Gynecology-Oncology Initiative

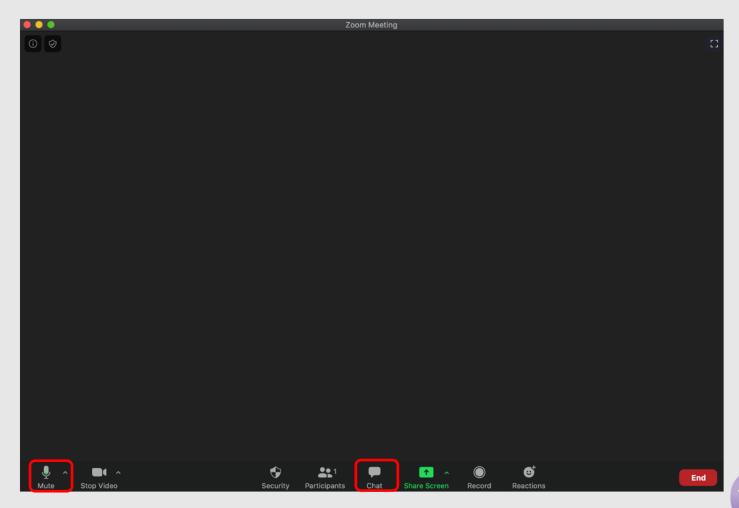
9:00 am - 12:00 pm

November 14, 2020





Virtual Meeting





Continuing Education Credits



Accreditation Statements

The University of Michigan Medical School is accredited by the Accreditation Council for Continuing Medical Education (ACCME) to provide continuing medical education for physicians. The University of Michigan Medical School designates this live activity for a maximum of **2.5** *AMA PRA Category 1 Credits*™. Physicians should claim only the credit commensurate with the extent of their participation in the activity.



Continuing Medical Education

Learning objectives

- Utilize learned gynecologic oncology quality measures
- Integrate relevant content to provide costeffective health care that does not compromise care quality
- Integrate relevant content to ensure multispecialty/multidisciplinary coordination of care
- Analyze and implement experience and improve practice

Competencies

- Practice-based learning and improvement
- Professionalism
- Systems-based practice



Agenda

Time		Presenter
9:00 am	Welcome	Vanessa Aron, BA
9:05 am	POQC Introduction Survivors Teaching Students	Amanda Itliong, MA
9:10 am	Data & Updates	Shitanshu Uppal, MD
10:00 am	10 Minute Break	
10:10 am	MiGHT Grant Update	Jennifer J. Griggs, MD, MPH
10:25 am	Ovarian Cancer Grant	Vanessa Aron, BA Audra Putt, MPH, CPH
11:05 am	10 Minute Break	
11:15 am	Next Steps/Open Discussion	Shitanshu Uppal, MD



POQC Updates/STS

Amanda Itliong, MA





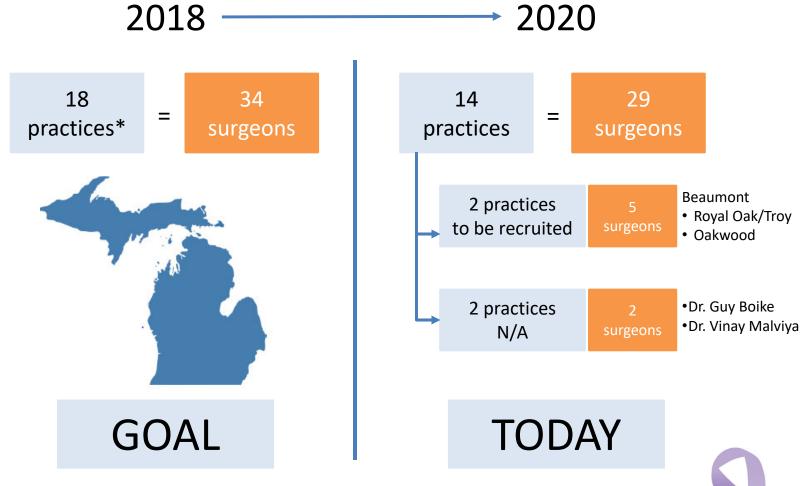


Data and Updates Shitanshu Uppal, MD





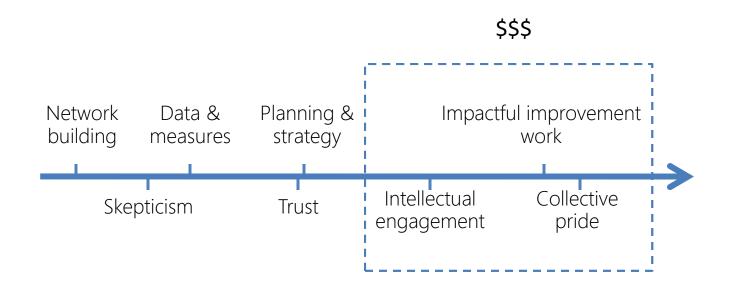
Our Group – Where Are We Today?





*Adjusted from 17 to 18 practices with addition of Midland MI Note: 2020 # of surgeons = 36; includes 1 locum & recruited surgeons into State N/A=not reachable or have indicated to MOQC not interested in joining

Evolution of Successful CQI Programs







Round 1 2020 Important Dates Charts abstracted January 7 – June 4, 2020

	Patients in Initial Therapy/Treatment (all cross cutting & disease measures)	Patients who have Died (End of Life)
Dx dates	December 2, 2018 – March 31, 2020	Dx with invasive cancer on or before March 31, 2020
First office visit	December 1, 2018 – May 31, 2020 Not required to be within office visit window (below) – occur within dx window and end of visit window date	December 1, 2018 – May 31, 2020
Two visits with provider	October 1, 2019 – May 31, 2020	Two office visits within 9 months of date of death (DoD) DoD occurred between June 1, 2018 and March 31, 2020

Notes for Graph Interpretation:

0% and no bar graph = "0" in number / "x" number in denominator No percentage (%) and no bar graph = no denominator for calculation

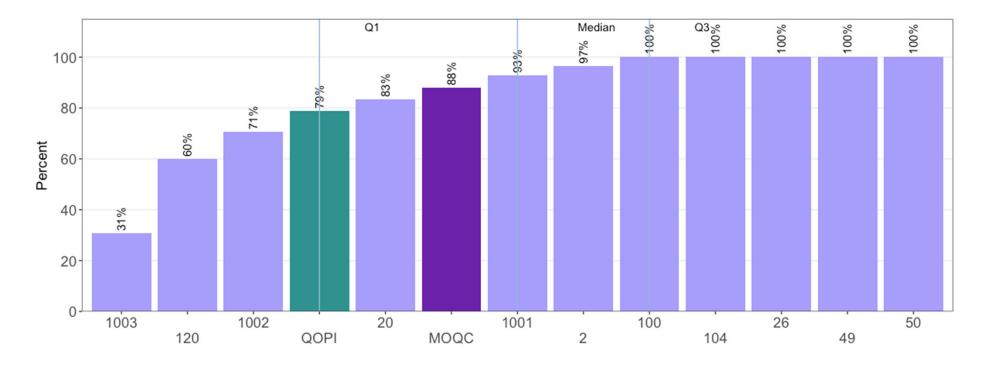




GynOnc November 2020 Meeting R1 2020

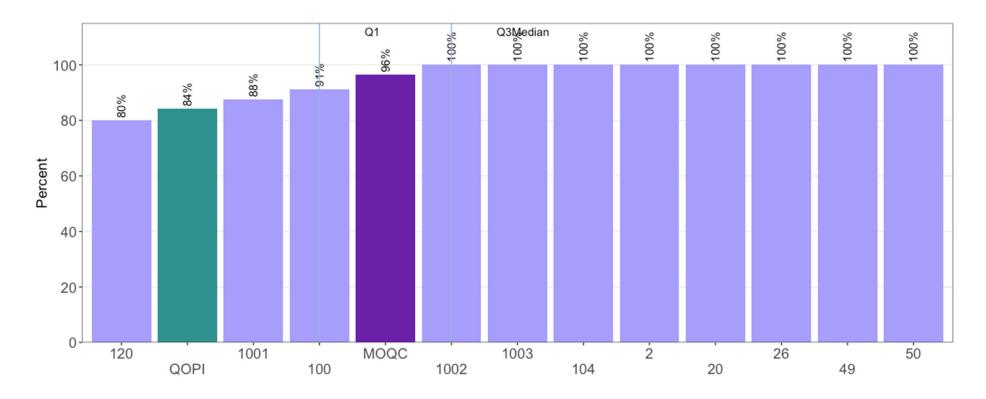
Pain addressed appropriately by second office visit and during most recent office visits

N = 182



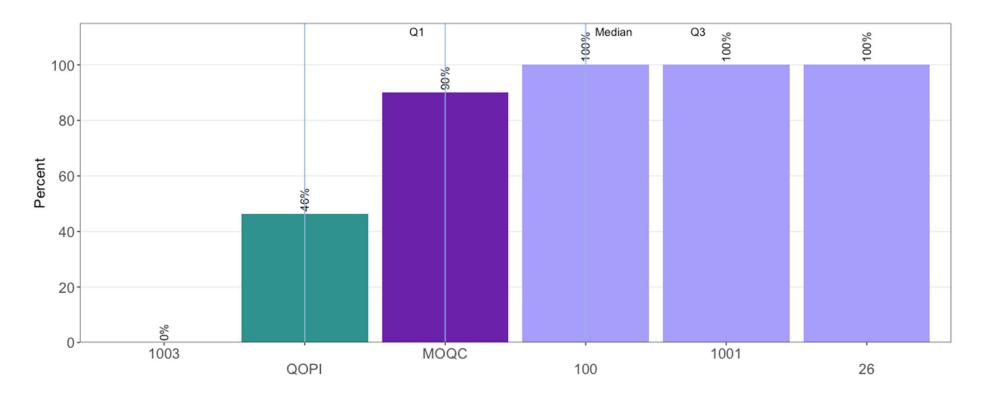
QOPI Measure CORE6e - Practice and Comparative Groups R1 2020

Signed patient consent for chemotherapy N = 141



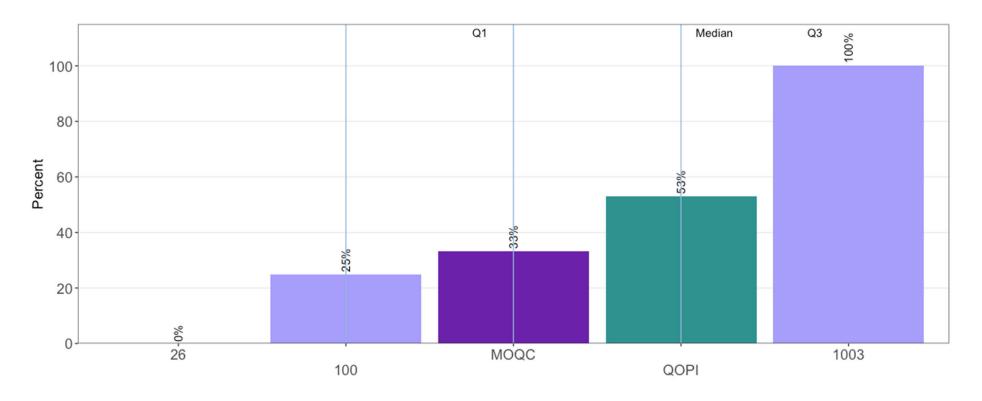
QOPI Measure CORE14 - Practice and Comparative Groups R1 2020

Tobacco cessation counseling administered or patient referred in past year N = 10



QOPI Measure CORE22bb - Practice and Comparative Groups R1 2020

Infertility risks discussed prior to chemotherapy with patients of reproductive age N = 6



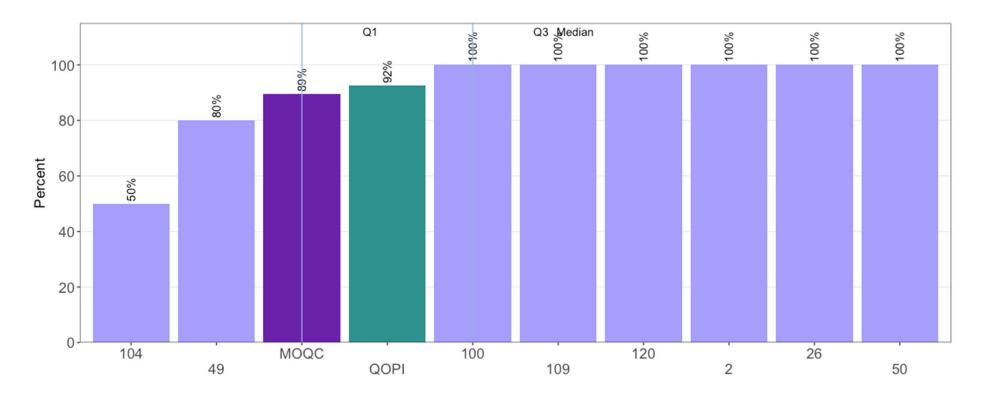
QOPI Measure SMT33 - Practice and Comparative Groups R1 2020

End of Life Measures



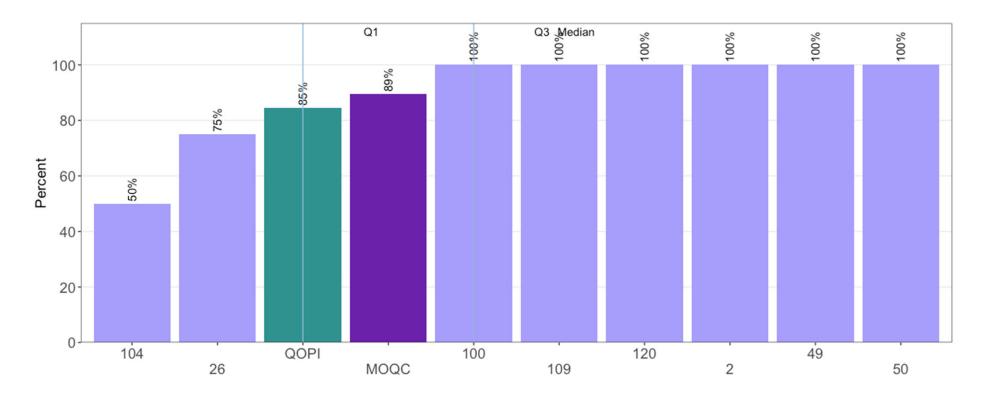


Pain addressed appropriately N = 19



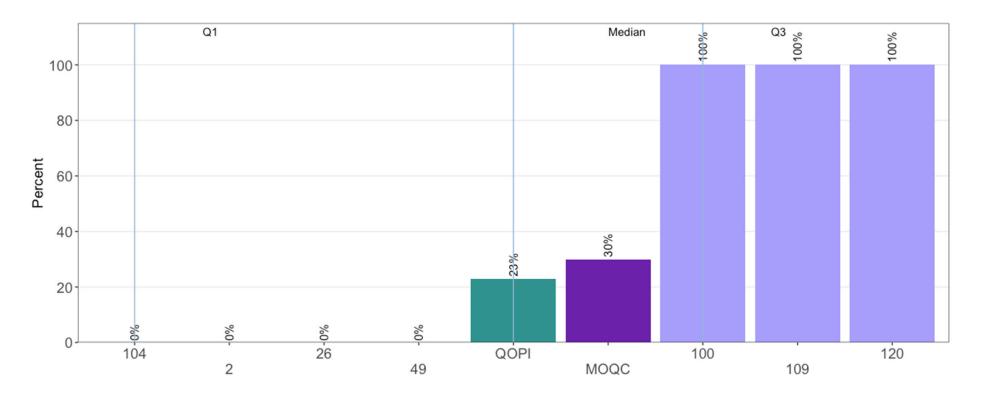
QOPI Measure EOL38 - Practice and Comparative Groups R1 2020

Dyspnea addressed appropriately N = 19



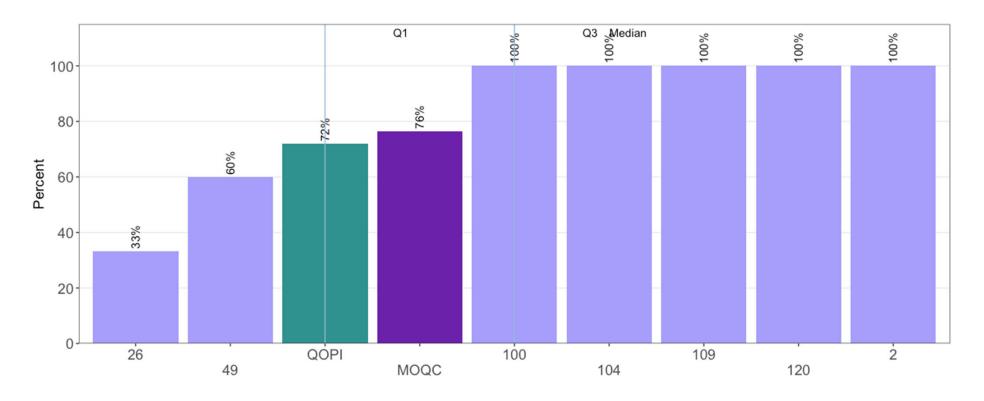
QOPI Measure EOL41 - Practice and Comparative Groups R1 2020

Hospice enrollment within 3 days of death (Lower score better) N = 10



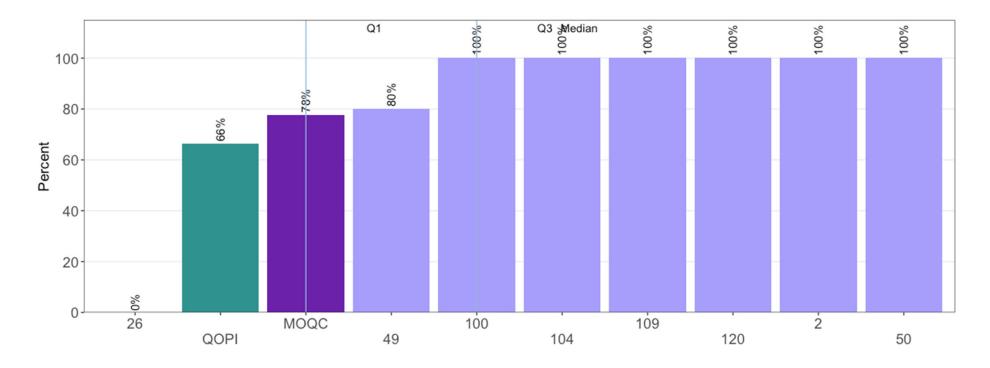
QOPI Measure EOL44 - Practice and Comparative Groups R1 2020

Hospice enrollment, or documented discussion (Combined measure 42 or 46) N = 17



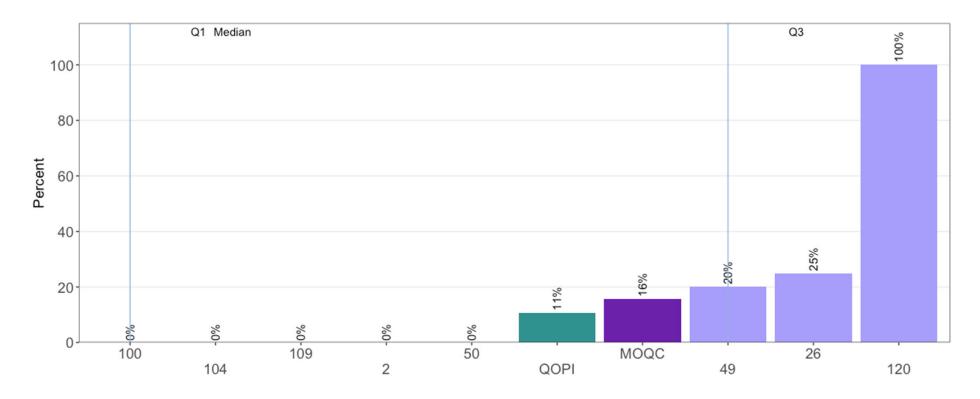
QOPI Measure EOL47a - Practice and Comparative Groups R1 2020

Palliative care referral/services, or documented discussion (Combined measure 43 or 46)
N = 18



QOPI Measure EOL47b - Practice and Comparative Groups R1 2020

Chemotherapy administered within the last two weeks of life N = 19



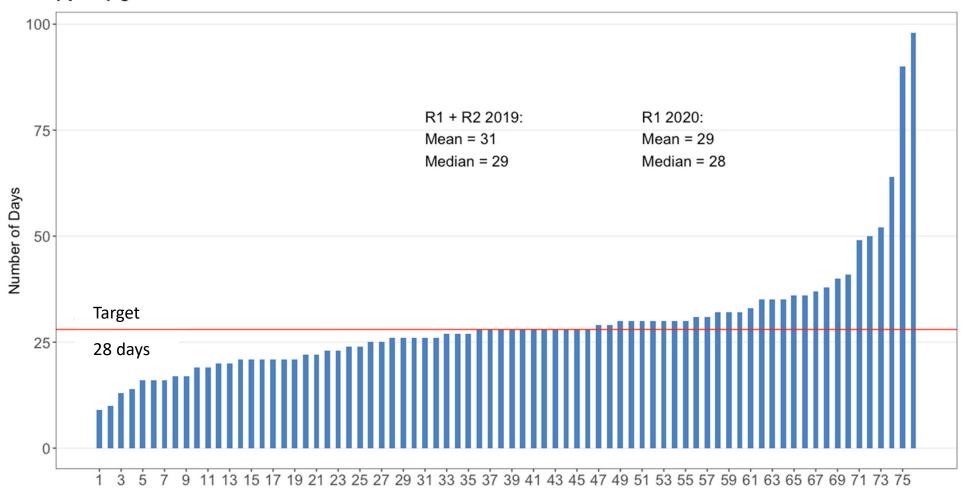
QOPI Measure EOL48 - Practice and Comparative Groups R1 2020

Gyn Onc Measures - Ovarian





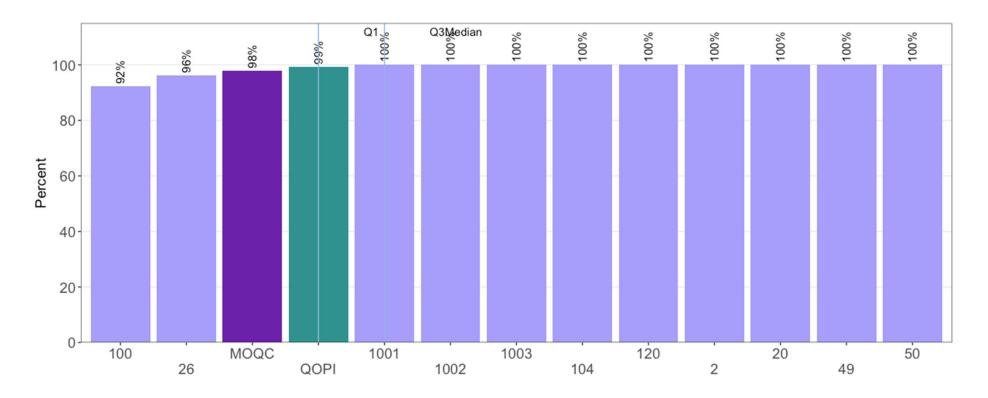
Days between Cytoreduction and 1st Day of Chemotherapy N = 76



QOPI CORE Measure GynOnc #1 Round 1 2020

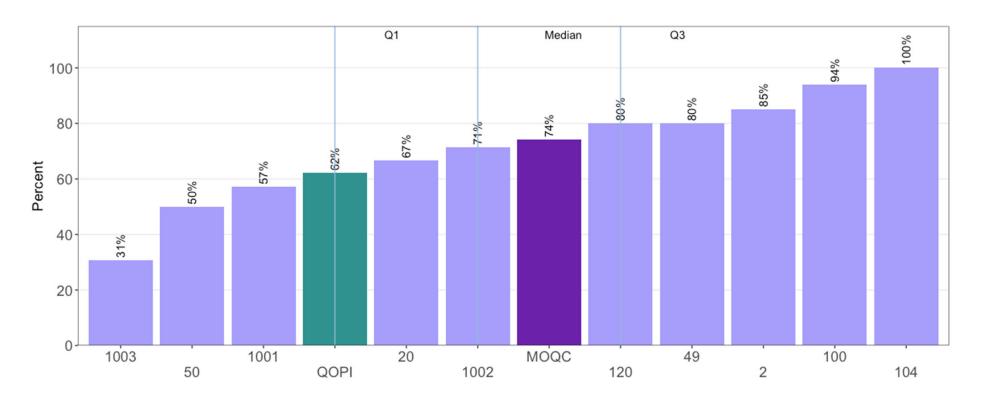
Each bar = 1 Patient

Patients with ovarian cancer referred to genetic testing/counselling N = 144



QOPI Measure GynOnc #2 - Practice and Comparative Groups R1 2020

Operative report with documentation of residual disease (Optimal/Suboptimal) N = 135



QOPI Measure GYNONC90g - Practice and Comparative Groups R1 2020

Value Based Reimbursement Summary

- Criteria for qualification
 - 1. 100% Attendance at Biannual Meetings (1 physician per practice)
 - 2. Operative report with documentation of residual disease within 48 hours of cytoreduction

• Current: 74%

• Goal: 70%

3. Days from cytoreduction to chemotherapy

• Current: 29 days

• Goal: 28 days or less





Future State

- Database Updates
 - Retired Measures
 - Consensus?
 - Abstracting all gynecologic charts, not just ovarian
 - Consensus?
 - MSQC Partnership (hysterectomy database)
- Possible VBR Measures
 - Opioid Utilization
 - Surgical Site Infections/Readmission/Reoperations etc. from MSQC
 - VTE
 - VTE Calculator
 - https://moqc.github.io/vte-calculator/



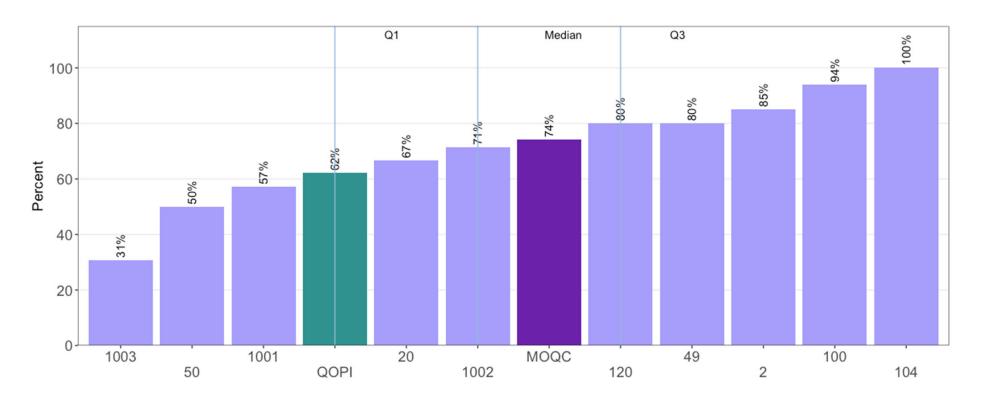


Quality Project: Operative Note



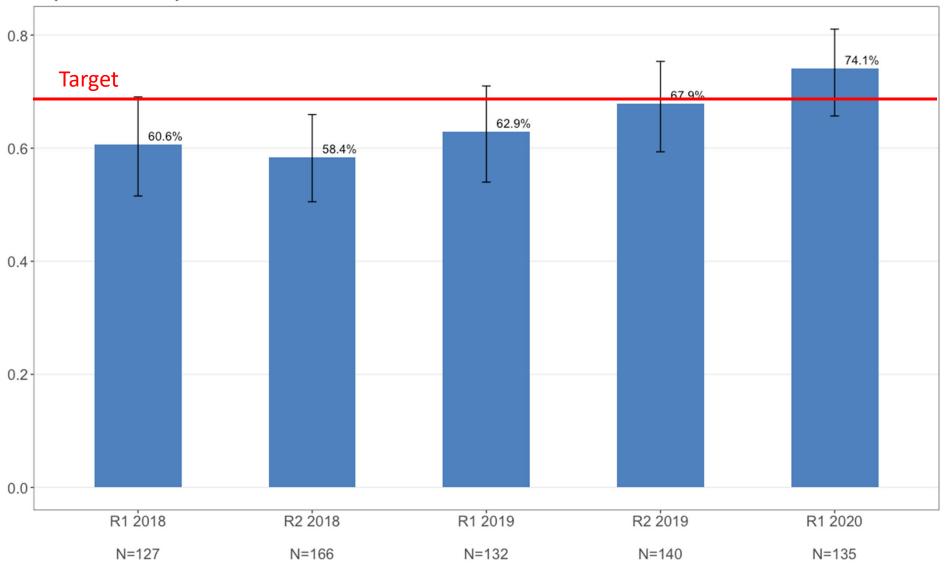


Operative report with documentation of residual disease (Optimal/Suboptimal) N = 135



QOPI Measure GYNONC90g - Practice and Comparative Groups R1 2020

Operative report with documentation of residual disease within 48 hours



QOPI CORE Measure GYNONC90g

<u>Interventions</u>

- Presentation by B Rosen, Beaumont, June 2018 → Op Note Templates
- Discussed data and re-vamped website, October 2019
- Developed templates
 - Website
 - www.moqcopnote.org
 - Lab Coat Pocket Card sent to practices, contact Vanessa Aron for additional, if desired
- Op Note Measure → VBR





www.moqcopnote.org

Standardized Operative Note

Checklist for Ovarian Cancer Operative Note Dictation



CLICK HERE for an online operative note generator

Please make sure to include the following data elements in your operative note.

- Debulking Status Primary vs. Interval Debulking
- Staging Information If available based on imaging (for example, at least stage IIIc for a patient with a biopsy-proven lesion of the omentum) Link to ovarian cancer staging
- Surgery Type Open/Robotic/Laparoscopic
- Residual Disease Status Please specify if:
 - No residual disease (R0 or no visible disease)
 - Optimally debulked (1-5 mm largest visible disease)
 - Optimally debulked (6-10 mm visible disease)
 - Sub-optimally debulked (>10 mm disease residual)
 - For suboptimally debulked patients, specify the size and location of residual disease
- Surgical Complexity Scoring Use the calculator below to get the score

 (Aletti GD, Dowdy SC, Podratz KC, Cliby WA. Relationship among surgical complexity, short-term morbidity, and overall survival in primary surgery for advanced ovarian cancer. Am J Obstet

 Gynecol. 2007;197(6):676.e1-e7.)

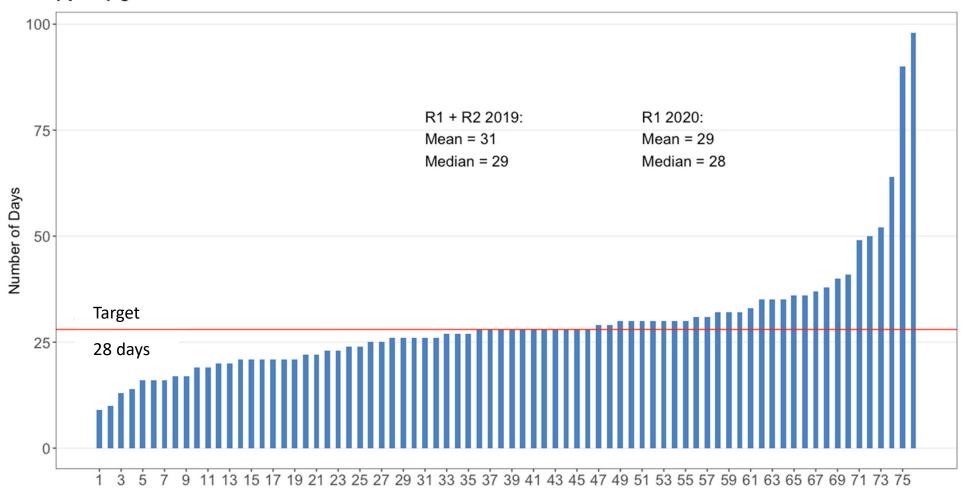


Quality Project: Days from Cytoreduction to Chemotherapy





Days between Cytoreduction and 1st Day of Chemotherapy N = 76



QOPI CORE Measure GynOnc #1 Round 1 2020

Each bar = 1 Patient

Days from Cytoreduction to Chemotherapy

- Literature
- Scheduler expectations/education
- What should we do next?



Table 1SGO guidelines for classification of urgency in gynecologic surgery.

Emergent/urgent Immediate	Semi-urgent 1–4 weeks	Non-urgent >4–12 weeks	Elective >3 months
Emergent: procedure performed without delay to preserve life or limb. Urgent: Procedure that is acutely time-sensitive and performed when the patient is medically stable.	Procedure performed in order to preserve the patient's life or prevent expected progression of disease/morbidity. Designation determined by specialty.	Progression of disease or symptoms, or readmission within 3 months is unlikely, or nonsurgical treatments available	Procedure that does not involve a medical emergency. The procedure can be delayed without meaningful disease progression or morbidity.
 Viscus perforation Closed-loop bowel or colonic obstruction Incarcerated hernia with gynecologic tumor Vaginal, uterine or pelvic hemorrhage Molar pregnancy Pelvic mass with torsion or with urinary or intestinal obstruction *CRS = cytoreductive surgery *PEG = percutaneous gastrostomy surgery *VAIN = vaginal intra-epithelial neoplasia *VIN = vulvar intraepithelial neoplasia *CIN = cervical intraepithelial neoplasia *CAH/EIN = complex atypical hyperplasia/endometrial intra-epithelial neoplasia 	 Establishment of cancer diagnosis when high suspicion exists (i.e., diagnostic laparoscopy, D&C Hysteroscopy etc.) Grade 1 endometrial cancer when hormonal therapy is contra-indicated or not possible High grade uterine cancers, all stages (i.e., epithelial and sarcoma histotypes) Cervical and vulvar cancers—surgery with curative intent Cervical and vaginal malignancies requiring radiation applicators Cervical AIS or inadequate colposcopy and concern for invasive cancer Advanced ovarian cancer, particularly interval CRS Abdominopelvic masses concerning for malignancy Symptomatic gynecologic cancer in pregnancy requiring surgery Patients with recurrent disease without non-surgical options Symptomatic patients with inoperable primary or recurrent cancer requiring palliative cancer procedures (i.e., diverting colostomy, venting PEG tubes, select exenteration) Moderate-severe anemia requiring repeated transfusion 	 Benign-appearing ovarian cysts/masses VAIN/VIN 2-3 CIN 2-3 CAH/EIN; Grade 1 endometrial cancer when hormonal therapy is not contraindicated Completion surgery for early-stage ovarian cancer Recurrent cancer requiring palliative resection 	 Risk reducing surgery for genetic predisposition to gynecologic cancer Hysterectomy for benign disease in absence of anemia Uncomplicated endometriosis Pelvic organ prolapse Urinary incontinence



Priority	Description	Examples
1	Curative therapy with a high (>50%) chance of success	Germ cell tumors Gestational trophoblastic disease
	 Adjuvant or neoadjuvant therapy which adds at least 50% chance of cure versus surgery or radiotherapy alone or treatment given at relapse 	Concurrent chemoradiation for cervical cancers
2	 Curative therapy with an intermediate (15-50%) chance of success Adjuvant or neoadjuvant therapy which adds 15-50% chance of cure versus surgery or radiotherapy alone or treatment given at relapse 	Patients with high grade serous or endometrioid ovarian cancer, particularly in patients known to have a BCRA mutation, low volume disease or good performance status
3	 Curative therapy with a low (10-15%) chance of success Adjuvant or neoadjuvant therapy which adds 10-15% chance of cure versus surgery or radiotherapy alone or treatment given at relapse Non-curative therapy with a high (>50%) chance of >1 year of life extension 	 Patients with high grade serous or endometrioid ovarian cancer, newly diagnosed or first platinum-sensitive relapse Patients with advanced, high-grade endometrial cancer
4	 Curative therapy with a low (0-15%) chance of success Adjuvant or neoadjuvant therapy which adds <10% chance of cure versus surgery or radiotherapy alone or treatment given at relapse Non-curative therapy with an intermediate (15-50%) chance of >1 year of life extension 	 Chemotherapy for cervical and endometrial cancer in first recurrence with good performance or advanced previously untreated disease Some patients with platinum-sensitive relapsed ovarian cancer
5	Non-curative therapy with a high (>50%) chance of palliation / temporary tumor control but <1 year life extension	Platinum-resistant ovarian cancer Recurrent endometrial cancer
6	Non-curative therapy with an intermediate (15-50%) chance of palliation / temporary tumor control and <1 year life extension	Chemotherapy for metastatic or recurrent cervical cancer or endometrial cancer in second recurrence



BREAK

Return in 10 minutes







MiGHT Grant

Jennifer J. Griggs, MD, MPH





Michigan Genetic Hereditary Testing (MiGHT)



Elena M. Stoeffel, MD, MPH Jennifer J. Griggs, MD, MPH Kenneth Resnicow, PhD Shitanshu Uppal, MD, MS



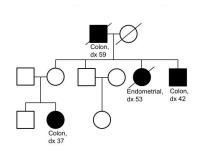


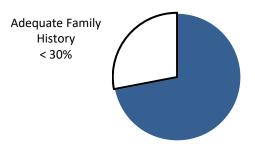








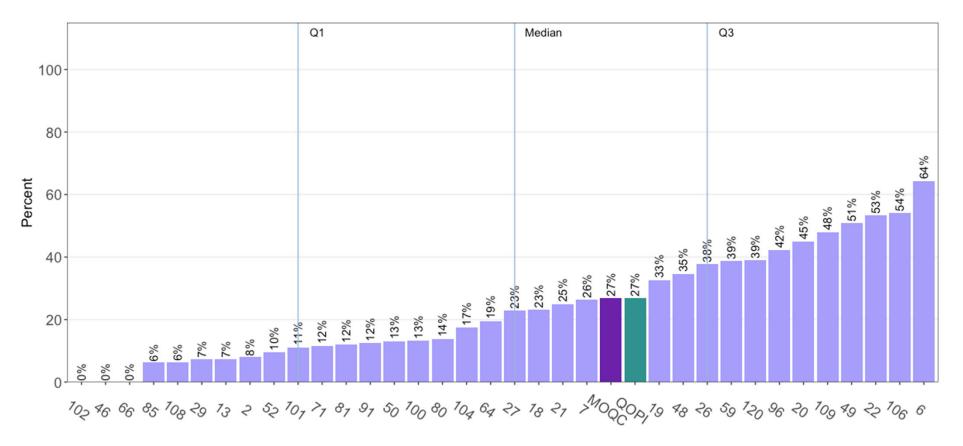






How Are We Doing?

Complete family history documented in patients with invasive cancer, N = 3040



QOPI Measure MOQC PM2 - Practice and Comparative Groups Round 2 2019

Note: Practices with no eligible cases in the denominator are not shown

Study Aims

Practice-Level

1. Improving quality of the family history in all patients with cancer

Does a tablet-based family health history survey tool with output for clinicians improve completion of the family history?

In conjunction with genetics information support for oncology teams & practices

Study Aims

Patient- and Family-Level

2. Increasing the proportion of people who get genetic testing

Can we improve the proportion of patients getting guideline-concordant genetic risk assessment & testing?

3-arm study of tailored messaging via mobile optimized web interface (app) vs genetic counseling with motivational interviewing vs usual care

Exploratory question: Can we improve the uptake of cascade testing?

VIRTUAL GENETIC COUNSELOR APP (AIM2)

CONTENT

Why should I get tested?

- Knowledge
- Myths & realities
- Attitudes
- Norms
- Motivation
- Barriers

How/Where do I get tested?

- Clinic
- Direct-to-consumer
- Payment/Insurance

Understanding Results Communication w/ Family

Cascade Testing Tips

FUNCTION

- Link to Family History Tool
- Tailored Content
- Reminders To Test
- Communicate With PCP/Oncologist
- Push Button Genetic Counselor
- Geolocated Testing And Genetic Counselor
- Searchable FAQ

Partnership with State Ovarian Cancer Grant

Vanessa Aron, BA MOQC Project Manager

Audra Putt, MPH, CPH State of Michigan – Department of Health and Human Resources





Federal Funding for Ovarian Cancer



≈ 650 New cases per year



≈ 500 Found cases



150 Case gap

Grant Projects

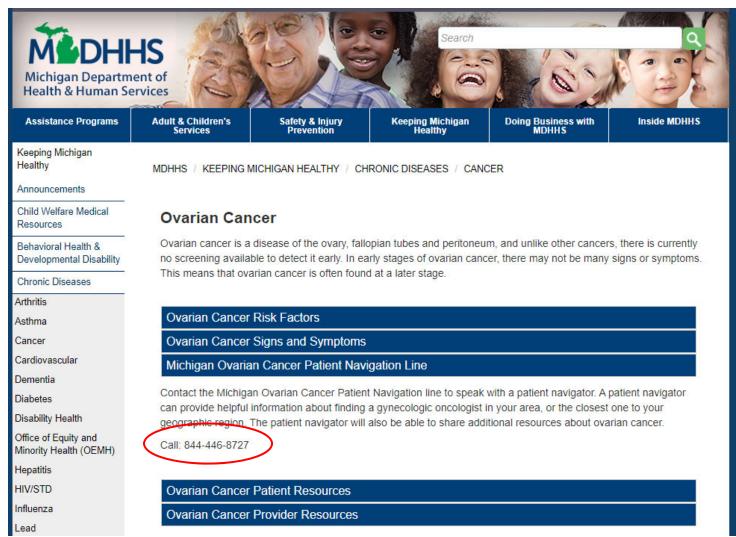






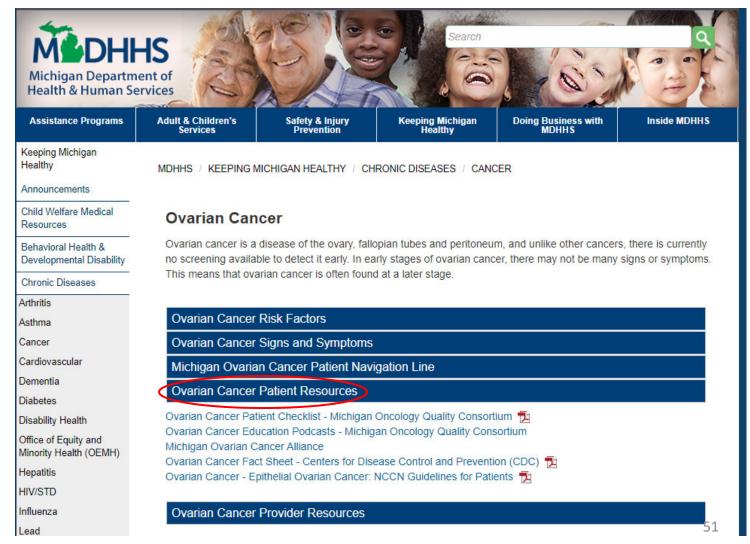
Patient Navigation Line

https://www.michigan.gov/ovariancancer

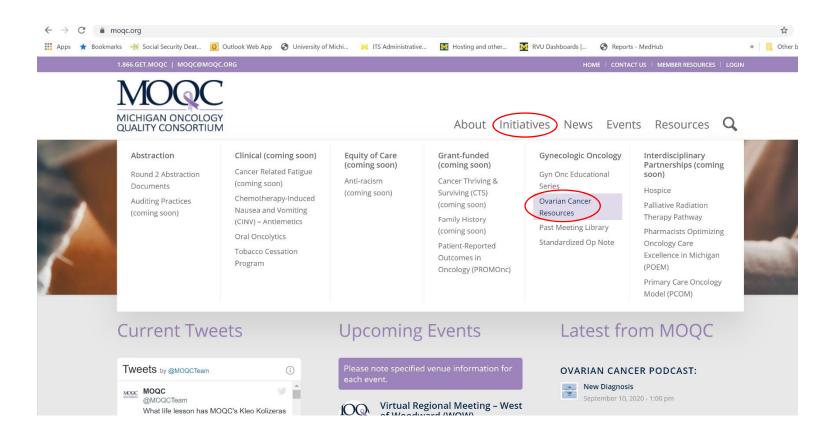


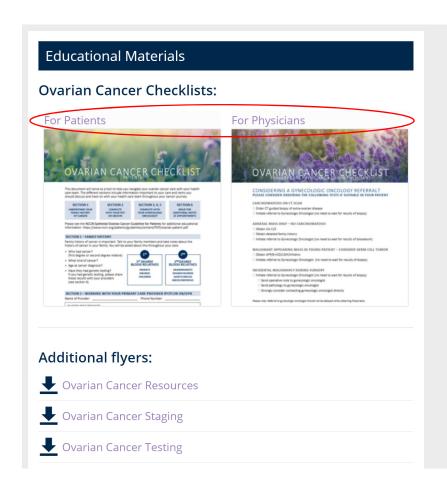
Patient Navigation Line

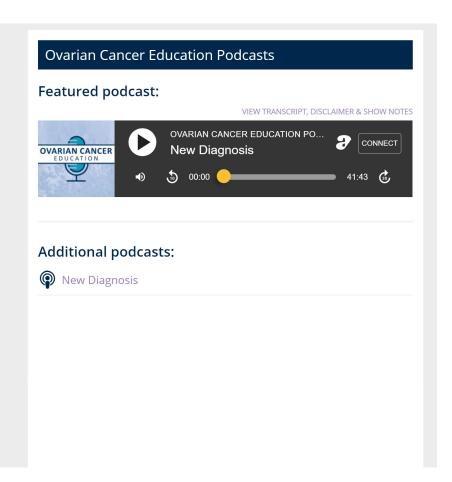
https://www.michigan.gov/ovariancancer



https://moqc.org/



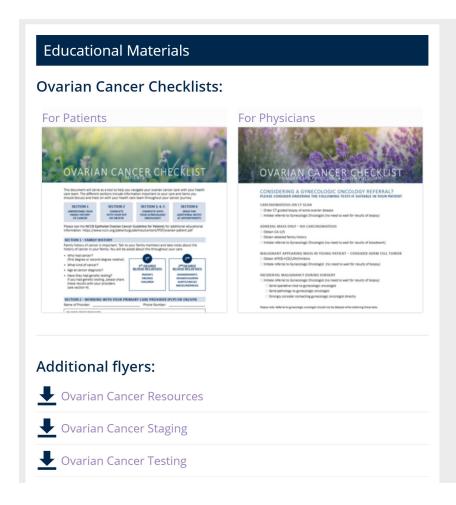


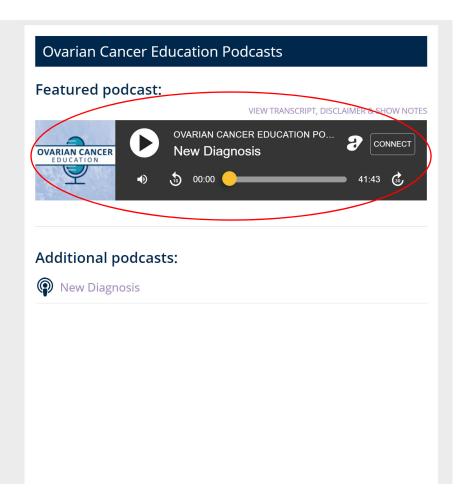






Podcast

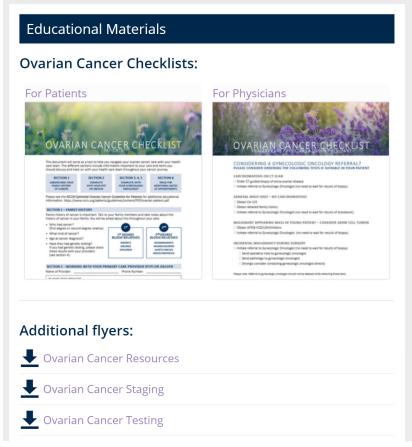


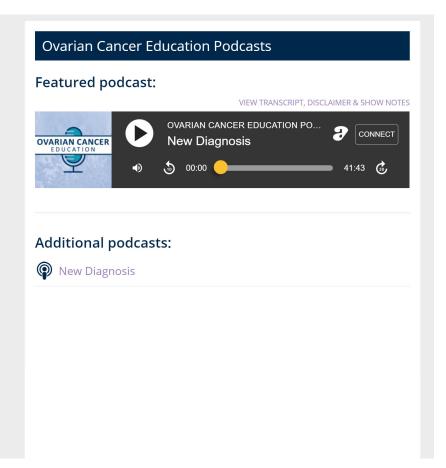


Podcast



Podcast







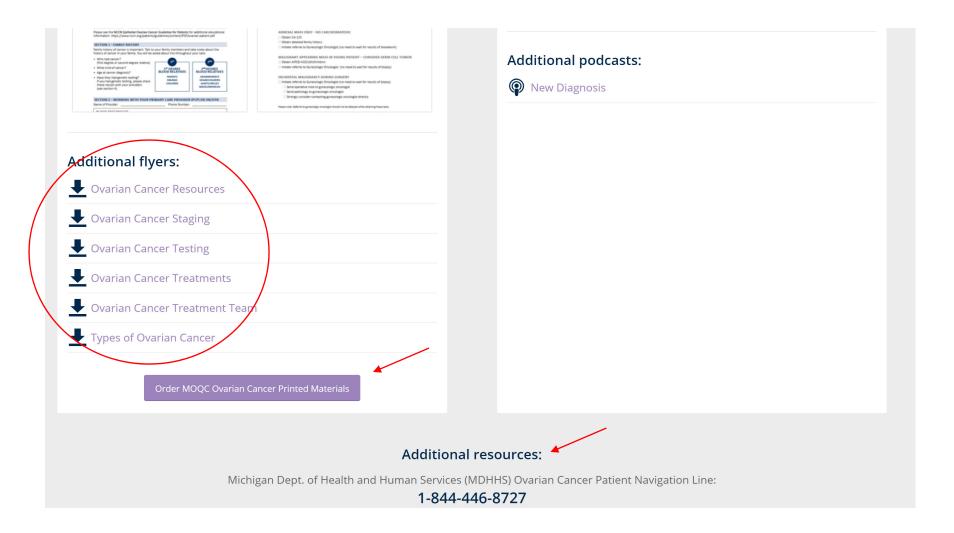




Acast







Next Steps







Thank you!

Questions?

BREAK

Return in 10 minutes







Next Steps/Open Discussion Shitanshu Uppal, MD





Next Steps

- Future MOQC Gyn-Onc Meetings
 - Saturday still best?
- POQC Recruitment
- MOQC Biannual Meeting: January 15
 - Integrated Oncology & Palliative Care
 - Keynote Speaker:

Jennifer Temel, MD

Clinical Director of Thoracic Oncology

Co-Director, Cancer Outcomes Research and Education Program

Professor of Medicine at Harvard Medical School



Discussion







