung patients met both constraints 50-64% of lung patients met both constraints <p>For SBRT treatment of lung cancer with a single PTV, the Paddick Conformity Index is ≥0.85. ≥80% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion</p> <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion		6		
Timely Submission of High-Quality Physics & Dosimetry Data Three Metrics Met Timely Submission of High-Quality Physics & Dosimetry Data Three Metrics Met Two Metrics Met One Metric Met None Met In node-positive breast cancer patients, the irradiated nodal group(s) is(are) contoured and named per TG-263 naming convention AND the dose to the SCV, ICV (Axillary Level 3), Axilla (Level 1 & 2), and/or IMN is reported. Contours and dose reported in ≥70% of patients Contours and dose reported in <50% of patients Contours and dose reported in <50% of patients For lung cancer patients treated with conventional fractionation, the mean esophageal dose is <29 Gy AND the esophageal max dose (D2cc) is <61 Gy. 4 10 ≥65% of lung patients met both constraints 50-64% of lung patients met both constraints <p>≤50% of lung patients met both constraints For SBRT treatment of lung cancer with a single PTV, the Paddick Conformity Index is ≥0.85. ≥80% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion</p> <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion				6
Timely Submission of High-Quality Physics & Dosimetry Data Three Metrics Met Trow Metrics Met One Metric Met None Met In node-positive breast cancer patients, the irradiated nodal group(s) is(are) contoured and named per TG-263 naming convention AND the dose to the SCV, ICV (Axillary Level 3), Axilla (Level 1 & 2), and/or IMN is reported. Contours and dose reported in ≥70% of patients Contours and dose reported in 50-69% of patients Contours and dose reported in <50% of patients Contours and dose reported in <50% of patients For lung cancer patients treated with conventional fractionation, the mean esophageal dose is <29 Gy AND the esophageal max dose (D2cc) is <61 Gy. 10 ≥65% of lung patients met both constraints 50-64% of lung patients met both constraints <p>For SBRT treatment of lung cancer with a single PTV, the Paddick Conformity Index is ≥0.85. ≥80% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion</p> <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion	1		Two Metrics Met	4
Timely Submission of High-Quality Physics & Dosimetry Data Three Metrics Met Two Metrics Met One Metric Met None Met In node-positive breast cancer patients, the irradiated nodal group(s) is(are) contoured and named per TG-263 naming convention AND the dose to the SCV, ICV (Axillary Level 3), Axilla (Level 1 & 2), and/or IMN is reported. Contours and dose reported in ≥70% of patients Contours and dose reported in 50-69% of patients Contours and dose reported in 50-69% of patients Contours and dose reported in 50-69% of patients For lung cancer patients treated with conventional fractionation, the mean esophageal dose is <29 Gy AND the esophageal max dose (D2cc) is <61 Gy. 4 10 ≥65% of lung patients met both constraints 50-64% of lung patients met both constraints <p>50-64% of lung patients met both constraints For SBRT treatment of lung cancer with a single PTV, the Paddick Conformity Index is ≥0.85. ≥80% of patients treated for lung cancer with a single PTV with SBRT met the criterion 60-79% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion</p> <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion			One Metric Met	2
Three Metrics Met Two Metrics Met One Metric Met None Met In node-positive breast cancer patients, the irradiated nodal group(s) is(are) contoured and named per TG-263 naming convention AND the dose to the SCV, ICV (Axillary Level 3), Axilla (Level 1 & 2), and/or IMN is reported. Contours and dose reported in ≥70% of patients Contours and dose reported in 50-69% of patients Contours and dose reported in <50% of patients Contours and dose reported in <50% of patients For lung cancer patients treated with conventional fractionation, the mean esophageal dose is <29 Gy AND the esophageal max dose (D2cc) is <61 Gy. 10 ≥65% of lung patients met both constraints <p>50-64% of lung patients met both constraints 50% of lung patients met both constraints For SBRT treatment of lung cancer with a single PTV, the Paddick Conformity Index is ≥0.85. ≥80% of patients treated for lung cancer with a single PTV with SBRT met the criterion 60-79% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion</p> <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion			None Met	0
Two Metrics Met One Metric Met None Met In node-positive breast cancer patients, the irradiated nodal group(s) is(are) contoured and named per TG-263 naming convention AND the dose to the SCV, ICV (Axillary Level 3), Axilla (Level 1 & 2), and/or IMN is reported. Contours and dose reported in ≥70% of patients Contours and dose reported in 50-69% of patients Contours and dose reported in <50% of patients Contours and dose reported in <50% of patients For lung cancer patients treated with conventional fractionation, the mean esophageal dose is <29 Gy AND the esophageal max dose (D2cc) is <61 Gy. ≥65% of lung patients met both constraints 50-64% of lung patients met both constraints <p><50% of lung patients met both constraints</p> For SBRT treatment of lung cancer with a single PTV, the Paddick Conformity Index is ≥0.85. ≥80% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion <60% of patients treated for lung cancer with a single PTV with SBRT met the criterion				
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with SBRT met the criterion (Collaborative-Wide Goal Metric A ONLY) Use of shorter course radiotherapy for bone	3		PTV with SBRT met the criterion	7
Use of shorter course radiotherapy for bone			with SBRT met the criterion	0
A: The MROQC consortium-wide rate of single fraction use is ≥45% for uncomplicated patients B: Your site-level rate of ≤5 fraction treatment is at least 75% for all patients	6	10	Use of shorter course radiotherapy for bone metastasis treatment as shown by: A: The MROQC consortium-wide rate of single fraction use is ≥45% for uncomplicated patients B: Your site-level rate of ≤5 fraction treatment is at least 75% for all patients	10

2024 Michigan Radiation Oncology Quality Consortium (MROQC) Quality Initiative Performance Index Scorecard Measurement Period: 01/01/2024-09/30/2024

Measure#	Weight	Measure Description	Points
		Only B is met	8
		B is not met	0
7	10	For treatment of bone metastasis using stereotactic body radiotherapy (SBRT): A. Standardized dose constraints for organs at risk (OARs) are used B. Any violations of the standardized dose constraints are documented. ≥80% of patients who received bone metastasis treatment with SBRT met both criteria <80% of patients who received bone metastasis treatment	10
8	10	Percentage of patients with intermediate risk prostate cancer as defined by NCCN treated with EBRT or brachytherapy who received "high value radiotherapy", defined as moderately hypofractionated EBRT (28 fractions or less) OR ultrahypofractionated EBRT/SBRT (7 fractions or less) OR brachytherapy monotherapy. Patients with unfavorable intermediate risk prostate cancer may also receive a brachytherapy boost.	U
		≥70% of patients received high value radiotherapy	10
		50-69% of patients received high value radiotherapy	7
		<50% of patients received high value radiotherapy	0
		Prostate Working Group Performance Goal: Completion of 12-month follow-up form (P6)	
9	10	≥60% of prostate 12-month follow-up forms completed	10
		40-59% of prostate 12-month follow-up forms completed	7
		<40% of prostate 12-month follow-up forms completed	0
	6	Collaborative-Wide Meeting Participation – Clinical Champion (per MROQC CC Attendance Policy)*	
10		All meetings or two meetings with one meeting attended by an acceptable designee	6
		Two meetings only	4
		One meeting or none attended	0
11	6	Collaborative-Wide Meeting Participation – Physics Lead (or designee)	
		All meetings	6
		Two meetings	4
		One meeting or none attended	0
12	6	Collaborative-Wide Meeting Participation – Clinical Data Abstractor (or designee)	

2024 Michigan Radiation Oncology Quality Consortium (MROQC) Quality Initiative Performance Index Scorecard Measurement Period: 01/01/2024-09/30/2024

Measure#	Weight	Measure Description	Points
	I	A11 (*)	
		All meetings	6
		Two meetings	4
		One meeting or none attended	0
	100		
		MROQC Physician Engagement	
Bonus	10	 Lead author on an MROQC publication (Counts as 2 items) Lead a skills workshop (Counts as 2 items) Present at an MROQC collaborative-wide meeting (Non-leadership role only) Present on MROQC at a national meeting (Cannot be a resident) Attend 5 working group meetings in 2024 (Total across practice physicians; 1 physician counts per meeting (i.e., no double points if 2 attend the same meeting) Coauthor on an MROQC publication Participate in 3 case review sessions Propose a new quality measure: provide reasoning to implement the measure, work with the MROQC data team to review supporting data and present the measure to the WG. 	
		5 or more items achieved	10
		3-4 items achieved	5
		1-2 items achieved	1

202	4 Michiga	n Surgical Quality Collaborative (MSQC) Performance Index Scorecard Project Time Period: 1/1/2024 – 12/31/2024	
Measure#	Weight	Measure Description	Points
		Collaborative Meetings (4 offered) – Surgical Clinical Quality Reviewer (SCQR)	
1	6	3 or more meetings	6
		2 meetings	3
		1 meeting	1
	6	Collaborative Meetings (3 offered) – Surgeon Champion (SC)	
2		2 or more meetings	6
		1 meeting	3
		0 meetings	0
		Conference Calls (3 offered) – SCQR	
3	4	2 or more calls	4
		1 call	2
		0 calls	0
	4	SCQR Participation/Engagement Participated in at least one MSQC activity listed in the	
4		supplement document.	4
4		No Contribution: Did not participate in any activities listed in	
		the supplement document.	0
		SC Participation/Engagement	
		Participated in at least one MSQC activity listed in the	
5	4	supplement document.	4
		No Contribution: Did not participate in any activities listed in	0
		the supplement document.	U
		Completeness of Data (maximum 6 pts available)	
		Sampled and incomplete cases ≤ 0.5% total volume	1
		Case Selection Audit with ≥ 95% agreement	1
		30 day follow-up rate ≥ 80% for 4th quarter 2023 (October	1
6	6	- December cases)	
6	6	30 day follow-up rate ≥ 80% for 1st quarter 2024 (Jan – March cases)	1
		30 day follow-up rate ≥ 80% for 2nd quarter 2024 (April –	
		June cases)	1
		30 day follow-up rate ≥ 80% for 3rd quarter 2024 (July –	_
		September cases)	1
7		Collaborative Wide Measure*: Preop Optimization for	
		elective abdominal hernia surgery:	
	20	• Reduce rate of persons with body mass index (BMI) ≥	
		40kg/m2 undergoing elective surgery to < 10%.	
		Reduce rate of persons with active tobacco use undergoing elective surgery to < 15%	
		undergoing elective surgery to < 15%.	
		Most both massives	20
		Meet both measures Meet one measure	20 10
		No measures met	0
		ino measures met	U

2024 Michigan Surgical Quality Collaborative (MSQC) Performance Index Scorecard Project Time Period: 1/1/2024 – 12/31/2024			
Measure#	Weight	Measure Description	Points
8	5	Complete documentation of designated cancer variables (CRC, Breast, Whipple, Thyroid)	
		90 - 100%	5
		< 90%	0
9	40	Quality Improvement Initiative (QII) - choose from one of the following: Option A: SUCCESS OR Option B: Frailty OR Option C: Breast Surgery OR Option D: Preoperative Testing (by invitation only)	40
10	5	Submit project report detailing local processes and structures and outcomes for improving adherence to Collaborative Wide Measure	5
		Bonus points to be added [^] to reflect active participation in MOQC over-sampling of hysterectomy cases. Points available when:	
Optional	5	Site fully participates by over-sampling and abstracting all gyn-onc hysterectomy cases	5
		Site partially participates by over-sampling gyn-onc hysterectomy cases only	2 100
Total Available Points			

[^]Earned bonus points will be added to Scorecard total, with final score not to exceed 100 points overall. See MSQC/MOQC Cross-Collaborative Project Summary document for details.

^{*}These goals may be updated at the end of 2023 once more data is available. Revised: 11/17/2023



Quality Improvement Implementation, Option A: SUCCESS 2024 Project Time Period: 1/1/2024 – 12/31/2024

Background - Although infections related to urinary catheters have received a great deal of attention due to public reporting and hospital penalties, non-infectious complications of bladder catheters are also a serious concern. These include trauma from catheter placement and/or removal, which is as common as urinary tract infections. Furthermore, surgeons' feedback revealed that the most common catheter-related problem seen in their practices is urinary retention, for which there is a lack of standardized management.

Project Goal and Summary – The 2024 SUCCESS project will build on the work begun in 2023. In collaboration with the Surgical Champion and the multidisciplinary team, this intervention aims to (1) reduce inappropriate perioperative urinary catheter use, (2) reduce catheter-associated trauma, and (3) improve the management of postoperative urinary retention. The project focuses on four common general surgery procedures: appendectomy, cholecystectomy, colorectal surgery, and hernia repair. This project will include the implementation and/or evaluation of toolkit elements that will address clinician knowledge and urinary catheterization skills, as well as communication and implementation challenges anticipated to affect catheter use in different types of perioperative clinical settings. Continuing sites will continue the implementation of the toolkit elements, address barriers, meet process/outcome measures, and refine the care pathway.

Eligibility – Continuing sites that participated in the 2023 SUCCESS QI project and new sites that capture data in the MSQC SUCCESS tab starting from Cycle 35, 2023 are eligible to select this project as their 2024 QI Project. There is required variable training for the SCQRs new to the project.

QI Implementation Goals and Requirements: (40 points total)

- 1. Capture all <u>SUCCESS data in MSQC Workstation</u> for eligible cases (3 points)
- 2. <u>Multidisciplinary team</u>: Participating hospitals will establish a multidisciplinary team to review data, guide quality improvement and toolkit element implementation plans, and implement the MSQC SUCCESS urinary care pathway (6 points total)
 - a. Suggested participants include surgeon leadership/surgeon champion, surgeons/residents (general & urology), executive leadership, anesthesiology, nursing supervisors for ER, Perioperative, PACU, and surgical units, quality department manager, patient safety, nursing education, and patient experience officer.
 - b. Hold three (3) <u>multidisciplinary meetings</u>. Submit minutes and attendees to the coordinating center with your 2024 Project Summary.
 - i. Kickoff meeting by March 29, 2024, to review project requirements and preliminary data. The SUCCESS Value Proposition/Leadership Engagement Briefing should be utilized during this meeting. New sites will complete the Readiness Assessment: Urinary Catheter Care Guide to Patient Safety (GPS) with the team (2 points).
 - ii. Two (2) additional multidisciplinary meetings (minimally) before December 1, 2024, which include a review of SUCCESS data and a quality review for cases in goal 6 (2 points each).

3. New sites only: Implement the elements of the <u>SUCCESS toolkit</u>. Submit a narrative of how the following toolkit elements were implemented/utilized (15 points).

Continuing sites only: Meet the below <u>process/outcomes measures</u> for appropriate catheter use and urinary retention diagnosis and management (15 points total, 3 points each measure). Measurement period 1/1/2024- 12/31/2024 OR dates:

- Catheter use measures
 - a. Avoid indwelling catheters intraoperatively in Category A*: < 10% of cases have an indwelling catheter
 - b. Catheter is removed in OR for > 75% of Category B* cases
- Urinary retention diagnosis and management measures
 - c. Bladder scan volume is documented \geq 90% of the time if urinary catheterization for retention was performed
 - d. No urinary catheter is used for bladder scan volumes < 300 ml for > 90% of cases
 - e. ISC was performed as opposed to an indwelling catheter (unless volume ≥ 500) for ≥ 90% of cases
- 4. With the multidisciplinary team, develop (new sites) or refine (continuing sites) the MSQC SUCCESS urinary care pathway template for your hospital's practices. This will be implemented and utilized by the care team to ensure the use of each element of the SUCCESS toolkit. Submit the final product to MSQC with your 2024 Project Summary (8 points).
 - a. Continuing sites only: Include a narrative of how any processes and toolkit elements were modified from 2023. Also include in the modified care pathway the process of coudé catheter training and comfort of use by nurses, and how patients at higher risk for urinary catheter trauma are identified.
- 5. Perform a <u>quality review</u> of each case that meets any of the criteria below, new sites from 4/1/2024 to 12/1/2024 OR dates, continuing sites from 1/1/2024 to 12/1/2024 OR dates. An overall findings summary (trends identified, action plans implemented) should be submitted with your 2024 Project Summary (8 points).
 - a. Patients in Category A who have an indwelling urinary catheter placed in the OR.
 - b. Retention is assigned for patients who had a urinary catheter (ISC or indwelling) placed when < 300 ml is documented via a bladder scanner or the catheter use
 - c. Patients who return to ED with Retention
 - d. Patients who were discharged with an indwelling catheter or need for ISC
 - e. Patients who have Urinary Catheter-Related Trauma assigned
- 6. Submit the 2024 SUCCESS Project Summary to the MSQC Coordinating Center no later than January 15, 2025. An additional 0-10 implementation points may be granted based on the detail of the project narrative, tracking log, and analysis; to be added to achieve the maximum of 40 project points.

* Category Definitions:

Category A: Avoid Placement: Avoid placing indwelling urinary catheter for these procedures: inappropriate to use a catheter or risks outweigh benefits (includes lap chole, lap/open appy, open groin hernia repair) Category B: Remove in OR: Consider removing indwelling urinary catheter before leaving the operating room (includes open/lap abdominal hemicolectomy, open/lap transanal rectal tumor excision, open/lap enterectomy, ostomy, MIS groin/ventral hernia repair, open ventral hernia repair <3 hrs) ventral hernia repair >3 hrs)



Quality Improvement Implementation, Option B: Frailty Project Time Period: 1/1/2024 – 12/31/2024

Background: Frailty develops through an accumulation of deficits over time that place patients at increased risk of suboptimal postoperative outcomes. Screening for frailty and discussion of the risks and benefits of moving forward with a surgical or non-surgical treatment pathway have been shown to improve both objective outcomes and patient satisfaction.

Project Goal and Summary: The 2024 frailty project will build on the work begun in 2023 which introduced a system to assess vulnerable patients for frailty using a validated screening tool with an informed discussion regarding the results of frailty screening during the surgical planning phase.

For 2024, the goal is to improve the surgical experience for patients identified as frail through interventions shown to potentially improve outcomes. This year will see an enhanced informed decision-making discussion with surgical candidates and caregivers and add interventions aimed at improving the surgical experience for candidates identified as frail. These interventions will include establishing patient overarching health goals and documenting this in the medical record, developing educational materials for patients and caregivers regarding care and expectations, and involvement of support persons throughout the surgical encounter to support those patients identified as frail.

QI Implementation Goals and Requirements (40 points total)

- Continuing sites: Measurement period is 1/1/2024-12/31/2024 OR dates
- New sites: Measurement period is 4/1/2024 to 12/31/2024 OR dates
- 1. Sites selecting the frailty pathway will collect MSQC data in the Frailty tab for eligible cases.
- 2. Hold three (3) multidisciplinary meetings. Submit minutes and attendees to the coordinating center with the final project submission (6 points):
 - a. Kickoff meeting by March 29, 2024, to review project requirements and preliminary data. (2 points).
 - b. Two (2) additional multidisciplinary meetings (minimally) before December 1, 2024, which include a review of data (2 points each).
- 3. Use of Frailty Tool for preoperative frailty screening in ≥ 75% of eligible patients (10 points). Sites will distribute and implement a frailty screening tool that will be completed during the preoperative planning period and be completed prior to the final decision to proceed with surgery on all elective surgery* patients meeting any of the following criteria:
 - Age ≥ 60 years on the day of surgery
 - Current dialysis as defined in MSQC core variable definitions.
 - Current cancer as defined in MSQC core variable definitions.
 - Functional health status identified as "Not Independent" in MSQC core variable definitions
 - Current CHF as defined in MSQC core variable definitions.

If frailty screening is not completed for a patient who qualifies:

- The SCQR will discern the rationale for failure to use the frailty tool.
- If this information is unavailable in the chart, the SCQR may also obtain this information by communication with office staff.

*A subset of participating surgeons may be identified if system-wide adoption is not feasible. This subset is to be defined using physician organization or affiliation. If a subset is to be used, a list of physicians who will be included in the data will be submitted when the quality improvement project declaration is provided to the coordinating center. All elective MSQC surgical cases performed by surgeons affiliated with the selected practice(s) will be assessed for use of the frailty screening tool.

4. A conversation between surgeon and patient/caregivers occurs for ≥ 75% of patients who screened as frail or pre-frail (7 points). The discussion will occur during the surgical planning process by a surgeon or an appropriate healthcare professional designee acting on behalf of the surgeon. This guided discussion and an attestation statement in the medical record are required and ensure that a conversation of the risks and benefits has taken place with the patient and/or caregivers.

The conversation and attestation statement regarding frailty screening, goals, and surgical plan must include the following elements:

- 1. General information on frailty
- 2. Interpretation of the scoring results (pre-frail or frail) according to the frailty screening
- 3. Discussion of the potential impact that frailty can have on surgical outcomes
- 4. The participants in the discussion, i.e., patient, family, caregiver
- 5. Any adjustments decided on in the plan of care (if applicable)

Appropriate designees for this conversation are defined as:

- An advanced practice provider (Nurse Practitioner or Physician Assistant) working in collaboration with the surgical team within the clinic setting.
- A provider (MD, DO, NP, CRNA, or PA) acting in partnership with the surgical team to provide preoperative screening, evaluation, or treatment that may occur outside the surgical clinic, on a day prior to the surgical date to facilitate surgical planning.
- 5. Patient/caregiver goals for surgery are documented for ≥ 75% of patients who screened as frail or pre-frail (7 points).
- 6. Provide preoperatively patient and/or caregiver education (10 points) *measurement period 4/1/2024-12/31/2024 to ≥ 75% of patients who screen frail or pre-frail (positive for frailty). The education may be completed by the surgeon or an appropriate healthcare professional designee which may include a Registered Nurse or Advanced Practice provider and must
 - be provided verbally and in writing regarding the impact of frailty on surgical outcomes.
 - include how increased risk could be mediated pre- and postoperatively.

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Submit the educational materials that are being used to the MSQC coordinating center for approval by February 15, 2024.

7. Submit the 2024 Frailty Project Summary to the MSQC Coordinating Center no later than January 15, 2025. An additional 0-10 implementation points may be granted based on the detail of the project narrative, tracking log, and analysis; to be added to achieve the maximum of 40 project points.



Quality Improvement Implementation, Option C: Breast Surgical Quality Measures Project Time Period: 1/1/2024-12/31/2024

Summary: This project focuses on improving the performance of evidence-based quality measures for patients undergoing partial mastectomy and mastectomy. MSQC began capturing breast surgery data in 2023, so this project will lay the groundwork and encourage engagement from the multidisciplinary team to promote high-quality treatment to improve short- and long-term outcomes.

Project Goals: Each site will designate a surgeon lead who performs breast surgery for this project who, along with the multidisciplinary team, will help disseminate information at the hospital, and be actively engaged in developing and implementing a care pathway that includes the perioperative QI implementation goals.

QI Implementation Goals and Requirements: (40 points total)

- 1. Data collection: For elective partial mastectomy and mastectomy surgical patients, participating hospitals will perform supplemental data collection that will allow the measurement of breast surgical quality.
- 2. Surgeon Champion: Each site will designate a surgeon champion who performs breast surgery to lead this project. The expectations are the surgeon lead will help the SCQR disseminate information at the hospital, be actively engaged in developing and implementing a care pathway, and be engaged with MSQC for the Breast Surgery QI project, which includes being a member of the Breast Care Committee meetings (either in person or virtual).
- 3. Multidisciplinary team (6 points total):
 - a. Participating hospitals will form a multidisciplinary team to review baseline data, guide quality improvement plans, and implement the care pathway. The multidisciplinary team should include the breast cancer surgeon champion, other surgeons who perform breast cancer surgery, nursing, patient navigator, plastics and reconstructive, breast radiology and others as relevant.
 - b. Hold a multidisciplinary meeting before March 29, 2024. Meeting notes, including attendees, must be submitted to the coordinating center with the final project submission. (2 points).
 - c. Two (2) additional multidisciplinary meetings (minimally) before December 1, 2024, which include a review of breast data (2 points each).
- 4. Implement Breast Surgery Care Pathway (4 points): Within the multidisciplinary team, create and modify the care pathway template to your hospital's practices. This will be implemented and utilized by the care team beginning in the preoperative period and extending into the postoperative period to ensure the implementation of each element of the breast surgery care pathway. Submit the final product to MSQC.

5. Perioperative Process Goals (20 points): Implement all the following process measures for each elective breast surgical patient as detailed below. Measurement Period is 4/1/2024 – 12/31/2024.

Preoperative Goals (8 points total)

- 5a: Preadmission teaching that discusses expectations after surgery, including multimodal pain management ≥ 70%, discussion of opioid-free surgery (if applicable), and expected use of surgical drains (if applicable) (4 points)
- 5b: Patient optimization discussion related to smoking cessation (if applicable) ≥ 80% (4 points)

Intraoperative Goals (4 points total)

• 5d: Use of intraoperative multimodal pain management ≥ 80% (4 points)

Postoperative Goals (8 points total)

- 5e: Postoperative <u>order</u> for multimodal pain management ≥ 70% (4 points)
- 5f: Opioid prescriptions meeting M-OPEN recommendations ≥ 70% (4 points)
- 6. Cancer-Specific Goals (10 points): Sites will pick two goals to focus on that need improvement. Included diagnosis codes: Cancer and DCIS diagnoses which are listed in the breast tab of 2024 Program Manual. Measurement Period is 4/1/2024 12/31/2024.
 - a. Preoperative MRI rate to $\leq 30\%$
 - b. Reduction of use of SLNB in women >70 years old to < 50%
 - c. Reduction of re-excision rates for positive margin after lumpectomy to ≤ 15%
 - d. Increase in the use of outpatient mastectomy to > 25%
- 7. Submit a QII Project Summary on or before <u>January 15, 2025</u>, which includes a narrative and activity tracking of the steps to implementation of the breast cancer surgery care pathway, successes and barriers, and analysis and next steps (a template will be available on the MSQC website). An additional 0-10 implementation points may be granted based on the detail of the project narrative, tracking log, and analysis, to be added to achieve the maximum of 40 project points.

Included CPT Codes:

CPT®	CPT® Description
Code	·
19301	19301: Mastectomy, partial (eg, lumpectomy, tylectomy, quadrantectomy, segmentectomy)
19302	19302: Mastectomy, partial (eg, lumpectomy, tylectomy, quadrantectomy, segmentectomy); with axillary lymphadenectomy
19303	19303: Mastectomy, simple, complete
19305	19305: Mastectomy, radical, including pectoral muscles, axillary lymph nodes
19306	19306: Mastectomy, radical, including pectoral muscles, axillary and internal mammary lymph nodes (Urban type operation)
19307	19307: Mastectomy, modified radical, including axillary lymph nodes, with or without pectoralis minor muscle, but excluding pectoralis major muscle



Quality Improvement Implementation, Option D (by invitation only): Preoperative Testing for Low-Risk Surgeries Project Time Period: 1/1/2024 – 12/31/2024

Background: The Preoperative Testing for Low-Risk Surgeries QII project is a cross-collaborative project between MSQC, the Michigan Value Collaborative (MVC) and the Michigan Program on Value Enhancement (MPrOVE). Routine preoperative testing before low-risk surgery has no known benefit and is an important target for de-implementation as it is overused, costly, and can lead to downstream care cascades involving invasive diagnostic testing¹.

As part of the Choosing Wisely® campaign the American Society of Anesthesiologists, Society of General Internal Medicine, American College of Surgeons (ACS), and the American Society for Clinical Pathology recommend against the use of routine laboratory studies before low-risk surgery (Table 1). Given the high prevalence of these services, eliminating unnecessary preoperative testing before low-risk surgery represents a key opportunity to improve quality, safety, and value in surgery. The pilot study in 2023, amongst MSQC participating sites, demonstrated wide variation in preoperative testing in low-risk surgeries, and continued work is needed.

Project Goal and Summary: In collaboration with MVC and MPrOVE, this project will continue to work toward reducing unnecessary, routine preoperative testing for low-risk surgeries, as well as implement interventions to heighten awareness and reduce variation among hospitals. Continuing sites who participated in the pilot project in 2023 are eligible if their average rate of preoperative testing is >20% from 4/1/2023 to 6/30/2023, and sites who did not select preoperative testing as their QII in 2023 will be eligible to participate.

Through a multi-faceted approach, invited sites will: 1) abstract preoperative testing variables on low-risk surgical cases, 2) implement a standard protocol defining appropriate use of preoperative testing, 3) employ strategies to promote adoption of the protocol, and 4) analyze MSQC, MVC, and internal data reports to monitor progress.

Eligible low-risk surgery cases will meet the procedure inclusion criteria:

- Minor hernia, laparoscopic cholecystectomy, and breast lumpectomy (Table 2), AND
- ASA classes 1 and 2, AND
- Surgical Priority = Elective, AND
- Surgical Procedure Tab: Is the CPT code the intended primary procedure = Yes

QI Implementation Goals and Requirements (40 total project points)

Goal #1: Data collection of 95% of preoperative tests for eligible procedures that were performed within 30 days prior to the surgery date, including obtained preoperatively on the day of surgery. The measurement period is 1/1/2024 - 12/31/2024 OR dates for all sites. (3 points)

The presence or absence of all of the following preoperative diagnostic tests on an eligible case must be captured in the MSQC Workstation to meet the numerator requirement:

- ECG
- Trans-thoracic echocardiography
- Cardiac stress test
- Chest Xray

- Urinalysis
- Complete blood count
- Basic metabolic panel
- Coagulation tests
- Pulmonary function tests

Goal #2 New sites only: Develop and implement a standard preoperative testing protocol for low-risk surgeries at your site. The protocol selected must be implemented no later than June 30, 2024. (20 total points)

- Goal 2a: Adopt a preoperative testing guideline protocol to implement at your site. Sites may choose the approach that fits best at their hospital. (10 points)
 - Adopt an existing protocol
 - American Society of Anesthesiologists' "Choosing Wisely" program (https://www.choosingwisely.org/clinician-lists/american-society-anesthesiologists-baseline-laboratory-studies-for-low-risk-surgery)
 - United Kingdom's NICE (National Institute for Health and Care Excellence) preoperative testing guidelines for elective surgery (https://www.nice.org.uk/guidance/ng45)
 - MPrOVE's "Waive the Workup" protocol https://sites.google.com/umich.edu/waivetheworkupmichigan/ho me\
 - Develop your own hospital preoperative testing protocol
 - Review and modify an existing protocol already in use at your hospital. You must also describe the process used for monitoring compliance, and interventions put into place to improve compliance.
 - Submit a copy of the preoperative testing protocol for low-risk surgery adopted by your hospital with your 2024 final project summary.
- Goal 2b: As part of the implementation process, sites must adopt clinical decision support tools to embed the preoperative testing protocol into practice. (10 points)
 - Examples of decision support tools selected for implementation include order sets, care pathways, CPOE pop-up messages/suggestions, BPA (Best Practice Advisory), documented inventory indicating review of existing order sets for concurrence with adopted testing guidelines. This list is not exhaustive.
 - Submit an example of at least one clinical decision support tool that was implemented or modified at your site with your 2024 final project summary.

Goal #2 Continuing sites only: include an in-depth analysis using a quality tool of your choice (e.g., A3, failure mode effects analysis, 5 Whys, Fishbone, etc.) with your 2024 final project summary of how the protocol and clinical decision support tools that were developed in the pilot year were modified or implemented differently to improve compliance with reducing preoperative testing. This should be completed with the multidisciplinary team early in the project year. (10 points)

Goal #3: Reduce the percentage of cases that receive one or more of the specified preoperative tests (as listed in Goal #1) by 20% (relative) as compared to the baseline rate, without having an increase of preoperative testing on the same day as surgery. (20 points continuing sites, 10 points new sites)

- Baseline period, continuing sites: 4/1/2023 12/31/2023 OR dates
- Baseline period, new sites: 1/1/2024 3/31/2024 OR dates
- Measurement period, all sites: 4/1/2024 12/31/2024 OR dates

Goal #4: Conduct a minimum of two multidisciplinary meetings with key stakeholders to review project requirements, implement project components and monitor project performance. (4 total points)

Goal 4a: host a project kickoff meeting held no later than March 31, 2024. (2 points)

- Goal 4b: host at least one follow-up multidisciplinary meeting between July and December 2024 to discuss protocol implementation, progress and barriers to implementation, and monitoring of compliance data (including MVC and MSQC preoperative testing data). (2 points)
- Meeting participants must include a general surgeon, anesthesiologist, MSQC/Quality dept representation; additional attendees can also include the hospital's MVC Site Coordinator (if applicable), a primary care provider (PCP), a representative from the preoperative clinic (if applicable), a surgical resident, and others as appropriate for your site.
- Meetings can be in person, virtual, or hybrid (project information shared over email, or multiple one- on-one meetings do not count toward this requirement).
- Designate a specific member of the team to serve as the Preoperative Testing for Low-Risk Surgeries QII project Point of Contact (POC). The POC will receive updated preoperative testing reports from MVC (in addition to your hospital's MVC point of contact). The POC will be responsible for sharing MVC reports with team members. The site must provide MVC (email address TBD) and MSQC (MSQC-Info@med.umich.edu) with the project designee's contact information no later than April 14, 2024.
- <u>For each meeting</u>, submit the meeting minutes and attendee list (with attendee name, credentials, and department represented) with your 2024 final project summary.

Goal #5 Performance Data Monitoring: Brief feedback regarding the value of the MVC and MSQC data reports and how the data was utilized must be submitted with your 2024 final project summary. (1 point)

- Sites will use data from several sources to monitor the progress of the protocol implementation.
 - MVC Preoperative Testing Reports (distributed to the POC and the MVC Site Contact)
 - MSQC case abstraction data on preoperative testing
 - o Internal hospital data collection for monitoring compliance and adoption of the preoperative testing protocol.

Goal #6: Submit an analysis on cases that received testing on the day of surgery, prior to In Room Time (2 points).

Goal #7: Submit the 2024 final project summary, due to the MSQC Coordinating Center no later than January 15, 2025.

- Improvement page of the MSQC website. The document will contain a narrative describing the adoption, implementation, and monitoring of a preoperative testing protocol for low-risk surgeries, along with successes, barriers, plans for moving forward with the project. Additional documents to be submitted with the summary include:
 - Copy of the preoperative testing protocol for low-risk surgery adopted by your hospital (from Goal 2a).
 - Example of at least one clinical decision support tool that was implemented at your site (from Goal 2b)
 - Continuing sites only: analysis using a quality tool of how the protocol and clinical decision support tools that were developed in the pilot year were modified or implemented differently to improve compliance with reducing preoperative testing (from Goal #2)
 - Meeting documents (minutes, participant list) from the project kickoff and subsequent follow- up multi-disciplinary meetings held during the project year (from Goal #4)
 - Feedback on the MVC and MSQC data reports (from Goal #5)
 - Analysis of cases with testing on day of surgery (from Goal #6)

Implementation Points

An additional 0-10 implementation points may be granted based on the detail of the project narrative, tracking log and analysis, to be added to achieve the maximum of 40 project points.

Professional Society	Recommendation
American Society of Anesthesiologists	Don't obtain baseline laboratory studies in patients without significant systemic disease undergoing low-risk surgery—specifically completed blood counts, metabolic panels, or coagulation studies.
	Don't obtain baseline diagnostic cardiac testing (e.g., echocardiography) in asymptomatic stable patients with known cardiac disease undergoing low or moderate risk surgery.
Society of General Internal Medicine	Don't perform routine pre-operative testing before low-risk surgical procedures.
American College of Surgeons	Avoid preoperative chest x-rays for ambulatory patients with unremarkable history and physical exam.
American Academy of Ophthalmology	Don't perform preoperative medical tests for eye surgery unless there are specific medical indications.
American Society for Clinical Pathology	Avoid routine preoperative testing for low-risk surgeries without a clinical indication.

Abdomin	al Hernias less than 3 cm and all Inguinal/Femoral Hernia Repairs ("Minor Hernia")
49505	49505: Repair initial inguinal hernia, age 5 years or older; reducible.
49507	49507: Repair initial inguinal hernia, age 5 years or older; incarcerated or strangulated.
49520	49520: Repair recurrent inguinal hernia, any age; reducible.
49521	49521: Repair recurrent inguinal hernia, any age; incarcerated or strangulated.
49525	49525: Repair inguinal hernia, sliding, any age.
49550	49550: Repair initial femoral hernia, any age; reducible.
49553	49553: Repair initial femoral hernia, any age; incarcerated or strangulated.
49555	49555: Repair recurrent femoral hernia; reducible.
49557	49557: Repair recurrent femoral hernia; incarcerated or strangulated.
49591	49591: Repair of anterior abdominal hernia(s) (ie, epigastric, incisional, ventral, umbilical, spigelian), any approach (ie, open, laparoscopic, robotic), initial, including implantation of mesh or other prosthesis when performed, total length of defect(s); less than 3 cm, reducible
49592	49592: Repair of anterior abdominal hernia(s) (ie, epigastric, incisional, ventral, umbilical, spigelian), any approach (ie, open, laparoscopic, robotic), initial, including implantation of mesh or other prosthesis when performed, total length of defect(s); less than 3 cm, incarcerated or strangulated

49613 49613: Repair of anterior abdominal hernia(s) (ie, epigastric, incisional, ventral, umbilical, spigelian), any approach (ie, open, laparoscopic, robotic), recurrent, including implantation of mesh or other prosthesis when performed, total length of defect(s); less than 3 cm, reducible 49614 49614: Repair of anterior abdominal hernia(s) (ie, epigastric, incisional, ventral, umbilical, spigelian), any approach (ie, open, laparoscopic, robotic), recurrent, including implantation of mesh or other prosthesis when performed, total length of defect(s); less than 3 cm, incarcerated or strangulated
umbilical, spigelian), any approach (ie, open, laparoscopic, robotic), recurrent, including implantation of mesh or other prosthesis when performed, total length of defect(s); less than 3 cm, incarcerated or strangulated
40050 40050 1
49650 49650: Laparoscopy, surgical; repair initial inguinal hernia
49651 49651: Laparoscopy, surgical; repair recurrent inguinal hernia
49659 49659: Unlisted laparoscopy procedure, hernioplasty, herniorrhaphy, herniotomy.
Laparoscopic Cholecystectomy
Laparoscopic Cholecystectomy 47562 47562: Laparoscopy, surgical; cholecystectomy
47562 47562: Laparoscopy, surgical; cholecystectomy
47562 47562: Laparoscopy, surgical; cholecystectomy 47563 47563: Laparoscopy, surgical; cholecystectomy with cholangiography
47562 47562: Laparoscopy, surgical; cholecystectomy 47563 47563: Laparoscopy, surgical; cholecystectomy with cholangiography

Resources

- Berlin NL, Yost ML, Cheng B, et al. Patterns and Determinants of Low-Value Preoperative Testing in Michigan. *JAMA Intern Med.* 2021;181(8):1115–1118. doi:10.1001/jamainternmed.2021.1653
- https://ihpi.umich.edu/featured-work/michigan-program-value-enhancement
- https://ihpi.umich.edu/news/routine-testing-surgery-remains-common-despite-low-value
- https://michiganvalue.org/our-work/mvc-value-coalition-campaigns-vccs/

References

¹Berlin NL, Yost ML, Cheng B, et al. Patterns and Determinants of Low-Value Preoperative Testing in Michigan. *JAMA Intern Med.* 2021;181(8):1115–1118. doi:10.1001/jamainternmed.2021.1653

2024 Michigan Spine Surgery Improvement Collaborative (MSSIC) Quality Initiative Performance Index Scorecard Hospital - Year 1 Measurement Period: 10/01/2023-09/30/2024, unless otherwise stated

Measure#	Weight	Measure Description	Points
Participation measures are based on calendar year meetings, calls, and audit (1/12/31/24).			
	15%	Meeting participation - Surgeon Champion	
_		Attended all 3 meetings	15
1		Attended 2 out of 3 meetings	10
		Attended 1 out of 3 meetings	5
		No Attendance	0
2	10%	Meeting and Abstractor Symposium participation – Clinical Data Abstractor. (It is required that <u>each</u> MSSIC Abstractor be present at MSSIC meetings and <u>all</u> abstractors are required to attend the Annual Abstractor Symposium.)	
		Attended all 4	10
		Attended 3 out of 4	6
		Attended 2 out of 4	3
		Attend 1 or none	0
	15%	Conference Calls Surgeon Champion (3 calls/year)	
2		Attended 3 calls	15
3		Attended 2 calls	10
		Attended 1 call	5
		No Calls	0
	10%	Conference Calls - Clinical Data Abstractor (8 calls/year)	
4		Participate on 8 calls	10
7		Participate on 7 calls	6
		Participate on 6 calls	3
		Participate on less than 6 calls	0
F	100/	Meeting participation - Administrative Lead (no designee)	
5	10%	Attend at least one triannual MSSIC meeting	10
		No Attendance	0
6	10%	Annual Audit Review – Data Review: Accuracy of data –	
		Complete and accurate 95-100% of the time	10
		Complete and accurate 90-94.9% of the time	5
		Complete and accurate < 90% of the time	0
7	15%	All official documents signed: IRB and Business Associate Agreement	

2024 Michigan Spine Surgery Improvement Collaborative (MSSIC) Quality Initiative Performance Index Scorecard Hospital - Year 1 Measurement Period: 10/01/2023-09/30/2024, unless otherwise stated

Measure#	Weight	Measure Description	Points
		Within 2 months of Coordinating Center approval date to	15
		proceed	
		Within 3 months of Coordinating Center approval date to proceed	12
		Within 4 months of Coordinating Center approval date to proceed	8
		Within 5 months of Coordinating Center approval date to proceed	4
		6 or more months of Coordinating Center approval date to proceed	0
		Hire Data Abstractor in a timely manner	
		Within 2 months of Coordinating Center approval date to proceed	15
	450/	Within 3 months of Coordinating Center approval date to proceed	12
8	15%	Within 4 months of Coordinating Center approval date to proceed	8
		Within 5 months of Coordinating Center approval date to proceed	4
		6 or more months of Coordinating Center approval date to proceed	0

Michigan Spine Surgery Improvement Collaborative (MSSIC)

2024 Performance Index Scorecard Measure Explanation:

Hospital Year 1

Measure number and description	Additional narrative describing the measure				
Participation measures are based on calendar year meetings, calls, and audit					
#1 - Meeting participation - Surgeon Champion: Three meetings per calendar year: #2 - Meeting and Abstractor Symposium participation – Clinical Data Abstractor. (It is required that each MSSIC Abstractor be present at MSSIC meetings, and all abstractors are	 12/31/24). April 26, 2024 August 9, 2024 (virtual) November 8, 2024 Please refer to the 2024 MSSIC Surgeon Champion Schedule for details and location of mandatory meetings and to the MSSIC Eligibility and Expectations Document for participation requirements. April 26, 2024 May 17, 2024 (Annual Abstractor Symposium) August 9, 2024 (virtual) November 8, 2024 Please refer to the 2024 MSSIC Data Abstractor 				
required to attend the Annual Abstractor Symposium.) Three meetings and Annual Abstractor Symposium per calendar year: #3 - Conference Calls Surgeon Champion (3 calls per year)	Schedule for details and location of mandatory meetings and to the MSSIC Eligibility and Expectations Document for participation requirements. Please refer to the 2024 MSSIC Surgeon Champion Schedule for mandatory calls and to the MSSIC Eligibility and Expectations Document for participation				
#4 - Conference Calls - Clinical Data Abstractor (8 calls per year)	requirements. Please refer to the 2024 MSSIC Data Abstractor Schedule for mandatory calls and to the MSSIC Eligibility and Expectations Document for participation requirements.				
#5 - Meeting participation - Administrative Lead (no designee), at least one meeting per year.	 April 26, 2024 August 9, 2024 (virtual) November 8, 2024 Please refer to the 2024 MSSIC Data Abstractor Schedule for details and location of mandatory meetings and to the MSSIC Eligibility and Expectations Document for participation requirements. 				
#6 - Annual Audit Review – Data Review: Accuracy of data	The abstractor(s) will participate in MSSIC Coordinating Center-led audits of charts of patients entered in the MSSIC registry to assure complete, quality data collection. Please see the MSSIC Manual of Operations, Section 2, "Abstractor Education and Training" for more details.				
#7 - All official documents signed: IRB and Business Associate Agreement	All required documents signed and returned to the MSSIC Program Manager. The timeframe associated with points earned begins with the email date from the				

	MSSIC Program Manager notifying the site of approval to proceed with documents & hire.
#8 - Hire Data Abstractor in a timely manner	It is the site's responsibility to notify the MSSIC Program Manager, in writing, when the data abstractor is hired. A start date for the abstractor must also be communicated. The timeframe associated with points earned begins with the email date from the MSSIC Program Manager notifying the site of approval to proceed with documents & hire.

MSSIC Patient questionnaires: Questionnaires are an essential data element and collection is expected and required as a condition of participation, described in the Eligibility and Expectations document. MSSIC defines a complete case as a fully abstracted medical record and entry into the registry as well as the collection and entry into the registry of a completed baseline questionnaire. All spine patients are asked to complete a validated health status questionnaire prior to surgery and then sampled patients in the MSSIC registry are asked to complete validated health status questionnaires at 3, 12, and 24 months after surgery. Each participating site is responsible for collecting this information. Patient-reported Outcome (PRO) data is an important measure of success for Quality Improvement Initiatives (QII).

2024 Michigan Spine Surgery Improvement Collaborative (MSSIC) Quality Initiative Performance Index Scorecard Hospital & ASF - Year 2 (1 site) Measurement Period: 10/01/2023-09/30/2024, unless otherwise stated

Measure#	Weight	Measure Description	Points
Participation measures are based on calendar year meetings, calls, and audit (1/1/2 12/31/24).			
		Meeting participation - Surgeon Champion	
		Attended all 3 meetings	15
1	15%	Attended 2 out of 3 meetings	10
		Attended 1 out of 3 meetings	5
		No Attendance	0
2	15%	Meeting and Abstractor Symposium participation – Clinical Data Abstractor. (It is required that <u>each</u> MSSIC Abstractor be present at MSSIC meetings and <u>all</u> abstractors are required to attend the Annual Abstractor Symposium.)	
		Attended all 4	15
		Attended 3 out of 4	10
		Attended 2 or less	0
		Conference Calls - Surgeon Champion (3 calls/year)	
		Attended 3 calls	15
3	15%	Attended 2 calls	10
		Attended 1 call	5
		No Calls	0
		Conference Calls - Clinical Data Abstractor (8 calls/year)	
4	10%	Participate on 8 calls	10
4	10%	Participate on 7 calls	6
		Participate on 6 calls	3
		Participate on less than 6 calls	0
5	15%	Meeting participation - Administrative Lead (no designee)	
5	15%	Attend at least one triannual MSSIC meeting	15
		No Attendance	0
		Annual Audit Review - Data Review: Accuracy of data -	
6	10%	Complete and accurate 95-100% of the time	10
	1070	Complete and accurate 90-94.9% of the time	5
		Complete and accurate < 90% of the time	0
Enhanced R	Enhanced Recovery After Surgery (ERAS), Phase 1 Performance Measures - (20 points below		
7	5%	Demonstration of multidisciplinary team engagement through the submission of quarterly meeting attendance sheet and minutes supporting discussion and establishment of ERAS.	
		4/4 meeting submissions	5
		3/4 meeting submissions	3
		2 or less/4 meeting submissions	0

2024 Michigan Spine Surgery Improvement Collaborative (MSSIC) Quality Initiative Performance Index Scorecard Hospital & ASF - Year 2 (1 site) Measurement Period: 10/01/2023-09/30/2024, unless otherwise stated

Measure#	Weight	Measure Description	Points
		No later than 9/30/24, each site will have submitted and obtained approval by the Coordinating Center, the following deliverables as evidence of a fully developed and implemented ERAS program:	
8	15%	ERAS protocol document outlining how each required component will be implemented. Template provided by the Coordinating Center.	7
		Submission of all ERAS supporting documents including pre-surgical patient education, order sets, protocols, applicable screen shots from EMR, discharge instructions, and risk-assessment tools implemented in support of the ERAS program.	8

Michigan Spine Surgery Improvement Collaborative (MSSIC)

2024 Performance Index Scorecard Measure Explanation:

Hospital & ASF Year 2

Measure number and description	Additional narrative describing the measure				
Participation measures are bas	ed on calendar year meetings, calls, and audit (1/1/24 -				
12/31/24).					
#1 - Meeting participation -	April 26, 2024				
Surgeon Champion:	August 9, 2024 (virtual)				
Three meetings per calendar	November 8, 2024				
year:	Please refer to the 2024 MSSIC Surgeon Champion				
	Schedule for details and location of mandatory meetings				
	and to the MSSIC Eligibility and Expectations Document				
	for participation requirements.				
#2 - Meeting and Abstractor	April 26, 2024				
Symposium participation –	May 17, 2024 (Annual Abstractor Symposium)				
Clinical Data Abstractor. (It is	August 9, 2024 (virtual)				
required that each MSSIC	November 8, 2024				
Abstractor be present at MSSIC	Please refer to the 2024 MSSIC Data Abstractor				
meetings, and all abstractors are	Schedule for details and location of mandatory meetings				
required to attend the Annual	and to the MSSIC Eligibility and Expectations Document				
Abstractor Symposium.)	for participation requirements.				
Three meetings and Annual					
Abstractor Symposium per calendar year:					
#3 - Conference Calls Surgeon	Diagon refer to the 2024 MCCIC Curgon Champion				
Champion (3 calls per year)	Please refer to the 2024 MSSIC Surgeon Champion Schedule for mandatory calls and to the MSSIC Eligibility				
Champion (3 cans per year)	and Expectations Document for participation				
	requirements.				
#4 - Conference Calls - Clinical	Please refer to the 2024 MSSIC Data Abstractor				
Data Abstractor (8 calls per year)	Schedule for mandatory calls and to the MSSIC Eligibility				
	and Expectations Document for participation				
	requirements.				
#5 - Meeting participation -	April 26, 2024				
Administrative Lead (no	August 9, 2024 (virtual)				
designee), at least one meeting	• November 8, 2024				
per year.	Please refer to the 2024 MSSIC Data Abstractor				
	Schedule for details and location of mandatory meetings				
	and to the MSSIC Eligibility and Expectations Document				
	for participation requirements.				
#6 - Annual Audit Review – Data	The abstractor(s) will participate in MSSIC Coordinating				
Review: Accuracy of data	Center-led audits of charts of patients entered in the				
	MSSIC registry to assure complete, quality data				
	collection. Please see the MSSIC Manual of Operations,				
	Section 2, "Abstractor Education and Training" for more				
details.					
Enhanced Recovery After Surger					
#7 - Demonstration of	During ERAS, Phase 1, Year 2 sites will demonstrate site				
multidisciplinary team	engagement through the submission of quarterly meeting				

engagement through the submission of quarterly meeting attendance sheet and minutes supporting discussion and establishment of ERAS. attendance and minutes which support the development and implementation of ERAS. The Coordinating Center will supply a "MSSIC Quarterly ERAS Meeting Minutes" template for sites to communicate meeting discussions concisely and provide a list of meeting attendees. Content should be high-level, and we are only interested in ERAS related discussion. The due dates for the **four deliverables** are as follows:

- Meeting between October 1 December 31, 2023.
 Submit form by January 5, 2024.
- Meeting between January 1 March 31, 2024.
 Submit form by April 5, 2024.
- Meeting between April 1 June 30, 2024. Submit form by July 5, 2024.
- Meeting between July 1 September 30, 2024.
 Submit form by October 5, 2024.

#8 - No later than 9/30/24, each site will have submitted and obtained approval by the Coordinating Center, the following deliverables as evidence of a fully developed and implemented ERAS program:

- MSSIC ERAS Protocol Document (template provided by the Coordinating Center) outlining the process of how each required component will be implemented at the site. The content should be high-level, and the template will provide fields for specific information that is requested.
- Submission of applicable <u>ERAS supporting</u> documents:
 - Order sets, protocols, pre-surgical patient education (booklets, class PowerPoints, online education links, etc.) and risk-assessment tools implemented in support of the ERAS program. These supporting documents will also be listed in each section of the ERAS Protocol Document to assist you.

MSSIC Patient questionnaires: Questionnaires are an essential data element and collection is expected and required as a condition of participation, described in the Eligibility and Expectations document. MSSIC defines a complete case as a fully abstracted medical record and entry into the registry as well as the collection and entry into the registry of a completed baseline questionnaire. All spine patients are asked to complete a validated health status questionnaire prior to surgery and then sampled patients in the MSSIC registry are asked to complete validated health status questionnaires at 3, 12, and 24 months after surgery. Each participating site is responsible for collecting this information. Patient-reported Outcome (PRO) data is an important measure of success for Quality Improvement Initiatives (QII).

2024 Michigan Spine Surgery Improvement Collaborative (MSSIC) Quality Initiative Performance Index Scorecard

Hospital & ASF - Year 3 & Older (29 sites)
Measurement Period: 10/01/2023-09/30/2024, unless otherwise stated

Measure#	Weight	Measure Description	Points	
Participation measures are based on calendar year meetings, calls, and audit (1/1/2 12/31/24).				
		Meeting participation - Surgeon Champion		
		Attended all 3 meetings	5	
1	5%	Attended 2 out of 3 meetings	3	
		Attended 1 out of 3 meetings	1	
		No Attendance	0	
2	3%	Meeting and Abstractor Symposium participation – Clinical Data Abstractor. (It is required that <u>each</u> MSSIC Abstractor be present at MSSIC meetings and <u>all</u> abstractors are required to attend the Annual Abstractor Symposium.)		
		Attended all 4	3	
		Attended 3 out of 4	2	
		Attended 2 or less	0	
		Conference Calls - Surgeon Champion (3 calls/year)		
	5%	Attended 3 calls	5	
3		Attended 2 calls	3	
		Attended 1 call	1	
		No Calls Conference Calls - Clinical Data Abstractor (8 calls/year)	0	
_	-0/	Participate on 8 calls	3	
4	3%	Participate on 7 calls	2	
		Participate on 6 calls	1	
		Participate on less than 6 calls	0	
F	4%	Meeting participation - Administrative Lead (no designee)		
5		Attend at least one triannual MSSIC meeting	4	
		No Attendance	0	
		Annual Audit Review - Data Review: Accuracy of data -		
6	10%	Complete and accurate 95-100% of the time	10	
J	5%	Complete and accurate 90-94.9% of the time	5	
		Complete and accurate < 90% of the time	0	
7		Each site: Dual collection rate of both a baseline and 90-day questionnaire (rounded to the nearest whole number) for cases with due dates 1/1/24 – 12/31/24.		
		Dual collection rate of 66% or more OR ≥ 20 percentage points of absolute improvement in the site's 2022 dual collection rate	5	

2024 Michigan Spine Surgery Improvement Collaborative (MSSIC) Quality Initiative Performance Index Scorecard Hospital & ASF - Year 3 & Older (29 sites) Measurement Period: 10/01/2023-09/30/2024, unless otherwise stated

Measure#	Weight	Measure Description	Points
		Dual collection rate of 51%-65% OR 10-19 percentage points of absolute improvement in the site's 2022 dual collection rate	3
		Dual collection rate of less than 50% AND < 10 percentage points of absolute improvement in the site's 2022 dual collection rate	0
		Each site: Dual collection rate of both a baseline and 1-year questionnaire (rounded to the nearest whole number) for cases with 1 year questionnaire due dates 1/1/24 – 12/31/24.	
8	5%	Dual collection rate of 45% or more OR ≥ 15 percentage points of absolute improvement in the site's 2022 dual collection rate	5
		Dual collection rate of 36%-44% OR 10-14 percentage points of absolute improvement in the site's 2022 dual collection rate.	3
		Dual collection rate of < 36% AND less than 10 percentage points of absolute improvement in the site's 2022 dual collection rate.	0
		Collaborative-wide measure: % of Opioid naïve, 1-2 level lumbar fusions meeting the MSSIC Opioid Prescribing guideline (≤ 320 MME) (score rounded to the nearest whole number)	
9	20%	80% or greater meeting the guideline of ≤ 320 MME or less	20
		65-79% of patients meet the guideline of ≤ 320 MME or less	10
		Less than 65% of patients meet the guideline of ≤ 320 MME or less	0
	20%	Demonstration of compliance for the following 3 MSSIC ERAS components: *Baseline rate is determined using cases with surgery dates 7/1/22-6/30/23.	
		1. Formal, pre-operative patient education that does not vary from surgeon to surgeon and is available to all spine patients	
10		80% or greater, OR ≥ 25 percentage points of absolute improvement in the *baseline rate	8
		50%-79%, OR 15-24 percentage points of absolute improvement from *baseline rate	4
		<50%, OR < 15 percentage points of absolute improvement from *baseline rate	0
		2. Limited fasting with a carbohydrate-rich drink up to two hours before surgery	

2024 Michigan Spine Surgery Improvement Collaborative (MSSIC) Quality Initiative Performance Index Scorecard

Hospital & ASF - Year 3 & Older (29 sites)
Measurement Period: 10/01/2023-09/30/2024, unless otherwise stated

Measure#	Weight	Measure Description	Points
		80% or greater, OR ≥ 25 percentage points of absolute improvement in the *baseline rate	8
		50%-79%, OR 15-24 percentage points of absolute improvement from *baseline rate	4
		<50%, OR < 15 percentage points of absolute improvement from *baseline rate	0
		3. Ambulation within 8 hours of surgery stop time	
		80% or greater, OR ≥ 25 percentage points of absolute improvement in the *baseline rate	4
		60%-79%, OR 15-24 percentage points of absolute improvement from *baseline rate	2
		<60%, OR < 15 percentage points of absolute improvement from *baseline rate	0
		Site Specific: Implementation of one Quality Improvement Initiative using MSSIC data. Sites may choose a site-specific initiative and submit it to the Coordinating Center no later than 9/8/23 for approval. The percentage goal of improvement is determined by the Coordinating Center and is communicated to the site via email after approval. The QI Plan was developed, implemented and there was	
		improvement in the target goal. In addition, both QI Reports were submitted on time. The breakdown is as follows:	00
		Site met 100% or greater of the target goal	20
11	20	Site met 75-99% of the target goal Site met 50-74% of the target goal	18 16
		Site met 1-49% of the target goal	14
		The QI Plan was developed and implemented, but there was no improvement to the target goal. In addition, both QI reports were submitted on time.	12
		The QI Plan was developed, implemented, and improvement could even have been made, but one of the QI reports was not submitted on time.	6
		The QI Plan was not developed or implemented; or both QI reports were not submitted on time.	0

2024 Michigan Spine Surgery Improvement Collaborative (MSSIC) Quality Initiative Performance Index Scorecard Hospital & ASF - Year 3 & Older (29 sites) Measurement Period: 10/01/2023-09/30/2024, unless otherwise stated

Measure#	Weight	Measure Description	Points
12	Bonus	Optional Bonus Performance Points: Bonus points are awarded to sites with above and beyond engagement and QI work as demonstrated by any of the following Coordinating Center approved activities: 1.) Host a MSSIC site visit 2.) Participate in, or develop and implement, an approved MSSIC pilot 3.) Complete an additional, approved site-level QI initiative and submit a MSSIC QI report template describing the initiative (see Support Document). 4.) Submit all 4 quarterly meeting minutes where multidisciplinary team (see Support Document) engagement is shown and MSSIC data and QI initiatives are being discussed (see Support Document). (*Sites will not exceed 100%. The bonus performance measure will only assist where points were lost on other performance measures.)	10

Michigan Spine Surgery Improvement Collaborative (MSSIC)

2024 Performance Index Scorecard Measure Explanation:

Hospital & ASF Year 3 & Older

Measure number and description	Additional narrative describing the measure				
Participation measures are based on calendar year meetings, calls, and audit (1/1/24 - 12/31/24).					
#1 - Meeting participation - Surgeon Champion: Three meetings per calendar year: #2 - Meeting and Abstractor	 April 26, 2024 August 9, 2024 (virtual) November 8, 2024 Please refer to the 2024 MSSIC Surgeon Champion Schedule for details and location of mandatory meetings and to the MSSIC Eligibility and Expectations Document for participation requirements. April 26, 2024 				
Symposium participation – Clinical Data Abstractor. (It is required that each MSSIC Abstractor be present at MSSIC meetings, and all abstractors are required to attend the Annual Abstractor Symposium.) Three meetings and Abstractor Symposium:	 May 17, 2024 (Annual Abstractor Symposium) August 9, 2024 (virtual) November 8, 2024 Please refer to the 2024 MSSIC Data Abstractor Schedule for details and location of mandatory meetings and to the MSSIC Eligibility and Expectations Document for participation requirements. 				
#3 - Conference Calls Surgeon Champion (3 calls per year)	Please refer to the 2024 MSSIC Surgeon Champion Schedule for mandatory calls and to the MSSIC Eligibility and Expectations Document for participation requirements.				
#4 - Conference Calls - Clinical Data Abstractor (8 calls per year)	Please refer to the 2024 MSSIC Data Abstractor Schedule for mandatory calls and to the MSSIC Eligibility and Expectations Document for participation requirements.				
#5 - Meeting participation - Administrative Lead (no designee), at least one meeting per year.	 April 26, 2024 August 9, 2024 (virtual) November 8, 2024 Please refer to the 2024 MSSIC Data Abstractor Schedule for details and location of mandatory meetings and to the MSSIC Eligibility and Expectations Document for participation requirements. 				
#6 - Annual Audit Review – Data Review: Accuracy of data	The abstractor(s) will participate in MSSIC Coordinating Center-led audits of charts of patients entered in the MSSIC registry to assure complete, quality data collection. Please see the MSSIC Manual of Operations, Section 2, "Abstractor Education and Training" for more details.				
#7 - Each site: Dual collection rate of both a baseline and 90-day questionnaire (rounded to the nearest whole number) for	MSSIC has modified its patient survey collection performance metric to help improve patient-reported outcomes data collection and quality. To measure change in patients and see if they improved in pain or physical				

cases with due dates 1/1/24 – 12/31/24.	function - both a baseline survey and a post- operative survey must be collected. Sites will maintain responsibility for collecting post-op surveys for all sampled patients (including those without a baseline) at all time periods, however the performance metric will target patients with pre and post surveys. Collection of post-operative surveys, even when the baseline is missing, is still critical for MSSIC data as it shares information on items like satisfaction, opioid usage, and return to work.
#8 - Each site: Dual collection rate of both a baseline and 1-year questionnaire (rounded to the nearest whole number) for cases with 1 year questionnaire due dates 1/1/24 – 12/31/24.	See explanation above for Measure #7.
#9 - Collaborative-wide measure: % of Opioid naïve, 1-2 level lumbar fusions meeting the MSSIC Opioid Prescribing guideline (≤ 320 MME). (Rounded to the nearest whole number.)	In recent years, MSSIC has collaborated the BCBSM CQI, Michigan-OPEN (https://michigan-open.org), whose mission is to support providers in combating opioid misuse. Both MSSIC data and the literature, including a study out of Mayo Clinic, support the judicious use of opiates as beneficial. Our data shows there is no evidence of decreased satisfaction or increased healthcare utilization with less prescribing. MSSIC analysis also showed a great deal of site variability in opioid prescribing patterns which represents an opportunity to make positive changes for opioid naïve patients.
	In 2023, MSSIC successfully implemented discharge prescribing guidelines in opioid naïve patients having 1-2 level lumbar decompression or 1-2 level Anterior Cervical fusion or arthroplasty. MSSIC Leadership and the MSSIC Executive Committee agreed that our opioid prescribing guidelines at discharge should be expanded to include lumbar fusion, 1-2 levels in 2024. It makes sense to broaden our scope of impact when it comes to discharge prescribing practices with opioid naïve patients.
#10 - Demonstration of compliance for the following 3 MSSIC ERAS components (*Baseline rate is determined using cases with surgery dates 7/1/22-6/30/23): 1.) Formal, pre-operative patient education that does not vary from surgeon to surgeon and is available to all spine patients	Demonstration of compliance for the following 3 MSSIC ERAS components: 1.) Formal, pre-operative patient education that does not vary from surgeon to surgeon and is available to all spine patients. For a site to mark "yes" for this variable, the patient must participate in that site's MSSIC approved, ERAS pre-operative patient education program. The aim is to educate the patient about ERAS protocols, to set realistic expectations for postoperative recovery, and to psychologically prepare the patient and family for the care program. Written and oral information must be taught and should be provided in detail. Patients that are

- 2.) Limited fasting with a carbohydrate-rich drink up to two hours before surgery
- 3.) Ambulation within 8 hours of surgery stop time
- admitted emergently as defined in the Master Variable List are excluded from the denominator.
- 2.) Limited fasting with a carbohydrate-rich drink up to two hours before surgery. Carbohydrate loading not only reduces insulin resistance but also improves muscle function by reducing nitrogen and protein loss. It is also associated with reduce preoperative thirst, hunger, and anxiety. Patients that are admitted emergently as defined in the Master Variable List, or are a Type 1 Diabetic, or are a Type 2 on insulin are excluded from the denominator.
- 3.) Ambulation within 8 hours of surgery stop time.

 Literature review of Enhanced Recovery After
 Surgery (ERAS) protocols, specific for spine surgery,
 all strongly support early ambulation as defined within
 hours of surgery stop time. A recently published
 MSSIC study correlated ambulation within 8 hours of
 spine surgery with fewer adverse events and
 improved outcomes. Exclusions from the denominator
 include wheelchair bound (non-ambulatory) before
 surgery, CSF leak, durotomy, and fusions four levels
 or greater.

#11- Site Specific: Implementation of one Quality Improvement Initiative using MSSIC data. Sites may choose a site-specific initiative and submit it to the Coordinating Center no later than 9/8/23 for approval. The Coordinating Center will determine the percentage goal of improvement and is communicated to the site via email after approval.

No later than 9/8/23, sites will submit for approval, a QII proposal to the Coordinating Center using the MSSIC Site-specific QI Project Submission Form. The measurement period for improvement is 10/1/23-9/30/24. Utilizing the MSSIC QI Report template, sites are to submit progress reports twice a year to communicate the project's development and movement. Sections 1-9 are due May 31, 2024, and the complete MSSIC QI Report, including the outcome of the project, is due January 31, 2025.

#12- Optional, Bonus Performance Points: Bonus points are awarded to sites with above and beyond engagement and/or QI work as demonstrated by any of the following Coordinating Center approved activities:

- 1.) Host a MSSIC site visit
- Participate in, or develop and implement, an approved MSSIC pilot
- 3.) Complete an additional, approved site-level QI initiative and submit a MSSIC QI report template describing the initiative

You can earn ten bonus points. The bonus is all or nothing. Descriptions for each numbered "Bonus" activity:

- 1.) A fully executed, in-person site visit must take place between 10/1/23-9/30/24. Sites will contact the MSSIC QI team to coordinate the details and obtain approval for the visit. It may be either a MSSIC Coordinating Center site visit or an expanded site visit where additional MSSIC sites are quests.
- 2.) Please contact the MSSIC QI team with a site-level pilot proposal or if interested in a pilot offered by the Coordinating Center.
- 3.) No later than 1/1/24, sites will submit for approval, an additional QII proposal to the Coordinating Center using the MSSIC Site-specific QI Project Submission Form. The Coordinating Center will determine the percentage goal of improvement. There must be at least nine full months of data in the measurement

- 4.) Submit all four quarterly meeting minutes where multidisciplinary team (see Support Document) engagement is shown and MSSIC data and QI initiatives are being discussed
- period to determine the outcome, ending with OR date 9/30/24. Sections 1-9 are due three months after project approval and the final MSSIC QI Report is due no later than January 31, 2025. A site must submit all QI reports on time and achieve at least 75% or greater of their goal in order to obtain bonus points.
- 4.) Sites will demonstrate, through the submission of quarterly meeting minutes and multidisciplinary attendance, MSSIC QI initiative discussion, and the sharing of MSSIC data. The Coordinating Center will supply a "MSSIC Quarterly Meeting Minutes" template for sites to communicate meeting discussions concisely and provide a list of meeting attendees. Content should be high-level, and we are only interested in MSSIC related discussion. All four due dates must be met to receive credit. The due dates for the **four deliverables** are as follows:
 - Meeting between October 1 December 31, 2023. Submit form by January 5, 2024.
 - Meeting between January 1 March 31, 2024.
 Submit form by April 5, 2024.
 - Meeting between April 1 June 30, 2024. Submit form by July 5, 2024.
 - Meeting between July 1 September 30, 2024.
 Submit form by October 5, 2024.

*Sites will not exceed 100% on the Performance Index. The bonus performance measure will only add ten points where points were lost on other performance measures. It will not assist where points were lost on participation measures.

<u>Performance Improvement Plans:</u> Sites performing at or below the zero-point threshold for any performance measure will be asked to complete a Performance Improvement Plan (PIP). The PIP will be used to guide additional coaching and determine the most helpful means of support and resources to the site.

MSSIC Patient questionnaires: Questionnaires are an essential data element and collection is expected and required as a condition of participation, described in the Eligibility and Expectations document. MSSIC defines a complete case as a fully abstracted medical record and entry into the registry as well as the collection and entry into the registry of a completed baseline questionnaire. All spine patients are asked to complete a validated health status questionnaire prior to surgery and then sampled patients in the MSSIC registry are asked to complete validated health status questionnaires at 3, 12, and 24 months after surgery. Each participating site is responsible for collecting this information. Patient-reported Outcome (PRO) data is an important measure of success for Quality Improvement Initiatives (QII).

2024 Michigan Society of Thoracic and Cardiovascular Surgeons (MSTCVS) Quality Collaborative Performance Index Scorecard

Measure#	Weight	Measure Description		
1	-	Accuracy of Data		
		5-star audit score	10	
	10	4-star audit score	8	
		3-star audit score	6	
		≤ 2-star audit score	0	
		Quarterly collaborative meeting participation – Surgeon and Data Manager Combined Attendance (January 1, 2024–December 31, 2024)		
		Surgeon and data manager attended 4 quarterly meetings	8	
2	8	Surgeon and data manager attended 3 quarterly meetings	6	
		Surgeon and data manager attended 2 quarterly meetings	4	
		Surgeon and data manager attended 1 quarterly meeting	2	
		Attended 0 quarterly meetings	0	
	4	Quarterly collaborative meeting participation – Alternate Surgeon (January 1, 2024–December 31, 2024)		
3		Alternate surgeon attended 1 quarterly meeting	4	
		Alternate surgeon attended 0 quarterly meetings	0	
		* Alternate surgeon performs cardiac surgery at the site and is not the physician champion		
	4	Quarterly data manager educational meeting - Data Manager (January 1, 2024–December 31, 2024)		
		Attended 4 data manager meetings	4	
4		Attended 3 data manager meetings	3	
		Attended 2 data manager meetings	2	
		Attended 1 data manager meeting	1	
		Attended 0 data manager meetings	0	
	4	Quarterly PERForm educational meeting - Perfusionist (January 1, 2024–December 31, 2024) *		
		Attended 3 PERForm meetings + Data Quality Report Submission	4	
5		Attended 2 PERForm meetings + Data Quality Report Submission	3	
		Attended 1 PERForm meetings + Data Quality Report Submission	2	
		Attended 0 PERForm meetings	0	
6	15	Collaborative-wide quality initiative 2024: Isolated CABG – Initial Ventilator Hours <6 (January 1, 2024–December 31, 2024)		
		Collaborative mean Initial Ventilator Hours <6 rate 70% or greater	15	
		Collaborative mean Initial Ventilator Hours <6 rate less than 70%	0	

2024 Michigan Society of Thoracic and Cardiovascular Surgeons (MSTCVS) Quality Collaborative Performance Index Scorecard

Measure#	Weight	Measure Description	Points
		Site specific quality initiative	
		Met improvement goal	15
7	15	Improved but did not meet goal	10
		Implemented plan but did not improve	5
		Improved but unable to implement plan or did not submit	0
		plan/progress report	
		Isolated CAB: O/E mortality for 12 months (January 1,	
		2024-December 31, 2024)	
8	20	O/E ≤ 1.0	20
		O/E ≤ 1.5	10
		O/E > 1.5	0
		Isolated Valve +/- CAB Mortality and Major Morbidity	
		OE for 36 months (January 1, 2022–December 31, 2024)	
9	20	O/E ≤ 1.0	20
		O/E ≤ 1.5	10
		O/E > 1.5	0
10		Extra Credit Opportunities: 1 point per approved	
10		activity for surgeons	
* Data Qualit	ty Reports	are a new on line tool with no baseline to report	

2024 Michigan Trauma Quality Improvement Program (MTQIP) Collaborative Quality Initiative Performance Index Scorecard

Measure#	Weight	Measure Description	Points
1		Data Submission Measurement period: 01/01/2024 – 12/31/2024	
	10	On time and complete 3 of 3 times	10
	10	On time and complete 2 of 3 times	5
		On time and complete 1 of 3 times	0
		Meeting Participation	
		Measurement period: 01/01/2024 – 12/31/2024	
		Surgeon, and Manager (TPM) or Abstractor (MCR) attend	9
		3 of 3 mtgs Surgeon, and Manager (TPM) or Abstractor (MCR) attend	
2	10	in 2 of 3 mtgs	6
		Surgeon, and Manager (TPM) or Abstractor (MCR) attend	0
		0-1 of 3 mtgs	0
		Registrar or Abstractor (MCR) attend annual data	1
		Data Validation Error Rate	
		Measurement period: 01/01/2024 – 12/31/2024	
		0-3.0%	10
3	10	3.1-4.0%	8
		4.1-5.0%	5
		> 5.0%	0
4	5	Performance Improvement (PI)Death Determination Documentation (defined as number of cases/people the hospital misses (does not submit) a PI death determination) Measurement period: 07/01/2023 – 06/30/2024	
		0-2 Missing Documentation	5
		3-4 Missing Documentation	3
		> 4 Missing Documentation	0
	8	Timely Low Molecular Weight Heparin (LMWH) Venous Thromboembolism (VTE) Prophylaxis Trauma Admits Measurement period: 07/01/2023 – 06/30/2024	
5a		≥ 52.5% of patients (≤ 48 hr)	8
		≥ 50.0% of patients (≤ 48 hr)	6
		≥ 45.0% of patients (≤ 48 hr)	3
		< 45.0% of patients (≤ 48 hr)	0
F.	2	Weight Based LMWH Protocol in Use Measurement period: 07/01/2023 – 06/30/2024	
5b		Yes	2
		No	0
6	10	Timely Surgical Repair Geriatric (Age ≥ 65) Isolated Hip Fracture (Fx)	
5		Measurement period: 07/01/2023 – 06/30/2024 ≥ 92.0% of patients (≤ 42 hr)	10
		≥ 92.0% or patients (≤ 42 nr)	10

2024 Michigan Trauma Quality Improvement Program (MTQIP) Collaborative Quality Initiative Performance Index Scorecard

Measure#	Weight	Measure Description	Points
		≥ 87.0% of patients (≤ 42 hr)	8
		≥ 85.0% of patients (≤ 42 hr)	5
		< 85.0% of patients (≤ 42 hr)	0
		RBC to Plasma Ratio in Massive Transfusion Measurement period: 01/01/2023- 06/30/2024	
7	10	Weighted Mean Points in Patients Transfused ≥ 5 Units 1st 4 hr	0-10
		Serious Complication Z-Score Trend Trauma Admits Measurement period: 07/01/2021- 06/30/2024	
8	10	< -1 (major improvement)	10
0	10	-1 to 1 or serious complications low-outlier (average or better rate)	7
		> 1 (rates of serious complications increased)	5
	10	Mortality Z-Score Trend Trauma Admits Measurement period: 07/01/2021- 06/30/2024	
9		< -1 (major improvement)	10
		-1 to 1 or mortality low-outlier (average or better)	7
		> 1 (rates of mortality increased)	5
	5	Patient Reported Outcomes Participation Measurement period: 07/01/2023 – 06/30/2024	
10		Signed agreement and >90% of patients contact information submitted	5
		No agreement OR Signed agreement and <90% of patients contact information submitted	0
		Collaborative: Timely Antibiotic Femur/Tibia Open Fx Measurement period: 07/01/2023 - 06/30/2024	
11	10	≥ 85% patients (≤ 90 min)	10
		< 85% patients (≤ 90 min)	0
		Total	100

2024 MTQIP P4P PI Scorecard Supporting Documentation

Measure 1: Data Submission: Partial/incomplete submissions receive no points. Complete data submission is defined as all cases submitted for the requested interval. To be considered complete, cohort 1 cases should have a missing rate of <10% for first name, last name, and MRN variables for 1/1/20 cases forward.

Measure 2: Meeting Participation: A surgeon may represent one trauma center only. Alternate surgeons are allowed but must be consistent (not rotating). The alternate surgeon must be an attending-level equivalent from the trauma call panel.

Measure 3: Data Validation Error Rate:

Centers not selected for validation this year will receive full points. Centers that are selected but do not schedule a visit will receive 0 points for the validation measure.

Measure 7: RBC to Plasma Ratio in Massive Transfusion

Step 1: Assign (weight) to each MTP patient's 4 hr PRBC/FPP ratio to the designated tier and points using the chart below.

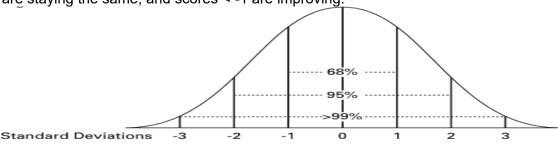
Step 2: Add the points and divide by the number of patients (weighted average). See the example below:

Step One				
PRBC to Plasma Ratio	Tier	Points		
<1.5	1	10		
1.6 – 2.0	2	10		
2.1 – 2.5	3	5		
>2.5	4	0		

Step Two (Example)					
Patient	PRBC	FFP	PRBC/FF	Tier	Points
			Р		
1	10	10	1.0	1	10
2	5	2	2.5	3	5
3	9	2	4.5	4	0
Total 15					
Total Points/Total #Patients = 15/3 = 5 points earned					

Measure 8 and 9: Z-Score Trend Calculation

The z-score measures a hospital's trend in #8 serious complications and #9 mortality over a three-year period. The z-score estimates the slope of a hospital's own linear trend line over time, standardized by the error estimate. The score indicates whether the hospital's performance is flat or trending upwards or downwards. If the z-score is one standard deviation away (either >1 or <-1), there is evidence that the hospital's performance is trending in one of these directions as opposed to being flat. Scores >1 are worsening, scores between 1 and -1 are staying the same, and scores < -1 are improving.



Measure 8: Serious Complication is Any Complication with a Severity Grade of 2 or 3

(Defined Below)

Complication Severity Grade 2

Definition: Potentially life-threatening complications

Complications: catheter-related bloodstream infection, central line-associated bloodstream infection, clostridium difficile, decubitus ulcer, deep vein thrombosis, enterocutaneous fistula, pneumonia, pulmonary embolism, unplanned return to ICU, unplanned return to OR

Complication Severity Grade 3

Definition: Life-threatening complications with a residual or lasting disability Complications: acute renal failure, acute respiratory distress syndrome, cardiac arrest, myocardial infarction, renal insufficiency, stroke/CVA, systemic sepsis, unplanned intubation, ventilator-associated pneumonia.

Collaborative Wide Measure:

Points are awarded based on the total collaborative result, not individual hospital result

Scoring When Center Has No Patients Meeting Measure Criteria

When a center has no patients to score for a measure, that measure will be excluded from its performance index denominator. Example: A center with no massive transfusion patients will have the measure (worth 10 points) excluded, and their maximum total numerator will be 90 points, the denominator will be 90 points, and a new % (points) calculated by dividing the numerator by the denominator

Filters

#4 PI Death Determination

Cohort: 2 (Admit Trauma Services)
No Signs of Life: Exclude DOAs

Default Period: Custom (7/1/23 to 6/30/24)

#5a: Timely LMWH VTE Prophylaxis in Trauma Service Admits

Practices > VTE Prophylaxis Metric

LMWH ≤ 48 hr

Cohort: 2 (Admit to Trauma Service) > 2-day LOS

No Signs of Life: Exclude DOAs Transfers Out: Exclude Transfers Out Default Period: Custom (1/1/23 to 6/30/24)

#5b: Weight-Based LWMH Protocol in Use

Points are awarded based on the submission of the following:

Screenshot of the center's protocol with the weight-based criteria visible in the image AND Screenshots of 5 patients using the protocol with the date and dosage visible in the image.

Submit screenshots to the MTQIP submission portal. For further instruction, see <u>Video</u> demonstration.

Default Period: Submit by 12/6/24.

<u>#6</u>: Timely Surgical Repair in Geriatric (Age ≥ 65) Isolated Hip Fracture

Cohort: 8 (Isolated hip fracture)

Age: ≥ 65

No Signs of Life: Exclude DOAs

Exclude: Transfers out, non-operative isolated hip fractures

Default Period: Custom (7/1/23 to 6/30/24)

#7: Red Blood Cell to Plasma Ratio in Massive Transfusion

Hemorrhage Cohort: 1 (All)

No Signs of Life: Include DOAs
Transfers Out: Include Transfers Out

Default Period: Custom (1/1/23 to 6/30/24)

#8: Serious Complication

Cohort: 2 (Admit to Trauma Service)
No Signs of Life: Exclude DOA
Transfers Out: Exclude Transfers Out
Default Period: Custom (7/1/21 to 6/30/24)

#9: Mortality

Cohort: 2 (Admit to Trauma Service)
No Signs of Life: Exclude DOA
Transfers Out: Exclude Transfers Out
Default Period: Custom (7/1/21 to 6/30/24)

#10: Patient-Reported Outcomes Participation

Points are awarded based on the following:

A signed agreement and \geq 90% of patients have contact information submitted defined as a validly formatted email or telephone number.

Cohort: 1 (All)

No Signs of Life: Exclude DOAs and all Deaths/Discharge to Hospice

Transfers Out: Include Transfers Out Default Period: Custom (7/1/23 to 6/30/24)

#11: Timely Antibiotic in Femur/Tibia Open Fractures - COLLABORATIVE WIDE MEASURE

Points awarded based on the total collaborative result, not the individual hospital result. Type of antibiotic administered along with date and time for open fracture of femur or tibia.

Eligible: Presence of acute open femur or tibia fracture based on AIS or ICD-10 codes (available on mtqip.org)

Exclude: Direct admissions. Transfers in. and Death in ED

Cohort: 1 (All)

No Signs of Life: Exclude DOAs Transfers Out: Include Transfers Out Default Period: Custom (7/1/23 to 6/30/24)

Abbreviations Key

AIS – abbreviated injury score CT- computed tomography CVA - cerebral vascular accident DOA - dead on arrival

ED - emergency department FFP - fresh frozen plasma FX - fracture

HR - hour

ICD - international classification of diseases

ICU - intensive care unit

LMWH - low molecular weight heparin LOS - length of stay

MCR - MTQIP clinical reviewer MIN - minute

MO - month

OR - operating room

PCP - primary care physician PI – performance improvement RBC – red blood cell

TBI – traumatic brain injury

TPM – trauma program manager VTE – venous thromboembolism YR - year

2024 Obstetrics Initiative (OBI) Collaborative Quality Initiative Performance Index Scorecard Measurement Period:01/01/2024 – 12/31/2024, unless otherwise specified

Measure#	Weight	Measure Description	Points
		OBI Semiannual Meeting Attendance	
		OBI Physician Champion (MD or DO actively practicing	
		inpatient maternity care) AND Midwife or Nurse Champion	
1	20%	(actively practicing inpatient maternity care), AND Clinical	
		Data Abstractor <i>combined</i> attendance (i.e. all three	
		individuals must attend each meeting to obtain full points)	
		Spring Semiannual meeting	10
		Fall Semiannual meeting	10
		Quality Initiative Engagement	
		Site-based Champion Team	
		1) Holds 6 quality meetings throughout the measurement	
2	5%	period AND	
_	0,0	2) Disseminates OBI data and discusses OBI initiative	5
		progress with unit staff and clinicians at least twice in the measurement period AND	
		Submits quarterly Program Progress and Monitoring	
		(PPM) Reports	
		Data Accuracy and Completeness of Abstracted Data	
		1) Data Quality Review Score ≥ 97%	
3	5%	2) 100% of cases from 01/01/2024 - 09/30/2024 completed	
3	5%	and submitted prior to 90 days postpartum	
		2 of 2 metrics met	5
		1 of 2 metrics met	3
		NTSV Cesarean Measures	
		Labor Dystocia Compliance	
		Site-Specific percentage of NTSV cesareans performed for	
		dystocia that meet national criteria for dystocia	
		≥ 80% compliance OR ≥ 20 percentage points of	15
		improvement* from baseline	
		≥ 70% compliance OR ≥ 15 percentage points of improvement* from baseline	10
		·	
4	50%	Management of Category II Fetal Tracings Site-Specific percentage of NTSV cesareans performed for	
		Cat II or indeterminate fetal heart rate tracing that have	
		documentation of management with an algorithm	
		Sites who previously selected Cat II as their OBI QII	
		project	
		≥ 80% compliance OR ≥ 25% improvement** from baseline to at least 60%	10
		Between 70 - 79% compliance OR ≥ 20% improvement**	
		from baseline to at least 50%	5
		≤ 69% compliance	0

2024 Obstetrics Initiative (OBI) Collaborative Quality Initiative Performance Index Scorecard Measurement Period:01/01/2024 – 12/31/2024, unless otherwise specified

		·	
Measure #	Weight	Measure Description	Points
		Sites who have NOT previously selected Cat II as their	
		OBI QII project	4.0
		≥ 60% compliance	10
		Between 50 - 59% compliance	5
		≤ 49% compliance	0
		Case Reviews	
		Group 1 Sites	
		Site-based Champion Team conducts 32 NTSV case	5
		reviews annually with submissions at least quarterly	5
		Group 2 Sites	
		Site-based Champion Team conducts 40 NTSV case	40
		reviews annually with submissions at least quarterly	10
		All Sites	
		At least 2 of 3 site Champions participate in a case review	
		training webinar at the beginning of the measurement	3
		period	
		Provide clinician to clinician feedback for those involved in	
		cases deemed likely inappropriate or indeterminate at least	4
		quarterly	
		Distribute summary feedback of case review findings to entire unit staff/ clinicians and to the OBI Coordinating	3
		Center at least quarterly	J
		Quality Improvement Planning	
		Group 1 Sites	
		Site-based Champion Team hosts in-person OBI site visit	
		within the first 2 quarters of 2024 AND meets virtually at	40
		least 3 times in the measurement period with an OBI	10
		Quality and Outreach Coordinator	
		Group 2 Sites	
		Site-based Champion Team meets virtually at least 2 times	
		in the measurement period with an OBI Quality and	5
		Outreach Coordinator	
		Pain Management Measures	
_	400/	Scheduled acetaminophen and oral NSAID	
		Scheduled acetaminophen AND oral NSAID ordered on patients without contraindication after NTSV cesarean birth	
5	10%	·	
		≥ 80% compliance	5
		Between 60 - 79% compliance	3
		≤ 59% compliance	0

2024 Obstetrics Initiative (OBI) Collaborative Quality Initiative Performance Index Scorecard Measurement Period:01/01/2024 – 12/31/2024, unless otherwise specified

Measure#	1		Points
		Total oral morphine equivalent (OME) prescribed appropriate with delivery method (excluding people with history of opioid use disorder) 1) 90% of vaginal births with no 3 rd /4 th degree laceration: total OME prescribed = 0 2) 90% of vaginal births with 3 rd /4 th degree laceration: total OME prescribed ≤ 75 3) 90% of cesarean births: total OME prescribed ≤ 113 3 of 3 metrics met	5 3
		≤ 1 of 3 metrics met	0
6	5%	Patient Voices Email address included on ≥ 90% OR ≥ 20 percentage points of improvement from baseline*** of patients abstracted from 01/01/2024 - 09/30/2024	2
		Race-Ethnicity Variable	
		Data element of "Race and Ethnicity" = "Unknown" ≤ 2%	3
		Severe Maternal Morbidity	-
7	5%	Submission of 2023 eCQM Severe Maternal Complications data to OBI Coordinating Center by 07/01/2024	5
		Optional Bonus Points Sites can earn up to 5 bonus points total. Sites cannot exceed 100 points total.	
Optional		Co-author an OBI publication Submit eCQM Severe Maternal Complications data to OBI Coordinating Center for first 2 quarters of 2024 before the end of the measurement period Share a patient story (with patient's permission) selected by	1
		OBI for dissemination to OBI members (e.g. via newsletter or social media)	
		100% of cases submitted and complete by 60 days postpartum for neonatal birthdates from 01/01/24 - 10/31/24	
		Participate on an OBI Committee Response rate for the OBI Patient Voices survey ≥ 40% Lead author (first/ senior) an OBI publication Present OBI data at a national meeting Present (or identify a patient who presents) at an OBI	2
		Semiannual meeting	

^{*}For sites with ≥ 30 births in the denominator for the measurement period; baseline set using data from 01/01/2023 - 09/30/2023

^{**}Baseline set using data from 01/01/2023 - 09/30/2023

^{***}Baseline set using data from 05/01/2023 - 09/30/2023