

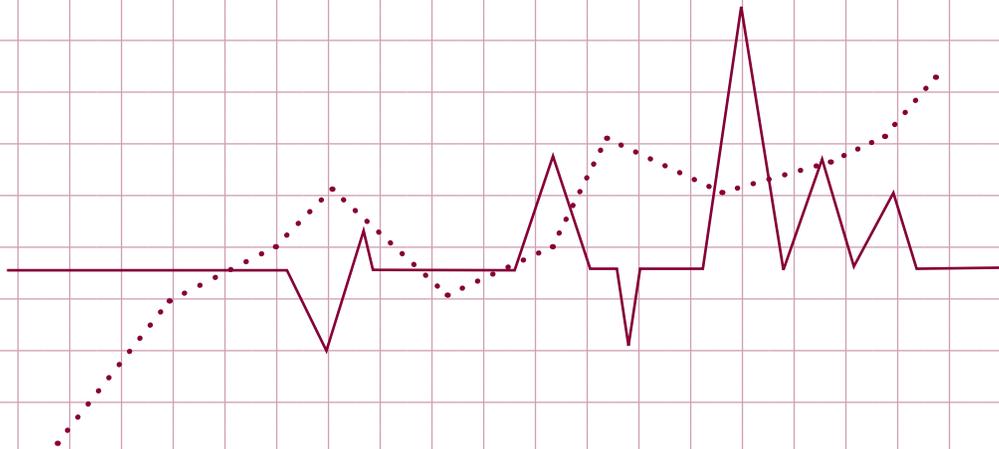
QUALITY IMPROVEMENT PROJECTS
DESIGNATED FOR MAINTENANCE OF CERTIFICATION, PART IV
UNIVERSITY OF MICHIGAN HEALTH SYSTEM

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MULTI-SPECIALTY 
PORTFOLIO APPROVAL PROGRAM



**American Board
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Multi-Specialty Portfolio Approval Program

The Multi-Specialty Portfolio Approval Program (Portfolio Program) offers a pathway for healthcare organizations to support physician involvement in organizational quality improvement (QI) initiatives and provide physicians from multiple specialties the opportunity to receive credit in their ABMS Programs for Maintenance of Certification (ABMS Programs for MOC).

The Portfolio Program began in 2009 with three ABMS Member Boards—the American Board of Family Medicine (ABFM), American Board of Internal Medicine (ABIM), and American Board of Pediatrics (ABP)—and the Mayo Clinic in Rochester, Minn., the first Portfolio Sponsor. It offers a streamlined approach for organizations that sponsor and support numerous well-designed QI efforts involving physicians across multiple disciplines to work with ABMS Member Boards to obtain MOC Part IV credit. It promotes organizational effectiveness and efficiency through team-based QI initiatives that are directly related to physicians' practice and influence the care they deliver. In doing so, the Portfolio Program delivers a more meaningful, relevant MOC experience for physicians that can be emulated in integrated multi-specialty systems across the country. Earlier this year, the Portfolio Program transitioned to ABMS.

Portfolio Program Fast Facts

19 ABMS Member Boards participate

33 healthcare organizations are approved Portfolio Sponsors

More than 60 healthcare organizations are in the Portfolio Sponsor application process

100 healthcare organizations are in the Portfolio Sponsors pre-application pipeline

650+ approved QI projects

4000+ physicians have received MOC Part IV credit

Why Organizations Participate

Organizations that become Portfolio Sponsors are committed to supporting physician involvement in both QI and MOC. Portfolio Sponsors:

- Can award their physicians MOC Part IV credit for participating in healthcare QI efforts that originate from within
- Can use MOC credit to involve physicians in QI efforts that are aligned with organizational priorities and goals
- Are able to approve their own QI efforts for MOC Part IV credit from participating ABMS Member Boards
- Reduce effort and time associated with applying to multiple ABMS Member Boards for approval of one or more QI efforts
- Reduce cost associated with obtaining MOC Part IV approval for multiple QI efforts that span multiple medical specialties
- Foster communication among each other to learn and share successful QI practices

How Physicians Benefit

Physicians participating in the Portfolio Program can:

- Avoid duplication of efforts by earning MOC Part IV credit for engaging in organizational QI initiatives
- Engage in a more meaningful, relevant MOC experience as QI initiatives are directly related to a physician's practice
- Promote organizational effectiveness and efficiency through team-based QI initiatives
- Work in an environment that supports their involvement in QI and MOC

Why ABMS Member Boards Participate

ABMS Member Boards that participate in the Portfolio Program:

- Reduce the administrative burden of approving multiple QI efforts that cross multiple specialties
- Reduce the cost of building and maintaining resources to facilitate a similar program
- Offer an established option for recognizing valid QI efforts in which their Diplomates are engaged through their organization
- Provide a vehicle for learning from, and collaborating with, other ABMS Member Boards
- Learn what QI efforts their Diplomates are working on or interested in
- Offer an opportunity to interact with leading organizations involved in QI and supportive of physician involvement in MOC

How Patients Benefit

When physicians participate in the Portfolio Program, their patients benefit because they:

- Receive care that is delivered using improved patient care processes and outcomes
- Receive care from physicians who are committed to improving patient care

For more information on the Portfolio Program, visit <http://mocportfolioprogram.org/approved-portfolio-sponsors/>

Examples of University of Michigan Health System QI Projects Designated for Part IV MOC

Summarized on the following pages are 8 of the more than 30 quality improvement (QI) projects performed at the University of Michigan Health System that were documented as meeting the requirements of the ABMS Multi-specialty Part IV MOC Portfolio Program. Participating physicians met their certifying Board's expectation for this Maintenance of Certification requirement. A list of all UMHS QI projects approved for Part IV MOC is available at <http://ocpd.med.umich.edu/moc/approved-umhs-projects>.

The descriptions are each composed of (1) the summary description of the project that is reported to the ABMS Multi-specialty Portfolio Program at the time the project is approved and (2) results and comments from the more detailed Application/Report that is prepared before participation is verified. An example of the full report for the first project summarized below is presented on the UMHS MOC website <http://ocpd.med.umich.edu/moc/part-iv-credit%20designation> (see section #3). A full report is available for all completed projects.

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I. Pregnancy Risk Assessment for Women with Congenital Heart Disease

(9/1/11 – 11/1/12)

Specific Aim

The aim of the project is to improve communication of risks of pregnancy to women with congenital heart disease as well as other physicians caring for these patients, including primary care physicians and obstetricians.

Problem

Woman with congenital heart disease often do not receive proper counseling regarding cardiovascular risks of pregnancy. Women with significant heart disease often present well into pregnancy in situations where a pre-pregnancy evaluation or therapeutic regimen might have improved the pregnancy outcome. It is the recommendation the 2008 American College of Cardiology/American Heart Association guidelines for the care of adults with congenital heart disease that women undergo pre-conception counseling and risk stratification prior to pregnancy.

Patient Population

The project includes women aged 18–45 with congenital heart disease. The project will involve women seen in the adult congenital cardiology clinic as well as adult women followed within the pediatric cardiology clinics. The physician will be responsible for identifying women aged 18–45 in their clinic. The average interval of follow-up for adults with congenital heart disease is one year, so it is not likely that there will be significant duplication of patients.

Measures

The measure will be documentation of the CARPREG score, a well-accepted method for risk stratification of women with heart disease. The numerator will be the number of patients for whom the CARPREG score is documented in the clinic letter. The denominator will be the number of women aged 18–45 seen by the physician practitioner. Both of the 2 adult congenital physician practitioners will participate in the project. All pediatric cardiologists who frequently see adult patients (>2–3 per month) will be encouraged to participate. For pediatric cardiologists, the pre-intervention compliance rate will be assumed to be 0 as pediatric cardiologists have not been educated as to pregnancy risk assessment tools.

Interventions

Education and calculation worksheet.

The initial phase of the intervention consisted of an education phase in which clinicians were educated regarding the CARPREG score. Participating physicians received the original paper describing the justification for the score, as well as how to calculate the score. They were also instructed in how to use a worksheet to simplify calculating the score. The physician lead for the project (TC) provided the physician education.

Process for identifying relevant patients and recording scores.

The operational process differed slightly between the adult congenital clinic and the pediatric cardiology clinic because clinic processes and resource vary. Implementing the process involved educating other health care team members regarding the process.

Adult Congenital Program Process

The adult congenital program assistant reviewed the clinic list for the day and identified women aged 18–45.

- The clinic nurse brought this patient list to clinic. As patients were brought back, the clinic nurse affixed the CARPREG calculation worksheet to the patients' paperwork
- The attending physician calculated the CARPREG score at the time of the visit, and dictated the CARPREG score into the diagnosis list of the clinic letter

Pediatric Clinic Process

- Currently, pediatric cardiologists review their patient list the day prior to or the morning of clinic and note what studies the patients will require at their visit. This is then written on a dry erase board next to the patients name during clinic
- When pediatric cardiologists reviewed their patient lists, they identified women aged 18–45 and noted "CARPREG" next to the patients name
- The medical assistants then transcribed this to the dry erase board. This served as a reminder to do the CARPREG assessment
- CARPREG calculators were made available in bins in the clinic team rooms. This was done by the physician lead (TC)
- The attending physician calculated the CARPREG score at the time of the visit, and dictated the CARPREG score into the diagnosis list of the clinic letter. If a fellow performed the dictation, the fellow would dictate the CARPREG score

Results

Time Period	Eligible Women Seen (Aged 18–45 with Congenital Heart Disease)	
	N	% with CARPREG Score
Baseline: 9/1/11 – 12/31/11	150	3%
Post-intervention 4/1/12 – 6/30/12	117	74%
Post-adjustment 9/1/12 – 10/31/12	96	80%

Comments/Plans

The continued reminder to physicians to document the CARPREG score should make the process sustainable. During the last time period a problem encountered by both clinics was the transition to a new electronic medical record EPIC™ (MI-Chart) that did not have a location for recording CARPREG scores. A documentation template was ultimately created to facilitate recording of CARPREG scores. This resulted in compliance rates of greater than 90% in the last month of the period.

2. Improving Diabetic Foot evaluation rates in UMHS Family Medicine Clinics

(6/30/13 – 11/30/13)

Specific Aim

To improve the rate of diabetic patients having had their feet examined and documented each year.

Problem

Screening for peripheral neuropathy is an important part of the management of diabetic patients. Diabetic neuropathy is one of the leading contributors to foot infections and ultimately amputations and other morbidities. Therefore yearly diabetic foot exams are an important quality measure monitored by multiple 3rd party payors and are a component of most if not all diabetes guidelines. In the past, the Department of Family Medicine used a population management system (Cielo[®]) which offered point of care (POC) prompts to help remind physicians to perform the diabetic foot exam when one was due. With the implementation of EPIC (MI-Chart) in August of 2012, Cielo was no longer available as a POC reminder. In its place, the reminder system within MI-Chart, “Best practice advisory” (BPA), required a different workflow that was initially not proactively planned as a standard process. We are now working to develop a standard workflow and plan to monitor adherence to that workflow in an effort to improve the rates of foot exams performed and documented.

Patient Population

Adult (age ≥ 18) diabetic patients that see a UM family physician at one of 5 Family Medicine outpatient clinics.

Measures

Percent of patients from the above patient population that have a diabetic foot exam performed and recorded in MI-Chart, as a Health Maintenance Modifier, in the last 365 days. Numerator is # of patients that have had the diabetic foot exam done and documented properly in the last 365 days. Denominator is the total number of diabetic patients fitting the above patient population.

Interventions

- Feedback of performance data to those performing care (8/28/13)
- Educational intervention with the faculty on how to properly document foot exams in MI-Chart
- Develop/update standard procedures for medical assistants to see if a patient is due for the foot exam (BPA) and how to prepare the patient during the visit (remove shoes and socks, have monofilament available)
- Educate medical assistants regarding these procedures
- Educate medical assistants on initiation of documentation of the foot exam for the physician
- Panel managers review GAP reports to correct errors, and contact patients overdue to schedule appointment or note on upcoming appointment to have it done
- Discussion with the family physicians at their faculty meeting to educate them on the methods and importance of properly documenting their foot exams
- Panel managers notifying patients that they were due for the foot exam on an individual basis and noting in the appointment summary to alert medical assistants and providers
- In some sites, medical assistants performed and documented part of the foot exam (the monofilament test) to help improve provider efficiency

**This was a brand name, previously called clinfotracker and now integrated into Crimson which is a product of the Advisory Board company. It was developed by members of the UMHS Department of Family Medicine.*

Site/Group	Baseline 6/30/13		Post-Intervention 9/16/13–10/4/13		Post-Adjustment 10/25/13–11/15/13	
	N Patients	% with foot exam	N Patients	% with foot exam	N Patients	% with foot exam
Fam Med Center 1	817	65%	839	73%	840	72%
Fam Med Center 2	809	56%	828	59%	836	67%
Fam Med Center 3	299	60%	300	76%	304	74%
Fam Med Center 4	452	53%	476	82%	492	81%
Fam Med Center 5	629	52%	655	61%	671	60%
All Family Medicine	3,006	58%	3,098	68%	3,143	70%

Note: Number of patients is the number with diabetes in the UM Health System's registry of diabetic patients on the last day of the intervention period. The percent is the number of these patients who, on the last day of the intervention period, have had a diabetic foot exam performed in the previous 365 days.

Comments/Plans

Providers now know how to correctly document the intervention. Staff training has occurred to assure proper patient preparation for the foot exam. The number and percent of diabetic patients with foot exams within the past year should increase as more of these patients are seen at periodic visits. We will need to continue to train new staff to maintain our recent gains, and continue to require excellence from them.

3. Eliminating Non-Medically-Indicated Planned Delivery

(1/1/11 – 9/30/12)

Specific Aim

Von Voightlander Women's Hospital aims to eliminate non-medically-indicated early term planned delivery.

Problem

Non-Medically-Indicated early (prior to 39 weeks') planned delivery is common nationally and at the VonVoightlander Women's Hospital (VVWH). It is associated with excess neonatal morbidity. Appropriately delaying non-medically-indicated planned deliveries to 39 weeks reduces neonatal morbidity, reduces admission to the NICU, and lowers costs.

Patient Population

Parturients with no fetal or maternal medical indications for early delivery.

Measures

Denominator all deliveries (cesarean or vaginal) 37 weeks + 0 days to 38 weeks +6 days gestation, with Leapfrog/Joint Commission PC-01 exclusion criteria applied, and in addition. (Roughly, this group is spontaneous labor + elective deliveries). Numerator: elective deliveries (induced labor or planned cesarean).

Interventions

Education

- Educate providers about evidence that suggests early elective delivery is associated with significant neonatal risk
- Educate about the multiple national efforts directed toward reducing the rate of elective early delivery
- Educated about standard of care that has been defined due to these widespread efforts
- Define department expectations (0 early elective deliveries)
- Methods
 - M&M presentation (August 2012)
 - Division meeting presentations (June 2012; July 2012)
 - Department-wide communication (September, 2012)
 - e-mail to all disciplines (including staff nurses)

Define QI process (peer review) that will address non-compliance.

- Early non-medically-indicated delivery is now a Department of Ob-Gyn QI Indicator (June 2012)
- Prenatal Joint Practice Committee Guideline endorses the avoidance of early non-medically indicated delivery (May 2012)
- All Early non-medically-indicated deliveries (QI Indicator) reviewed by QI Committee peer review. Determination and action taken (verbal, written feedback to surgeons, department action via OPPE when necessary) QI 'Advisory Letters' sent to non-compliers

Inform department about rates

- Present to individual faculty their 2011 data and department data (blinded) (August, 2012)

Plan Birth Center O.R process improvement: “Scheduled Cesarean Team” (Nov., 2012–ongoing)

- Committee comprised of Clinical leaders, Residency leaders, Residents, Nursing leaders, Staff nurses, Scrub techs
- Improved access: 3 spots per day instead of 2 to 3
- Defined scheduling guidelines
- Maintain direct access of surgeons to the schedule (no intermediary)
- Separate Team (outside of on-duty call team) for scheduled cases (separate faculty, residents, anesthesia, operating room staff)

QI Director presented data to faculty at M&M, Faculty meetings. Faculty feedback invited and received. (Nov. and Dec., 2012)

- Individual rates reported to faculty, QI Letters sent to non-compliers (ongoing)
- QI Director sent resources to faculty (January 2013)
 - These included patient education materials to help convince patients of the reason to wait
 - Posters sent to all offices from March of Dimes
 - Communication sent via email about the success of the effort, reminding everyone of the goals
- QI Director worked with individual faculty to identify barriers, plan strategies for compliance

New PJPC Guideline (Feb 2013)

- with updates, mainly in language
- Also added a required consultation with Maternal Fetal Medicine in order to schedule an early planned non-medically-indicated delivery. (“Hard Stop”)

Time Period	N Eligible Deliveries 37+0 to 38+6 Weeks Gestation	N of These Elective	% Elective
Baseline: 1/1/11 – 12/31/11	358	54	15%
Post intervention: 07/01/2012 – 09/30/2012	90	2	2.2%
Post-adjustment: 01/01/2013 – 06/30/2013	129	2	1.6%

Comments/plans

Ob/Gyn Database will be fully operational by August, 2013. This will automate much of the data gathering and make manual analysis/review easier to accomplish and to track.

- Implementation of Scheduled Cesarean Team July 1, 2013
- Statit® OPPE Dashboard will display individuals performance, real-time
- This project will continue with quarterly reporting within the department, then twice-yearly reporting to the Joint Commission, annual reporting to Leapfrog

*This is a brand name product used by the UMHS Office of Clinical Affairs to track Joint commission Ongoing Professional Performance Evaluation Metrics as defined by a Department.

†Leapfrog website: <http://www.leapfroggroup.org/>

4. Enhancing Access to Primary Care at the VA Ann Arbor Healthcare System

(5/1/11 – 9/30/12)

Specific Aim

As a result we wanted to improve the access of VA patients to care as demonstrated by the following national VA performance measures:

- Percentage of appointments made on the same day as the desired date by the patient (target of 66%)
- Percentage of appointments made within 7 days of the desired date by the patient (target 90%)
- Percentage of all visits to primary care that include telehealth or telecommunication with either the primary care physician or primary care nurse case manager (target 20%)

Problem

Nationally the VA has a significant interest and focus on assuring that all patients can be seen in a timely fashion (both new patients and established patients). The VA Ann Arbor is not currently meeting the targets for access as established by the Department of Veterans Affairs. Additionally the VA Central Office wants primary care providers to integrate telephone care/visits into their daily management of patients. The VA Ann Arbor is not currently meeting targets for telephone visits.

Patient Population

The population of patients is all veterans utilizing primary care clinic at the VA Ann Arbor Healthcare System. This includes established patients and new patients to primary care.

Measures

National VA performance measures for Access

- Percentage of appointments made on the same day when the patient desires to be seen the day they call for an appointment (target of 66%). Numerator: appointments made the day of the request when same day requested/ Denominator: all primary care appointments requested to be scheduled specifically on the day of the request/call
- Percentage of appointments made within 7 days of the desired date by the patient (target 90%). Numerator: date of appointment is within 7 days of the desired date as determined by the patient/ Denominator: all primary care appointments for a provider or a nurse manager in primary care

National VA performance measures for Telephone Visits:

- Percentage of all visits to primary care that include telehealth or telecommunication with either the primary care physician or primary care nurse case manager (target 20%). Numerator is phone visits by provider or assigned nurse case manager/ Denominator is all visits to the primary care provider or primary care nurse manager

Interventions

- Providers were educated about the concepts of panel management, return visit intervals and open access scheduling. This occurred during scheduled administrative meetings prior to the intervention period beginning
- Development and Implementation of a panel management process. This involved providing primary care providers prospective lists of their upcoming appointments two weeks in advance. In addition, providers were given lists of all patients with future appointments who were currently scheduled beyond 7 days of their desired date. Providers reviewed this list to evaluate if any of the current scheduled patients were inappropriately scheduled (i.e. didn't truly need an appointment) or could be seen by other members of the primary care team to provide the appropriate clinical care (i.e. Nurse Case Management face-to-face or phone visit, Clinical Pharmacist clinic face-to-face or phone visit, integrated primary care mental health clinic or specialty clinics). Providers were also encouraged to create additional access by opening clinics if possible
- The clerical staff was taught how to print the data for the providers they support
- Physicians received data on return visit interval for their patients as well as peer comparison data

Additional telephone visit interventions:

- Primary care providers and nurses were instructed on what types of patients would be appropriate for phone visits versus face-to-face visits
- PACT (Patient Aligned Care Team) phone clinics were created and mandated for all primary care providers and primary care nurse managers. These clinics could be utilized in place of traditional face-to-face visits

Measurement Period	Access			Telephone Visits	
	N of Appointments	% on Desired Date	% within 7 Days of Desired Date	N of Visits	% with Telehealth or Telecommunication
Baseline: 5/1/11–8/31/11	190	39%	74%	7,181	4%
Post-Intervention 10/1/11–3/31/12	155	60%	88%	6,227	34%
Post-adjustment: 4/1/12–8/31/12	269	27%	80%	5,583	35%

Comments/Plans

This project has fundamentally helped to change the manner in which our group of physicians and support staff view access to care. Panel management will be an ongoing part of standard weekly and daily work within each primary care teamlet. Additional LPN and RN support staff are being hired to help with panel management when a provider does not have consistent staff support. The goals will continue to be to create

timely access for all our veteran patients. Additionally, we have been successful in meeting our telephone visit measure and anticipate that this will remain a standard part of our access to patients in the coming years. Telephone visits will remain a part of the standard physician schedule and we will continue to monitor their use as will the VA nationally and regionally.

5. Use of Point of Care Clinical Reminders to Improve Tdap Vaccination Rates

(12/1/08 – 11/30/11)

Specific Aim

This project aims to improve Tdap (Tetanus, diphtheria, acellular pertussis) immunization rates for patients with no prior Tdap documented.

Problem

The Advisory Committee on Immunization Practice (ACIP) added a new recommendation for Tdap immunization booster for all adults in 2006 (MMWR 226;55 (RR-17):1-33). Despite this, there was only minimal increase in Tdap vaccination for eligible patients ages 11–64 who obtained care at the University of Michigan Family Medicine clinics subsequent to this new guideline.

Implementing substantial changes in clinical practice, even with clear cut, agreed upon guidelines, is challenging to accomplish. Even when physicians agree with the changes, clinical practice does not easily follow suit. At UMHS, the Department of Family Medicine uses a patient population management database program (Cielo Clinics), which has been in place since 2004 to facilitate clinical care which follows evidence-based guidelines. Among the tools available with this program is a point of care reminder system which prompts specific interventions for a defined patient population.

This project was undertaken to both improve Tdap vaccination rates in our patient population, and more broadly, to evaluate the effectiveness of a point of care reminder system which can be applied to multiple clinical areas if shown to be an effective intervention.

Patient Population

All patients without prior documentation of Tdap vaccination, ages 11–64, receiving care at UMHS Family Medicine clinics.

Measures

Denominator is number of patients seen without Tdap documentation during base year and project time period. Numerator is number of these patients receiving a Tdap vaccine during base year and project time period. The fraction = response rate to cielo prompt for Tdap vaccine.

Interventions

Given the CDC recommendation to improve Tdap Vaccination rates, in discussion during faculty meetings we established a point of care reminder with the following simple wording: “Tdap Vaccine”. The agreed upon method of addressing this reminder was that when it occurred, the medical assistant would confirm with the patient if they had not had a Tdap vaccine administered elsewhere. If so, they would enter this information into the electronic medical record (Careweb[®]). If not, the providers agreed to offer the Tdap vaccine to the patient. Providers agreed to inform patients of the differences between Td and Tdap when this question arose. If the patient agreed to the immunization, the vaccine would be administered and recorded in careweb.

**This is a “home grown product”, electronic health record from UMHS that is no longer actively used as of June 2014.*

Results

Time Period	Number of Patients in Overall Population	Number of Patients with Tdap documented	Percentage of patients with Tdap
12/1/2008–11/30/2009	42,738	13,733	32%
12/1/2009–11/30/2010 (post intervention)	42,361	24,781	58%
12/1/2010–11/30/2011 (post-adjustment)	43,081	30,298	70%

Cielo Responses

Time Period	Number of Patients Seen	Cumulative Tdap Administered	Prompt Addressed and Tdap given or documented	Prompt Addressed and Tdap not given	Prompt not addressed
12/1/2009–11/30/2010	16,273	6,828	6,828	5,030	4,415
12/1/2010–11/30/2011	21,776	12,861	6,033	1,516	2,369

Note: 5517 patients had additional documentation of Tdap during the post-adjustment period. Cielo data indicate 6033 had Tdap addressed or documented during this same time frame. This discrepancy is due to a process issue, not all providers indicated outside Tdap by entering the data directly into careweb, instead, this was only indicated on the cielo form. Indicating a Tdap on a cielo form does not automatically result in documentation in Careweb.

Comments/Plans

To facilitate the process further, we are establishing standing orders for Tdap administration by the medical assistant when a prompt occurs. The other causes are variations that are typically individual (not systemic). To address individual variation monthly team meetings will occur at which individual process issues will be addressed.

6. Improving the Appropriateness of Packed Red Blood Cell Transfusions for Patients admitted on the General Medicine Services

(2/9/10 – 4/1/12)

Specific Aim

This project aims to improve the overall appropriateness of packed red blood cell (RBC) transfusions given to patients admitted to the general medicine services. This will be accomplished by improving physicians' understanding and awareness of institutional guidelines that are in place to help direct RBC transfusion practices, as well as allowing physicians to review their own transfusion practices and compare them to their peers' in addition to monitoring the appropriateness of their RBC transfusions based on lab parameters.

Problem

A growing body of data shows that judicious use of packed red blood cells is associated with improved patient outcomes as well as decreased cost to the hospital system. To this end, institutional guidelines were developed to promote a more restrictive transfusion practice here at the U of M. Despite this, transfusion practices vary significantly between individual physicians and need to be monitored and reviewed.

Patient population

General medicine adult patients admitted to inpatient medicine services at University Hospital.

Measures

Percentage of total orders for red blood cell transfusions that do not meet pre-transfusion lab criteria for hemoglobin and hematocrit

Denominator: All RBC transfusions for patients on the General Medicine service during the measured quarter. Excluding RBC transfusions used to treat bleeding episodes (i.e., transfusions of four or more units within four hours or less, active GI bleeding, etc)

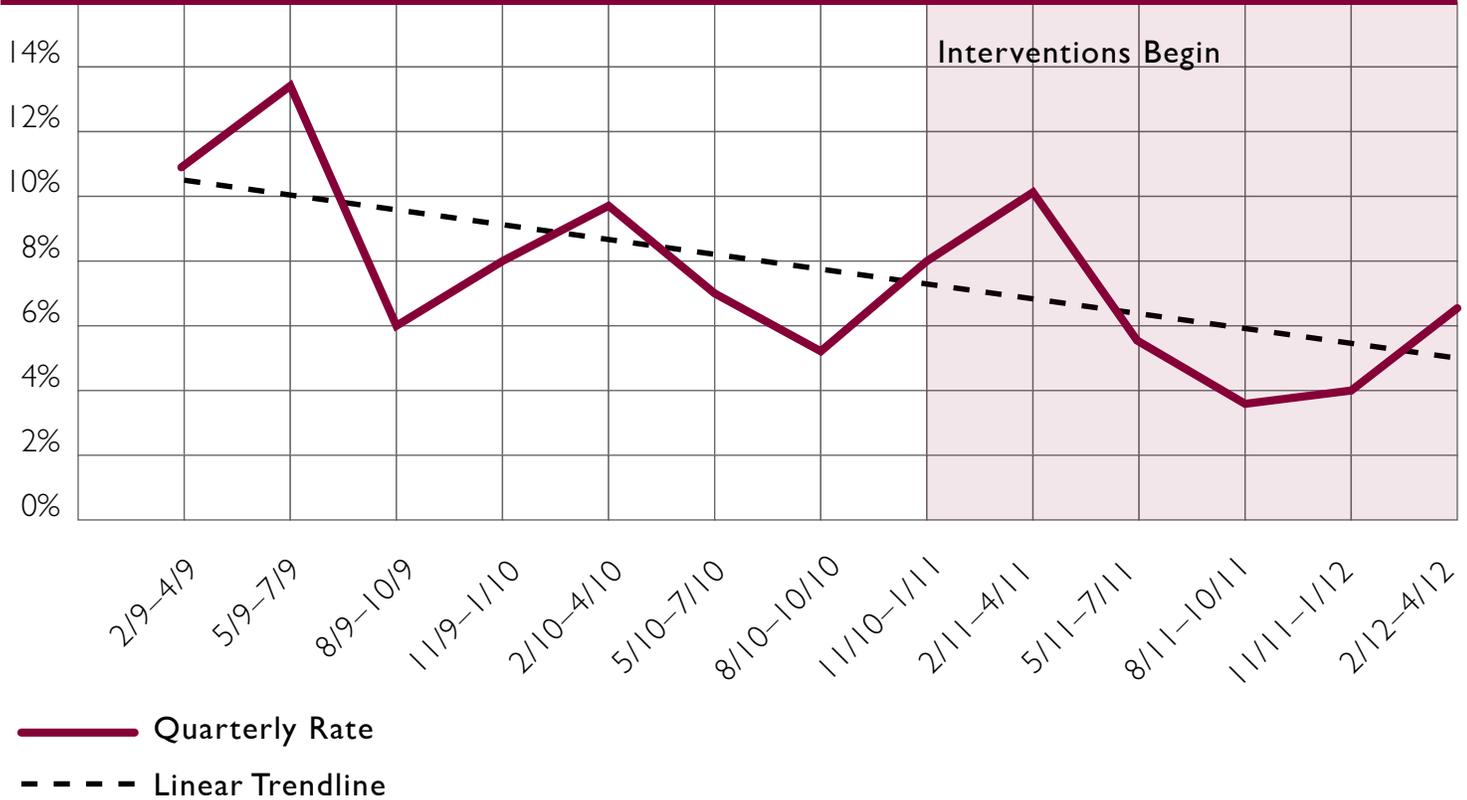
Numerator: RBC transfusions from the denominator where the patient's pre-transfusion hemoglobin was less than 8.3 g/dL or hematocrit was less than 25% during the measured quarter.

In addition to calculating the percentage for the total orders by general medicine hospitalists, the percentage not meeting criteria was calculated for each hospitalist in order to provide individual feedback.

Interventions

1. An educational campaign was presented to the group drawing attention to data indicating improved outcomes with a restrictive transfusion protocol as well as drawing attention to institutional guidelines.
2. Individual-specific feedback was presented to each individual physician by email on a quarterly basis. Physicians could then assess how they are performing as compared to their peers as well as enabling them to go back and review specific cases that were felt to be outliers based on laboratory parameters.
3. Each physician was encouraged to communicate any suggestions and potential improvements in this QI initiative with each emailing, as well as at multiple group-wide conferences.

Blood Transfusions Not Meeting Lab Criteria



Comments/Plans

This effort will continue to occur via a continuation of the educational campaign (as new physicians are coming into the group on a regular basis), regular quarterly review of data by each physician, as well as culture change within the group. Additionally, new guidelines have been published that further support a restrictive blood transfusion protocol, which should help to standardize this process.

7. Identifying Overweight and Obese Patients in Pediatric Primary Care Clinics

(3/1/11 – 2/1/12)

Specific Aim

This project aims to increase the percentage of pediatric patients with an annually documented BMI.

Problem

Obesity is reaching “epidemic” proportions among children in the U.S. Evidence-based guidelines from the American Academy of Pediatrics and the Endocrine Society recommend that as a first step in developing a systematic program to address obesity in pediatric patients, annually their height and weight should be documented and Body Mass Index (BMI) calculated and documented. The annual BMI measurement provides an empirical basis to initiate treatment and over time to evaluate treatment success and need for change in treatment. However, BMI is currently not documented annually for many of our patients. This project to document children’s BMI annually is the first in a series of linked projects to improve the clinical management of overweight and obese pediatric patients.

Patient Population

The population in this project includes all patients aged 3 years and older seen in primary care pediatrics clinics within the University of Michigan Health Care System.

Measures

For all pediatric patients aged 3 years or greater, the numerator will be the number with a documented BMI within the past 12 months, and the denominator will be all patients in that age range seen for any doctor or nurse practitioner visit during the time frame.

Interventions

The intervention objective was to have the medical assistant measure each child’s height at all visits in pediatric primary care clinic, both at health maintenance exams as well as sick and follow-up visits. Measuring weight was already a system standard for all visits. However, previously the system standard was to measure a height only at health maintenance exams. Also measuring height at sick and follow-up visits provided BMI information at all visits. This provided an annual BMI measure for most patients since most patients are seen at least annually for some type of visit.

To accomplish this objective, a physician QI site lead was identified at each clinical site. The QI site lead was responsible for working with the site medical director, administrator, physicians, and other site staff to define and implement local standard procedures for height to be measured at all visits. This was in addition to the site’s current standard procedure for measuring weight at all visits.

Site	Time Period					
	Baseline March–May 2011		Post Intervention July–Sept 2011		Post Adjustment Oct–Dec 2011	
	N	% with BMI*	N	% with BMI*	N	% with BMI*
Peds Health Center 1	2,372	85%	2,357	98%	2,413	99%
Peds Health Center 2	1,308	77%	1,275	79%	1,427	81%
Peds Health Center 3	2,243	77%	2,114	95%	2,059	95%
Peds Health Center 4	1,812	86%	1,835	98%	1,825	97%
Peds Health Center 5	1,682	91%	1,550	95%	1,609	95%
Peds Health Center 6	1,471	88%	1,366	95%	1,381	93%
Peds Health Center 7	1,134	73%	1,097	90%	1,106	94%
Peds Health Center 8	595	81%	599	88%	612	85%
Peds Health Center 9	1,198	85%	1,199	90%	1,233	96%
General Peds Overall	13,815	83%	13,374	93%	13,665	94%

Comments/Plans

The next intervention will be to utilize the data on BMI to target appropriate counseling and education for children with elevated BMI. Sites that continue to fall below the benchmark will continue to work on the identified barriers to obtaining a BMI.

8. PM&R Lean Initiative to Improve Patient Care Delivery

(6/1/11 – 11/1/11)

Specific Aim

Reduce delay in patient scheduling, optimize clinic visit time and avoid delays in charge entry.

Problem

1. Excessive delay in scheduling patient visits
2. Unnecessary extension of patient visits
3. Delay in charge entry (PEF)

Patient Population

Patients seen by the department of physical medicine and it's subspecialties at the Burlington Office Complex.

Measures

% new patients scheduled and seen within four weeks:
Denominator is total # of patients scheduled. Numerator is # of these patients seen within four weeks.

% of calls to the call center that are answered within 3 minutes:
Denominator is # of calls to the call center. Numerator is # of these calls that are answered within 3 minutes.

% of patient appointments completed within their scheduled time:
Denominator is number of patient appointments scheduled. Numerator is # of these appointments that are completed within their scheduled time. This can be calculated separately for new patients and for returning patients.

% of PEFs that are entered within three days:
Denominator is total number of PEF entered. Numerator is # of these PEFs entered within three days.

Interventions

1. Modify/add physician clinics
2. Restructure our prescription refill and call system for better telephone efficiency
3. Provide physician awareness of the importance of:
 - a. maintaining appropriate clinic visit time and
 - b. charge entry

Time Period	New patients scheduled/seen within 4 weeks	Calls answered within 3 min.	Appointments completed within scheduled time		PEFs entered within 3 days
			New Patients	Revisits	
Baseline 6/11–7/1/11	32% of 299	< 68% of 4,989	23% of 328	14% of 1,215	91% of 1,502
Post-Intervention 8/1–9/1/11	47% of 464	78% of 5,239	48% of 418	40% of 1,091	94% of 2,237
Post-Adjustment 10/1–11/1/11	61% of 438	84% of 5,489	51% of 419	40% of 1,458	91% of 2,159

Comments/Plans

Ongoing reviews of performance review will occur at monthly physician meetings, implementing recommended process changes, and reviewing the results at subsequent physician meetings. Future plans include consideration of Saturday clinics to enhance access further.

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American Board of Anesthesiology
American Board of Dermatology
American Board of Emergency Medicine
American Board of Family Medicine
American Board of Internal Medicine
American Board of Medical Genetics and Genomics
American Board of Obstetrics and Gynecology
American Board of Ophthalmology
American Board of Orthopaedic Surgery
American Board of Otolaryngology
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