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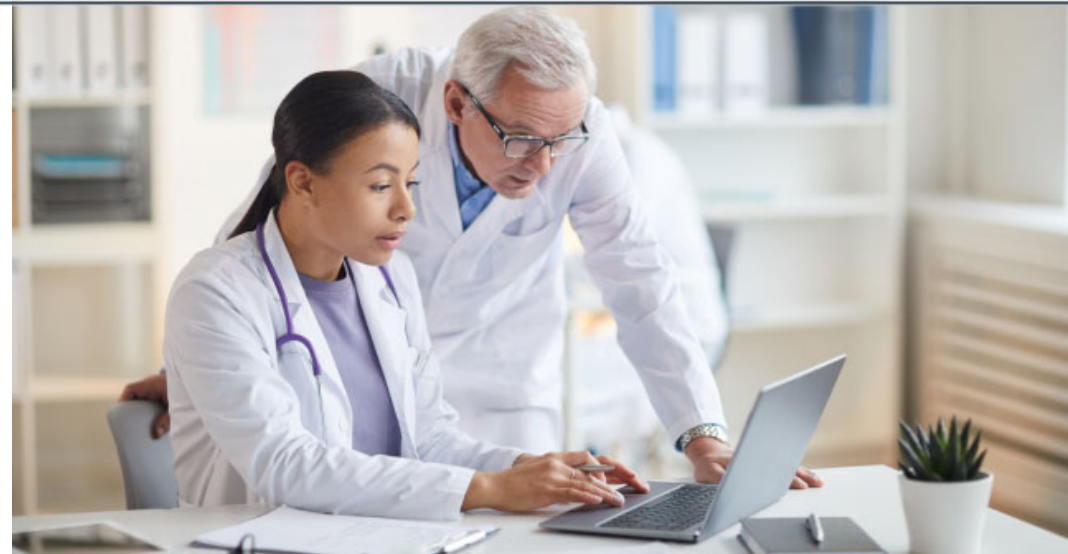
**STRONGER**TOGETHER

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for Clinical Pathology and Clinical Care Options, LLC

# Optimal Utilization of Ki67 Testing in HR-Positive/ HER2-Negative Early Breast Cancer: Education and Resources for the Oncology and Pathology Healthcare Teams

**Saturday, September 10, 2022 | 3:10 PM**  
**GLAONS 6<sup>th</sup> Annual Oncology Care Summit**  
**Los Angeles, CA**

This activity is supported by an educational grant from Lilly.



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## Program Chairs

### Joyce O'Shaughnessy, MD

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### Sara Tolaney, MD, MPH

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*Chief, Division of Breast Oncology*

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Breast Oncology Program

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## Faculty

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*Associate Chief, Hematology-  
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### **Susan Dent, MD, FRCPC**

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## Faculty

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## Faculty

### **Komal Jhaveri, MD, FACP**

*Assistant Professor of Medicine*  
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### **Michelle Melisko, MD**

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*Assistant Professor of Medicine*

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*Attending Physician, Medical*

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## Disclosure of Conflicts of Interest

Clinical Care Options, LLC (CCO) requires instructors, planners, managers, and other individuals who are in a position to control the content of this activity to disclose all financial conflicts of interest (COI) they may have with ineligible companies. All relevant COI are thoroughly vetted and mitigated according to CCO policy. CCO is committed to providing its learners with high-quality CME/CE activities and related materials that promote improvements or quality in healthcare and not a specific proprietary business interest of an ineligible company.

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**Joyce O'Shaughnessy, MD**, has disclosed that she has received consulting fees from AbbVie, Agendia, Amgen, Aptitude, AstraZeneca, Bristol-Myers Squibb, Celgene, Eisai, G1 Therapeutics, Genentech, Immunomedics, Ipsen, Jounce, Lilly, Merck, Myriad, Novartis, Ondonate, Pfizer, Puma, Prime, Roche, Seattle Genetics, and Syndax.

**Sara Tolaney, MD**, has disclosed that she has received funds for research support (paid to her institution) from AstraZeneca, Bristol-Myers Squibb, Cyclacel, Eisai, Exelixis, Genentech, Immunomedics/Gilead, Lilly, Merck, Nanostring, Nektar, Novartis, Odonate, Pfizer, Sanofi, and Seattle Genetics and consulting fees from 4D Pharma, AstraZeneca, Athenex, BeyondSpring, Bristol-Myers Squibb, Chugai, CytomX, Daiichi Sankyo, Eisai, Ellipses Pharma, Genentech, Immunomedics/Gilead, Infinity, Lilly, Mersana, Novartis, OncoPep, OncoSec, OncXerna, Pfizer, Puma, Sanofi, Samsung Bioepis, Seattle Genetics, Zymeworks, and Zentalis.

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**Adam M. Brufsky, MD, PhD**, has disclosed that he has received consultant/advisor/speaker fees from AstraZeneca, Daiichi Sankyo, Lilly, Merck, Novartis, Pfizer, and Roche.

**Susan Dent, MD, FRCPC**, has no relevant conflicts of interest to report.

**Komal Jhaveri, MD, FACP**, has disclosed that she has received consultant/advisor/speaker fees from AbbVie, AstraZeneca, Biotheranostics, Blueprint Medicines, Bristol-Myers Squibb, Genentech, Jounce Therapeutics, Lilly Pharmaceuticals, Novartis, Pfizer, Seattle Genetics, Sun Pharma Pvt Ltd, and Taiho Oncology and funds for research support from ADC Therapeutics, AstraZeneca, Clovis Oncology, Debio, Genentech, Immunomedics, Lilly, Merck/VelosBio, Novartis, Novita, Pfizer, Puma, and Zymeworks.

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**Erika P. Hamilton, MD**, has disclosed that she has received consultant/advisor/speaker fees from Arcus, Arvinas, AstraZeneca, Black Diamond, Boehringer Ingelheim, CytomX, Daiichi Sankyo, Dantari, Deciphera, Eisai, H3 Biomedicine, iTeos, Janssen, Lilly, Loxom, Merck, Mersana, Novartis, Pfizer, Puma, Relay Therapeutics, Roche/Genentech, Seattle Genetics, and Silverback and funds for research support from Abbvie, Acerta, ADC Therapeutics, Akesobio, Amgen, Aravive, ArQule, Arvinas, AstraZeneca, AtlasMedx, Black Diamond, Boehringer Ingelheim, Clovis Oncology, Compugen, Curis, CytomX, Daiichi Sankyo, Deciphera, eFFECTOR Therapeutics, Ellipses, EMD Serono, Fochon, Fujifilm, GI Therapeutics, Harpoon, H3 Biomedicine, Hutchinson MediPharma, Immunogen, Immunomedics, Incyte, InvestisBio, Jacobio, Karyopharm, Leap, Lilly, Lycera, MabSpace, MacroGenics, MedImmune, Merck, Mersana, Merus, Millennium, Molecular Templates, Myriad Genetics, Novartis, Nucana, Olema, OncoMed, Onconova, ORIC, Orinove, Pieris, PharmaMar, Pfizer, Plexxikon, Plonir, Radius Health, Regeneron, Repertoire Immune Medicines, Rgenix, Roche/Genentech, SeaGen, Sermonix, Shattuck Labs, Silverback, Stem CentRx, Sutro, Syndax, Syros, Taiho, Tapimmune, Tesaro, Treadwell Therapeutics, Verastem, Vincerx Pharma, Zenith Epigenetics, and Zymeworks.

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**Sara Hurvitz, MD, FACP**, has disclosed that she has received fees for contracted research from Ambrx, Amgen, Bayer, Cytomx, Daiichi-Sankyo, Dignitana, Genentech/Roche, Gilead, GlaxoSmithKline, Immunomedics, Lilly, MacroGenics, Novartis, OBI Pharma, Orinove, Phoenix Molecular Designs, Pfizer, Pieris, Puma, Radius, Sanofi, Seattle Genetics/Seagen, and Zymeworks.

**Virginia Kaklamani, MD, DSc**, has disclosed that she has received consultant/advisor/speaker fees from AstraZeneca, Genentech, Gilead, Novartis, Pfizer, Puma, and Seagen.

**Jane L. Meisel, MD**, has disclosed that she has received consulting fees from AstraZeneca, Genentech, GlaxoSmithKline, Lilly, Pfizer, Sanofi Genzyme, and Seagen.

**Michelle Melisko, MD**, has disclosed that she has received consultant/advisor/speaker fees from Biotheranostics and funds for research support from Novartis, OBI Pharmaceuticals, Puma, and Seattle Genetics.

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## Disclosure of Conflicts of Interest

The ***faculty*** reported the following relevant financial relationships or relationships to products or devices they have with ineligible companies related to the content of this educational activity:

**Ruth O'Regan, MD**, has disclosed that she has received consultant/advisor/speaker from Biotheranostics, Genentech, Genomic Health, Immunomedics, Lilly, MacroGenics, Novartis, Pfizer, and Puma.

**Ruta Rao, MD**, has disclosed that she has received consulting fees from Genentech, Novartis, and Sanofi.

**Laura M. Spring, MD**, has disclosed that she has received consulting fees from Novartis and Puma.

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## Staff Disclosures

The *planners/managers* reported the following relationships:

**Kemi Obajimi, PhD; Kristen Rosenthal, PhD; Gordon Kelley; Timothy A. Quill, PhD; Krista Marcello; Kelly G. Brandt, PharmD, BCOP, BCPS; and Kevin Obholz, PhD,** have no relevant conflicts of interest to report.

**Kristi Kay Orbaugh, RN, MSN, RNP, AOCN** has received speaker fees from BMS, Lilly, Sanofi, Regeneron, Gilead, Morphosys, Pfizer, AstraZeneca and DSI.

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This educational activity may contain discussion of published and/or investigational uses of agents that are not indicated by the FDA. The planners of this activity do not recommend the use of any agent outside of the labeled indications.

The opinions expressed in the educational activity are those of the faculty and do not necessarily represent the views of the planners. Please refer to the official prescribing information for each product for discussion of approved indications, contraindications, and warnings.

## Disclaimer

Learners have an implied responsibility to use the newly acquired information to enhance patient outcomes and their own professional development. The information presented in this activity is not meant to serve as a guideline for patient management. Any procedures, medications, or other courses of diagnosis or treatment discussed or suggested in this activity should not be used by healthcare professionals without evaluation of their patient's conditions and possible contraindications and/or dangers in use, review of any applicable manufacturer's product information, and comparison with recommendations of other authorities.

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## Goal

The goal of this educational curriculum is to improve the knowledge, competence, and performance of learners to optimally assess patients with HR-positive/HER2-negative early breast cancer for Ki67 and accurately report, interpret, and apply results of the testing to optimal therapy selection.

## Target Audience

This program is intended for oncologists, pathologists, histotechnologists, pharmacists, oncology nurses, and other healthcare professionals who care for patients with breast cancer.

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## Learning Objectives

At the conclusion of this activity, learners should be able to:

- Incorporate the evidence on the current use of Ki67 as a predictive and prognostic biomarker into clinical practice
- Evaluate the clinical validity, utility, and differences among Ki67 assessment techniques
- Integrate best practices for Ki67 testing methodology, standardization, and reporting
- Identify patients with high-risk features who may benefit from the addition of CDK4/6 inhibitor therapy to endocrine therapy in the adjuvant setting
- Foster collaborative discussions and workflows among oncologists, pathologists, and the multidisciplinary team on recent clinical evidence to guide personalized treatment planning for HR-positive/HER2-negative early breast cancer
- Identify patients with HR-positive/HER2-negative early breast cancer that expresses Ki67 who may be eligible for ongoing clinical trials

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## Accreditation Information



In support of improving patient care, Clinical Care Options, LLC (CCO) is jointly accredited by the Accreditation Council for Continuing Medical Education (ACCME), the Accreditation Council for Pharmacy Education (ACPE), and the American Nurses Credentialing Center (ANCC) to provide continuing education for the healthcare team.

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# Designation of Credit

## Physician Continuing Medical Education

CCO designates this live activity for a maximum of 1.0 *AMA PRA Category 1 Credit™*. Physicians should claim only the credit commensurate with the extent of their participation in the activity.

## Continuing Pharmacy Education

CCO designates this continuing education activity for 1.0 contact hour (0.1 CEUs) of the Accreditation Council for Pharmacy Education.

**UAN Live Meeting:** JA4008176-0000-22-006-L01-P

**UAN Virtual:** JA4008176-0000-22-007-L01-P

Type of Activity: Application

Upon successfully completing the activity evaluation form, transcript information will be sent to the NABP CPE Monitor Service within 60 days.

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## Designation of Credit

### Nursing Continuing Education

The maximum number of hours awarded for this Continuing Nursing Activity is 1.0 contact hour.

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Please refer to this card  
for instructions to  
receive CE credit and for  
additional resources



## Optimal Utilization of Ki67 Testing in HR-Positive/HER2-Negative Early Breast Cancer:

Education and Resources for the Oncology and  
Pathology Healthcare Teams

### EVALUATION AND REQUEST FOR CREDIT

Below are instructions for evaluating and requesting credit for the  
CME/CPE/CE-certified live meeting.

- Go to [www.clinicaloptions.com/event/Ki67ASCP22Eval](http://www.clinicaloptions.com/event/Ki67ASCP22Eval)  
to access the online evaluation and credit  
request system. You can use any computer  
or Internet interactive device or scan the  
QR code to complete the evaluation and  
credit request process.
- View and print your certificate upon  
completion of the online evaluation.



You have **30 days** before the link expires to complete the  
evaluation details and print your certificate.

### CLINICAL RESOURCES

#### Download slidesets from the live event

(available at: [www.clinicaloptions.com/event/Ki67ASCP22Slides](http://www.clinicaloptions.com/event/Ki67ASCP22Slides))

#### On-demand webcast

(available at: [www.clinicaloptions.com/event/Ki67ASCP22Webcast](http://www.clinicaloptions.com/event/Ki67ASCP22Webcast))

#### ClinicalThought

(available at: [www.clinicaloptions.com/event/Ki67ASCP22CCT](http://www.clinicaloptions.com/event/Ki67ASCP22CCT))



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## General Information

- Slides from today's presentation are now available on the CCO website

[www.clinicaloptions.com/event/Ki67ASCP22Slides](http://www.clinicaloptions.com/event/Ki67ASCP22Slides)

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**CME/CPE/CE-certified on-demand webcasts** of 2 live workshops (*coming soon*)

**Downloadable slideset** with slides from today's presentation, and updated throughout the course of the live meeting series (*available now*)

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