TARGET AUDIENCE
This activity is intended for medical oncologists, gynecologic oncologists and other healthcare providers involved in the treatment of ovarian cancer (OC).

OVERVIEW OF ACTIVITY
The pace of oncology drug development has accelerated in recent years to previously unmatched levels. Fueled by an increased understanding of the biologic underpinnings of tumor development and progression, clinical research platforms largely focused on evaluating the potential benefits of novel targeted therapeutics possessing unique mechanisms of action and safety profiles have led to improved outcomes in myriad large and rigorous clinical trials across many tumor types. The successes yielded by this rational approach to the design and evaluation of new therapies have in turn provided oncology healthcare professionals and patients with many additional and beneficial FDA-endorsed treatment options. Although this dynamic appears to be prevalent in many corners of oncology, recent advancements in the management of advanced OC have made it particularly relevant for this area of medicine. Perhaps the largest recent development in OC has been the introduction of PARP inhibitors into the therapeutic milieu, and the emergence of a handful of extremely promising data sets is stimulating significant enthusiasm in the expectation that several more novel approaches may soon become available to practicing clinicians.

These video proceedings from a CME symposium held during the 2018 ASCO Annual Meeting feature discussions with leading researchers with an expertise in ovarian cancer regarding actual cases from their practices and the published data that drive clinical decision-making for patients in those and diverse other situations. By providing information on the latest research developments and their potential application to routine practice, this activity is designed to assist medical oncologists, gynecologic oncologists and other healthcare providers with the formulation of up-to-date clinical management strategies.

LEARNING OBJECTIVES
• Appraise available guideline recommendations and consensus statements regarding the indications for genetic testing in OC, and use the results of these assessments to guide long-term treatment planning.
• Appreciate available clinical trial data with and approved indications for the use of FDA-approved PARP inhibitors for patients with OC in order to appropriately integrate these agents into routine clinical practice.
• Recognize the toxicities associated with PARP inhibitors commonly used in the care of patients with OC, and offer supportive management strategies to minimize and/or ameliorate these side effects.
• Recall the biologic rationale for and ongoing research efforts evaluating the role of PARP inhibitors in combination with chemotherapy, targeted therapy and immunotherapy, and refer appropriate patients for clinical trial participation.
• Review the mechanisms of action, emerging efficacy data and toxicity profiles of novel targeted agents and immunotherapeutic approaches under investigation in OC, and effectively prioritize clinical trial opportunities for appropriate patients.

ACCREDITATION STATEMENT
Research To Practice is accredited by the Accreditation Council for Continuing Medical Education to provide continuing medical education for physicians.

CREDIT DESIGNATION STATEMENT
Research To Practice designates this enduring material for a maximum of 1.5 AMA PRA Category 1 Credits™. Physicians should claim only the credit commensurate with the extent of their participation in the activity.

AMERICAN BOARD OF INTERNAL MEDICINE (ABIM) — MAINTENANCE OF CERTIFICATION (MOC)
Successful completion of this CME activity, which includes participation in the evaluation component, enables the participant to earn up to 1.5 Medical Knowledge MOC points in the American Board of Internal Medicine’s (ABIM) Maintenance of Certification (MOC) program. Participants will earn MOC points equivalent to the amount of CME credits claimed for the activity. It is the CME activity provider’s responsibility to submit participant completion information to ACCME for the purpose of granting ABIM MOC credit.

Please note, this program has been specifically designed for the following ABIM specialty: medical oncology.
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**HOW TO USE THIS CME ACTIVITY**

This CME activity consists of a video component. To receive credit, the participant should review the CME information, watch the video, complete the Post-test with a score of 80% or better and fill out the Educational Assessment and Credit Form located at ResearchToPractice.com/ASCOOvarian18/CME.

**CONTENT VALIDATION AND DISCLOSURES**

Research To Practice (RTP) is committed to providing its participants with high-quality, unbiased and state-of-the-art education. We assess conflicts of interest with faculty, planners and managers of CME activities. Conflicts of interest are identified and resolved through a conflict of interest resolution process. In addition, all activity content is reviewed by both a member of the RTP scientific staff and an external, independent physician reviewer for fair balance, scientific objectivity of studies referenced and patient care recommendations.

**FACULTY** — The following faculty (and their spouses/partners) reported relevant conflicts of interest, which have been resolved through a conflict of interest resolution process:

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**Advisory Committee:** AstraZeneca Pharmaceuticals LP, Genentech, Janssen Biotech Inc; **Consulting Agreements:** Abbott Laboratories, Astellas Pharma Global Development Inc, AstraZeneca Pharmaceuticals LP, Bayer HealthCare Pharmaceuticals, Bristol-Myers Squibb Company, Exelixis Inc, Genentech, GlaxoSmithKline, Incyte Corporation, Janssen Biotech Inc, Lilly, Merck, Novartis, Pfizer Inc, Pharmacyclics LLC, an AbbVie Company, Roche Laboratories Inc, Sanofi Genzyme, Takeda Oncology.


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**Hardware/Software Requirements:**
A high-speed Internet connection
A monitor set to 1280 x 1024 pixels or more
Internet Explorer 11 or later, Firefox 56 or later, Chrome 61 or later, Safari 11 or later, Opera 48 or later
Adobe Flash Player 27 plug-in or later
Adobe Acrobat Reader
(Optional) Sound card and speakers for audio

**Release date:** August 2018

**Expiration date:** August 2019
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