Cases from the Community

Clinical Investigators Provide Their Perspectives on the Use of Immune Checkpoint Inhibitors in the Management of Actual Patients with Genitourinary Cancers



A special audio program developed from a satellite CME conference held during the 2018 Genitourinary Cancers Symposium featuring expert comments on the application of emerging research to patient care

Faculty

Charles G Drake, MD, PhD Peter H O'Donnell, MD Elizabeth R Plimack, MD, MS Thomas Powles, MBBS, MRCP, MD David I Quinn, MBBS, PhD

Fditor

Neil Love, MD



Subscribe to Podcasts at ResearchToPractice.com/Podcasts



🗜 Follow us at Facebook.com/ResearchToPractice 🌛 Follow us on Twitter @DrNeilLove

Research To Practice®

Editor	Neil Love, MD
Director, Clinical Content and CPD/CME	Kathryn Ault Ziel, PhD
Scientific Director	Richard Kaderman, PhD
Editorial	Clayton Campbell
	Felix M Chinea, MD
	Marilyn Fernandez, PhD
	Adam P Hustad
	Gloria Kelly, PhD
	Kemi Obajimi, PhD
Creative Manager	Fernando Rendina
Graphic Designers	Jessica Benitez
	Tamara Dabney
	Silvana Izquierdo
Senior Manager, Special Projects	Kirsten Miller
Senior Production Editor	Aura Herrmann
Copy Editors	Rosemary Hulce
	Pat Morrissey/Havlin
	Alexis Oneca
	Kyriaki Tsaganis
Production Manager	Tracy Potter
Audio Production	Frank Cesarano
Web Master	John Ribeiro
Faculty Relations Manager	Stephanie Bodanyi, CMP
tinuing Education Administrator for Nursing	Karen Gabel Speroni, BSN, MHSA, PhD, RN
Contact Information	Neil Love, MD
	Research To Practice One Biscavne Tower
	2 South Biscayne Boulevard, Suite 3600 Miami, FL 33131
	Fax: (305) 377-9998
	Email: DrNeilLove@ResearchToPractice.com
For CME/CNE Information	Email: CE@ResearchToPractice.com

Copyright © 2018 Research To Practice. All rights reserved.

The compact disc, Internet content and accompanying printed material are protected by copyright. No part of this program may be reproduced or transmitted in any form or by any means, electronic or mechanical, including photocopying, recording or utilizing any information storage and retrieval system, without written permission from the copyright owner.

Conti

The opinions expressed are those of the presenters and are not to be construed as those of the publisher or grantors.

Participants 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 clinicians without evaluation of their patients' conditions and possible contraindications or dangers in use, review of any applicable manufacturer's product information and comparison with recommendations of other authorities.

Cases from the Community: Clinical Investigators Provide Their Perspectives on the Use of Immune Checkpoint Inhibitors in the Management of Actual Patients with Genitourinary Cancers

OVERVIEW OF ACTIVITY

The past several years have seen an explosion in the emergence of new therapies with the potential to leverage the natural ability of the human body to attack and treat cancer. Cancer immunotherapies are taking center stage at medical conferences and generating excitement all over the world. Not surprisingly, with the many advances rapidly occurring both within the field of genitourinary (GU) tumors and elsewhere, a number of vexing questions and clinical challenges are emerging simultaneously.

This CME program was developed from the proceedings of a satellite symposium held during the 2018 Genitourinary Cancers Symposium. It explores the most significant advances in the field of immunotherapy by using the perspectives of leading GU cancer experts on challenging cases and questions submitted by community oncologists to frame a discussion of how this information has aided in the refinement of current routine clinical practice and ongoing research. This activity will help medical oncologists and other allied healthcare professionals find answers to the individualized questions and concerns that they frequently encounter and in turn provide high-quality cancer care.

LEARNING OBJECTIVES

- Use patient and disease variables in addition to published research data to guide the selection of anti-PD-1/PD-L1 antibodies for patients with prostate, renal and urothelial bladder cancer.
- Compare and contrast the efficacy and safety/toxicity of approved and investigational immunotherapies for the treatment of prostate, renal cell and bladder cancer to determine the current and/or potential utility of these agents in clinical practice.
- Evaluate typical and atypical patterns of response to immune checkpoint inhibitors in an effort to identify patients who may or may not be benefiting from these agents.
- Recognize immune-related adverse events associated with immune checkpoint inhibitors, and use this information
 to develop supportive management plans for patients with GU cancers receiving these agents.

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 *AMA PRA Category 1 CreditTM*. 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 Medical Knowledge MOC point 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**. Personal information and data sharing: Research To Practice aggregates deidentified user data for program-use analysis, program development, activity planning and site improvement. We may provide aggregate and deidentified data to third parties, including commercial supporters. We do not share or sell personally identifiable information to any unaffiliated third parties or commercial supporters. Please see our privacy policy at **ResearchToPractice.com/Privacy-Policy** for more information.

HOW TO USE THIS CME ACTIVITY

This CME activity contains an audio component. To receive credit, the participant should review the CME information, listen to the audio tracks, complete the Post-test with a score of 80% or better and fill out the Educational Assessment and Credit Form located in the back of this booklet or on our website at **ResearchToPractice.com/GUCancers18/** Immunotherapy/CME.

This activity is supported by educational grants from Genentech BioOncology and Merck.

Release date: May 2018; Expiration date: May 2019

CME INFORMATION

FACULTY AFFILIATIONS



Charles G Drake, MD, PhD

Professor of Medicine Co-Director Cancer Immunotherapy Programs Columbia University Medical Center New York, New York



Thomas Powles, MBBS, MRCP, MD

Professor of Urology Oncology Director of St Bartholomew's Cancer Centre Queen Mary University of London London, United Kingdom



Peter H O'Donnell, MD Assistant Professor Department of Medicine Section of Hematology/Oncology Genitourinary Oncology Program The University of Chicago Chicago. Illinois



David I Quinn, MBBS, PhD

Medical Director, Norris Cancer Hospital and Clinics Head, GU Cancer Section Division of Cancer Medicine and Blood Diseases USC Norris Comprehensive Cancer Center Los Angeles, California



Elizabeth R Plimack, MD, MS Director, Genitourinary Clinical Research Associate Professor, Department of Hematology/Oncology Fox Chase Cancer Center Temple Health Philadelphia, Pennsylvania

COMMUNITY ONCOLOGISTS CONTRIBUTING CASES

Suzanne Cole, MD Oklahoma City, Oklahoma

Justin P Favaro, MD, PhD Charlotte, North Carolina

Philip T Glynn, MD Springfield, Massachusetts

EDITOR



Neil Love, MD Research To Practice Miami, Florida Joseph Martins, MD Tyler, Texas

Erik J Rupard, MD West Reading, Pennsylvania

CONTENT VALIDATION AND DISCLOSURES

Research To Practice (RTP) is committed to providing its participants with high-quality, unbiased and state-of-theart 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: Dr Drake — Consulting Agreements: Agenus Inc, Dendreon Pharmaceuticals Inc, Genentech BioOncology, ImmuneXcite, Janssen Biotech Inc, Lilly, Merck, NexImmune, Pierre Fabre, Roche Laboratories Inc; Contracted Research: Aduro Biotech, Bristol-Myers Squibb Company, Janssen Biotech Inc; Stockholder: Compugen, NexImmune, Potenza Therapeutics, Tizona Therapeutics Inc; Patents: AstraZeneca Pharmaceuticals LP, Bristol-Myers Squibb Company, Janssen Biotech Inc; Other: Bristol-Myers Squibb Company. Dr O'Donnell — Contracted Research: Acerta Pharma, AstraZeneca Pharmaceuticals LP/MedImmune Inc, Boehringer Ingelheim Pharmaceuticals Inc, Genentech BioOncology, Janssen Biotech Inc, Merck, Seattle Genetics; Expert Testimony: Temple Health, Trinity Health; Honoraria: Algeta ASA, American Medical Forum, Astellas Pharma Global Development Inc, AstraZeneca Pharmaceuticals LP, Genentech BioOncology, Harrison Consulting, Inovio Pharmaceuticals Inc. Janssen Biotech Inc, Kantar Health, Merck, Novartis, PAREXEL International Corporation, Quintiles, Seattle Genetics, Xcenda; Stock: Allergan Inc; Travel: Merck. Dr Plimack — Advisory Committee: AstraZeneca Pharmaceuticals LP, Bristol-Myers Squibb Company, Exelixis Inc, Genentech BioOncology, Horizon Pharma, Inovio Pharmaceuticals Inc, Lilly, Novartis, Pfizer Inc, Roche Laboratories Inc; Contracted Research: Acceleron Pharma, Agensys Inc, AstraZeneca Pharmaceuticals LP, Bristol-Myers Squibb Company, Merck, Peloton Therapeutics Inc, Pfizer Inc. Dr Powles — Consulting Agreements: AstraZeneca Pharmaceuticals LP, Bristol-Myers Squibb Company, Merck, Novartis, Pfizer Inc, Roche Laboratories Inc; Contracted Research: AstraZeneca Pharmaceuticals LP, Roche Laboratories Inc. Dr Quinn — Advisory Committee: Astellas Pharma Global Development Inc, AstraZeneca Pharmaceuticals LP, Bayer HealthCare Pharmaceuticals, Bristol-Myers Squibb Company, Dendreon Pharmaceuticals Inc, Exelixis Inc, Genentech BioOncology, Janssen Biotech Inc, Merck, Pfizer Inc, Sanofi Genzyme; Consulting Agreements: Astellas Pharma Global Development Inc, AstraZeneca Pharmaceuticals LP, Bayer HealthCare Pharmaceuticals, Bristol-Myers Squibb Company, Celgene Corporation, Dendreon Pharmaceuticals Inc, Exelixis Inc, Genentech BioOncology, Janssen Biotech Inc, Merck, Pfizer Inc, Sanofi Genzyme.

EDITOR — Dr Love is president and CEO of Research To Practice, which receives funds in the form of educational grants to develop CME activities from the following commercial interests: AbbVie Inc, Acerta Pharma, Adaptive Biotechnologies, Agendia Inc, Agios Pharmaceuticals Inc, Amgen Inc, Ariad Pharmaceuticals Inc, Array BioPharma Inc, Astellas Pharma Global Development Inc, AstraZeneca Pharmaceuticals LP, Baxalta Inc, Bayer HealthCare Pharmaceuticals, Biodesix Inc, bioTheranostics Inc, Boehringer Ingelheim Pharmaceuticals Inc, Boston Biomedical Pharma Inc, Bristol-Myers Squibb Company, Celgene Corporation, Clovis Oncology, CTI BioPharma Corp, Dendreon Pharmaceuticals Inc, Eisai Inc, Exelixis Inc, Foundation Medicine, Genentech BioOncology, Genomic Health Inc, Gilead Sciences Inc, Halozyme Inc, ImmunoGen Inc, Incyte Corporation, Infinity Pharmaceuticals Inc, Ipsen Biopharmaceuticals Inc, Janssen Biotech Inc, administered by Janssen Scientific Affairs LLC, Jazz Pharmaceuticals Inc, Kite Pharma Inc, Lexicon Pharmaceuticals Inc, Lilly, Medivation Inc, a Pfizer Company, Merck, Merrimack Pharmaceuticals Inc, Myriad Genetic Laboratories Inc, NanoString Technologies, Natera Inc, Novartis, Novocure, Onyx Pharmaceuticals, an Amgen subsidiary, Pfizer Inc, Pharmacyclics LLC, an AbbVie Company, Prometheus Laboratories Inc, Puma Biotechnology Inc, Regeneron Pharmaceuticals Inc, Sanofi Genzyme, Seattle Genetics, Sigma-Tau Pharmaceuticals Inc, Sirtex Medical Ltd, Spectrum Pharmaceuticals Inc, Taiho Oncology Inc, Takeda Oncology, Tesaro Inc, Teva Oncology and Tokai Pharmaceuticals Inc.

RESEARCH TO PRACTICE STAFF AND EXTERNAL REVIEWERS — The scientific staff and reviewers for Research To Practice have no relevant conflicts of interest to disclose.

This educational activity contains discussion of published and/or investigational uses of agents that are not indicated by the Food and Drug Administration. Research To Practice does not recommend the use of any agent outside of the labeled indications. Please refer to the official prescribing information for each product for discussion of approved indications, contraindications and warnings. The opinions expressed are those of the presenters and are not to be construed as those of the publisher or grantors.

If you would like to discontinue your complimentary subscription, please email us at **Info@ResearchToPractice. com**, call us at (800) 648-8654 or fax us at (305) 377-9998. Please include your full name and address, and we will remove you from the mailing list. Current and future integration of immune checkpoint inhibitors into the management of urothelial bladder cancer (UBC)

Tracks 1-4

to a checkpoint inhibitor Track 2 Side-effect profile and efficacy	woman with metastatic UBC experiences a complete response	Track 3	Selection of first-line therapy for patients with asymptomatic metastatic UBC		
		Track 4	Case (Dr Martins): A 68-year-old man has a minimal antitumor		
	of immune checkpoint inhibitors		response to neoadjuvant chemotherapy for bladder cancer		

Existing and emerging roles of anti-PD-1/PD-L1 antibodies in renal cell carcinoma (RCC)

Tracks 5-8

Track 5	Case (Dr Rupard): A 59-year-old man with metastatic clear cell RCC experiences disease progression	Track 7	Efficacy of immune checkpoint inhibitors in patients with RCC and brain metastases
Track 6	on first-line pazopanib Use of immune checkpoint inhibitors for metastatic RCC after disease progression on a tyrosine kinase inhibitor	Track 8	Choice of first-line therapy for patients with asymptomatic metastatic RCC

Role of immune checkpoint inhibitors in prostate cancer

Tracks 9-11

Track 9 Case (Dr Favaro): A 59-year-old Track 10 Activity of checkpoint inhibitors man with metastatic prostate in patients with CD274 (PD-L1) cancer that progresses through gene amplification multiple lines of therapy is found Track 11 Efficacy of immune checkpoint to have microsatellite instability inhibitors in patients with MSI-(MSI)-high disease and receives high prostate cancer pembrolizumab

Patterns of response to immune checkpoint inhibitors

Tracks 12-14

 Track 12
 Case (Dr Glynn): An 85-year-old man receives a checkpoint inhibitor as third-line therapy for metastatic RCC
 Track 13
 Pseudoprogression versus true progression in patients receiving immune checkpoint inhibitors

 Track 12
 Duration of therapy with anti-PD-1/PD-1 1 antibodies

Recognition and management of toxicities associated with immune checkpoint inhibitors

Tracks 15-17

Track 15	Management of dermatologic toxicities related to anti-PD-1/ PD-L1 antibodies	Track 17	Use of immune checkpoint inhibitors in patients with preexisting autoimmune disease
Track 16	Toxicity of anti-PD-1/PD-L1–anti- CTLA-4 antibody combinations		

SELECT PUBLICATIONS

Balar AV et al; IMvigor210 Study Group. Atezolizumab as first-line treatment in cisplatinineligible patients with locally advanced and metastatic urothelial carcinoma: A single-arm, multicentre, phase 2 trial. *Lancet* 2017;389(10064):64–76.

Balar AV et al. First-line pembrolizumab in cisplatin-ineligible patients with locally advanced and unresectable or metastatic urothelial cancer (KEYNOTE-052): A multicentre, singlearm, phase 2 study. *Lancet Oncol* 2017;18(11):1483-92.

Bellmunt J et al; KEYNOTE-045 Investigators. **Pembrolizumab as second-line therapy for advanced urothelial carcinoma.** *N Engl J Med* 2017;376(11):1015-26.

Bellmunt J, Barjorin DF. **Pembrolizumab for advanced urothelial carcinoma.** N Engl J Med 2017;376(23):2304.

Carbone DP et al; CheckMate 026 Investigators. First-line nivolumab in stage IV or recurrent non-small-cell lung cancer. N Engl J Med 2017;376(25):2415-26.

Danlos FX et al. Safety and efficacy of anti-programmed death 1 antibodies in patients with cancer and pre-existing autoimmune or inflammatory disease. *Eur J Cancer* 2018;91:21-9.

Escudier B et al. CheckMate 214: Efficacy and safety of nivolumab + ipilimumab (N+I) v sunitinib (S) for treatment-naïve advanced or metastatic renal cell carcinoma (mRCC), including IMDC risk and PD-L1 expression subgroups. *Proc ESMO* 2017;Abstract LBA5.

Goodman A et al. Analysis of over 100,000 patients with cancer for CD274 (PD-L1) amplification: Implications for treatment with immune checkpoint blockade. *Proc ASCO-SITC* 2018;Abstract 47.

Haanen JBAG et al; ESMO Guidelines Committee. Management of toxicities from immunotherapy: ESMO Clinical Practice Guidelines for diagnosis, treatment and follow-up. *Ann Oncol* 2017;28(Supp 4):iv119-42.

Lemery S et al. First FDA approval agnostic of cancer site — When a biomarker defines the indication. N Engl J Med 2017;377(15):1409-12.

McDermott DF et al. A phase II study of atezolizumab (atezo) with or without bevacizumab (bev) versus sunitinib (sun) in untreated metastatic renal cell carcinoma (mRCC) patients (pts). Genitourinary Cancers Symposium 2017;Abstract 431.

Minchot JM et al. Immune-related adverse events with immune checkpoint blockade: A comprehensive review. *Eur J Cancer* 2016;54:139-48.

Motzer RJ et al. IMmotion151: A randomized phase III study of atezolizumab plus bevacizumab vs sunitinib in untreated metastatic renal cell carcinoma (mRCC). Genitourinary Cancers Symposium 2018;Abstract 578.

Patel MR et al. Avelumab in metastatic urothelial carcinoma after platinum failure (JAVELIN Solid Tumor): Pooled results from two expansion cohorts of an open-label, phase 1 trial. *Lancet Oncol* 2018;19(1):51-64.

Powles T et al. Atezolizumab versus chemotherapy in patients with platinum-treated locally advanced or metastatic urothelial carcinoma (IMvigor211): A multicentre, open-label, phase 3 randomised controlled trial. *Lancet* 2017;391(10122):748-57.

Powles T et al. Efficacy and safety of durvalumab in locally advanced or metastatic urothelial carcinoma: Updated results from a phase 1/2 open-label study. *JAMA Oncol* 2017;3(9):e172411.

Powles T et al. IMmotion150: Novel radiological endpoints and updated data from a randomized phase II trial investigating atezolizumab (atezo) with or without bevacizumab (bev) vs sunitinib (sun) in untreated metastatic renal cell carcinoma (mRCC). *Proc ESMO* 2017;Abstract LBA39.

Richter MD et al. Brief report: Cancer immunotherapy in patients with preexisting rheumatologic disease: The Mayo Clinic experience. *Arthritis Rheumatol* 2018;70(3):356-60.

Rosenberg JE et al. Atezolizumab in patients with locally advanced and metastatic urothelial carcinoma who have progressed following treatment with platinum-based chemotherapy: A single-arm, multicentre, phase 2 trial. *Lancet* 2016;387(10031):1909-20.

Sharma P et al. Nivolumab in metastatic urothelial carcinoma after platinum therapy (CheckMate 275): A multicentre, single-arm, phase 2 trial. *Lancet Oncol* 2017;18(3):312-22.

Wallin JJ et al. Atezolizumab in combination with bevacizumab enhances antigen-specific T-cell migration in metastatic renal cell carcinoma. *Nat Commun* 2016;7:12624.

Weber JS et al. Safety profile of nivolumab monotherapy: A pooled analysis of patients with advanced melanoma. *J Clin Oncol* 2017;35(7):785-92.

POST-TEST

Cases from the Community: Clinical Investigators Provide Their Perspectives on the Use of Immune Checkpoint Inhibitors in the Management of Actual Patients with Genitourinary Cancers

QUESTIONS (PLEASE CIRCLE ANSWER):

- 1. Immune checkpoint inhibitors that are FDA approved in the first-line setting for cisplatinineligible patients with locally advanced or metastatic UBC include _____.
 - a. Pembrolizumab
 - b. Atezolizumab
 - c. Avelumab
 - d. All of the above
 - e. Both a and b
- Strategies for the management of dermatologic toxicities associated with immune checkpoint inhibitors include
 - a. Withholding the drug
 - b. Use of corticosteroids
 - c. Use of topical creams
 - d. All of the above

- Objective responses to immune checkpoint inhibitors ______ been observed in patients with brain metastases from RCC.
 - a. Have
 - b. Have not
- 4. What is the incidence of CD274 (PD-L1) gene amplification in patients with prostate cancer?
 - a. 10%
 - b. 4%
 - c. 0.2%
- The Phase II IMmotion150 trial compared _____ with or without bevacizumab to sunitinib for treatment-naïve advanced RCC.
 - a. Atezolizumab
 - b. Avelumab
 - c. Pembrolizumab

EDUCATIONAL ASSESSMENT AND CREDIT FORM

Cases from the Community: Clinical Investigators Provide Their Perspectives on the Use of Immune Checkpoint Inhibitors in the Management of Actual Patients with Genitourinary Cancers

Research To Practice is committed to providing valuable continuing education for oncology clinicians, and your input is critical to helping us achieve this important goal. Please take the time to assess the activity you just completed, with the assurance that your answers and suggestions are strictly confidential.

PART 1 — Please tell us about your experience with this educational activity

How would you characterize your level of knowledge on the following topics?

4 = Excellent $3 = Good$ 2		1 = Suboptimal			
	BEFORE	AFTER			
Selection of patients with newly diagnosed UBC who might benefit from treatment with pembrolizumab or atezolizumab	4321	4321			
Similarities and differences among approved anti-PD-1/PD-L1 antibodies for UBC	4321	4321			
Activity of immune checkpoint inhibitors in patients with prostate cancer	4321	4321			
Management of dermatologic toxicities associated with anti-PD-1/ PD-L1 antibodies	4321	4321			
Efficacy of immune checkpoint inhibitors in patients with RCC and brain metastases	4321	4321			
Practice Setting: Academic center/medical school Community cancer center/hospital Group practice Solo practice Government (eg, VA) Other (please specify)					
Approximately how many new patients with the following do you see per yes Prostate cancer: RCC: UBC:	ear?				
Was the activity evidence based, fair, balanced and free from commerc Yes No If no, please explain:					
 Please identify how you will change your practice as a result of complet apply). This activity validated my current practice Create/revise protocols, policies and/or procedures Change the management and/or treatment of my patients Other (please explain):					
The content of this activity matched my current (or potential) scope of Pes No If no, please explain:	practice.				
Please respond to the following learning objectives (LOs) by circling the	appropriate selec	tion:			
4 = Yes $3 = $ Will consider $2 = $ No $1 = $ Already doing $N/M = $ LO not met $N/A = $ Not applicable					
 As a result of this activity, I will be able to: Use patient and disease variables in addition to published research data guide the selection of anti-PD-1/PD-L1 antibodies for patients with prosta renal and urothelial bladder cancer. 	ite,	3 2 1 N/M N/A			
• Compare and contrast the efficacy and safety/toxicity of approved and investigational immunotherapies for the treatment of prostate, renal cell and bladder cancer to determine the current and/or potential utility of the agents in clinical practice.		3 2 1 N/M N/A			

EDUCATIONAL ASSESSMENT AND CREDIT FORM (continued)

As a result of this activity, I will be able to:

- Recognize immune-related adverse events associated with immune checkpoint inhibitors, and use this information to develop supportive management plans for patients with GU cancers receiving these agents.

Please describe any clinical situations that you find difficult to manage or resolve that you would like to see addressed in future educational activities:

.....

Would you recommend this activity to a colleague?

Yes
 No
 If no, please explain: ...

PART 2 — Please tell us about the faculty and editor for this educational activity

4 = Excellent 3	= Good	2	= Ade	quate	1 = Subo	ptimal		
Faculty	Knowled	lge of	subje	ct matter	Effective	ness	as an	educator
Charles G Drake, MD, PhD	4	3	2	1	4	3	2	1
Peter H O'Donnell, MD	4	3	2	1	4	3	2	1
Elizabeth R Plimack, MD, MS	4	3	2	1	4	3	2	1
Thomas Powles, MBBS, MRCP, MD	4	3	2	1	4	3	2	1
David I Quinn, MBBS, PhD	4	3	2	1	4	3	2	1
Editor	Knowled	lge of	subje	ct matter	Effective	ness	as an	educator
Neil Love, MD	4	3	2	1	4	3	2	1

REQUEST FOR CREDIT — Please print clearly

Name:	. Specialty:
Professional Designation: MD DO PharmD NP RN	PA Other
Street Address:	Box/Suite:
City, State, Zip:	
Telephone: Fax:	
Email:	
Research To Practice designates this enduring material for <i>Credit™</i> . Physicians should claim only the credit commensu the activity. I certify my actual time spent to complete this educational	urate with the extent of their participation in
Signature:	Date:
I would like Research To Practice to submit my CME crepoints. I understand that because I am requesting MOC creshare personally identifiable information with the ACCME and the ACCME	edit, Research To Practice will be required to
Additional information for MOC credit (required):	
Date of Birth (Month and Day Only):/ ABIM 6-Dig	git ID Number:
If you are not sure of your ABIM ID, please visit http://www.	.abim.org/online/findcand.aspx.
The expiration date for this activity is May 2019. To obt credit for this activity, please complete the Post-test, Credit Form and fax both to (800) 447-4310, or mail b Tower, 2 South Biscayne Boulevard, Suite 3600, Miar Post-test and Educational Assessment online at wy Immunotherapy/CME.	, fill out the Educational Assessment and oth to Research To Practice, One Biscayne mi, FL 33131. You may also complete the

Neil Love, MD Research To Practice One Biscayne Tower 2 South Biscayne Boulevard, Suite 3600 Miami, FL 33131 Copyright © 2018 Research To Practice. This activity is supported by educational grants from Genentech BioOncology and Merck.

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

Release date: May 2018 Expiration date: May 2019 Estimated time to complete: 1 hour

PRSRT STD U.S. POSTAGE PAID MIAMI, FL PERMIT #1317