

Cancer Q&A: Addressing Common Questions Posed by Patients with Relapsed/Refractory Multiple Myeloma

A Live Webinar for Patients, Developed in Partnership with CancerCare®

Wednesday, July 23, 2025

6:00 PM – 7:00 PM ET

Faculty

Natalie S Callander, MD

Sagar Lonial, MD, FACP

Moderator

Neil Love, MD

Faculty



Natalie S Callander, MD

Director, Myeloma Clinical Program
University of Wisconsin Carbone Cancer Center
Madison, Wisconsin



MODERATOR

Neil Love, MD

Research To Practice
Miami, Florida



Sagar Lonial, MD, FACP

Chair and Professor
Department of Hematology and Medical Oncology
Chief Medical Officer
Winship Cancer Institute
Emory University School of Medicine
Atlanta, Georgia

Survey Participants



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Chief Innovation Officer
Getz Family Professor of Cancer
Distinguished Mayo Investigator
Mayo Clinic in Arizona
Phoenix, Arizona



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Director, Center for Multiple Myeloma
Rita Kelley Chair in Oncology
Massachusetts General Hospital Cancer Center
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Harvard Medical School
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Robert Z Orlowski, MD, PhD

Florence Maude Thomas Cancer Research Professor
Department of Lymphoma and Myeloma
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Therapeutics
Vice Chair, Myeloma Translational Research
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The University of Texas MD Anderson Cancer Center
Houston, Texas



Paul G Richardson, MD

Clinical Program Leader and Director of
Clinical Research
Jerome Lipper Multiple Myeloma Center
Dana-Farber Cancer Institute
RJ Corman Professor of Medicine
Harvard Medical School
Boston, Massachusetts

Commercial Support

This activity is supported by an educational grant from GSK.

Dr Love — Disclosures

Dr Love is president and CEO of Research To Practice. Research To Practice receives funds in the form of educational grants to develop CME activities from the following companies: Aadi Bioscience, AbbVie Inc, ADC Therapeutics, Alexion Pharmaceuticals, Amgen Inc, Array BioPharma Inc, a subsidiary of Pfizer Inc, Arvinas, Astellas, AstraZeneca Pharmaceuticals LP, Aveo Pharmaceuticals, Bayer HealthCare Pharmaceuticals, BeOne, Black Diamond Therapeutics Inc, Blueprint Medicines, Boehringer Ingelheim Pharmaceuticals Inc, Bristol Myers Squibb, Clovis Oncology, Coherus BioSciences, CTI BioPharma, a Sobi Company, Daiichi Sankyo Inc, Eisai Inc, Elevation Oncology Inc, Exact Sciences Corporation, Exelixis Inc, Genentech, a member of the Roche Group, Genmab US Inc, Geron Corporation, Gilead Sciences Inc, GSK, Hologic Inc, ImmunoGen Inc, Incyte Corporation, Ipsen Biopharmaceuticals Inc, Jazz Pharmaceuticals Inc, Johnson & Johnson, Karyopharm Therapeutics, Kite, A Gilead Company, Kura Oncology, Legend Biotech, Lilly, MEI Pharma Inc, Merck, Mersana Therapeutics Inc, Mirati Therapeutics Inc, Mural Oncology Inc, Natera Inc, Novartis, Novartis Pharmaceuticals Corporation on behalf of Advanced Accelerator Applications, Novocure Inc, Nuvalent, Pfizer Inc, Pharmacyclics LLC, an AbbVie Company, Puma Biotechnology Inc, Regeneron Pharmaceuticals Inc, Rigel Pharmaceuticals Inc, R-Pharm US, Sanofi, Seagen Inc, Servier Pharmaceuticals LLC, SpringWorks Therapeutics Inc, Stemline Therapeutics Inc, Syndax Pharmaceuticals, Taiho Oncology Inc, Takeda Pharmaceuticals USA Inc, TerSera Therapeutics LLC, and Tesaro, A GSK Company.

Research To Practice CME Planning Committee Members, Staff and Reviewers

Planners, scientific staff and independent reviewers for Research To Practice have no relevant conflicts of interest to disclose.

Dr Callander — Disclosures Faculty

No relevant conflicts of interest to disclose.

Dr Lonial — Disclosures

Faculty

Advisory Committees and Consulting Agreements	AbbVie Inc, Amgen Inc, Bristol Myers Squibb, Genentech, a member of the Roche Group, GSK, Janssen Biotech Inc, Novartis, Pfizer Inc, Regeneron Pharmaceuticals Inc, Takeda Pharmaceuticals USA Inc
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Contracted Research	Bristol Myers Squibb, Janssen Biotech Inc, Novartis, Takeda Pharmaceuticals USA Inc
Stock Options/Stock — Public Companies	TG Therapeutics Inc

Dr Fonseca — Disclosures

Survey Participant

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Stock Options/Stock — Public Companies	Antengene, Caris Life Sciences

Dr Orlowski — Disclosures

Survey Participant

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Stock Options — Private Companies	Asyia Therapeutics Inc

Dr Raje — Disclosures

Survey Participant

Advisory Committees	Advisor to AstraZeneca Pharmaceuticals LP, Bristol Myers Squibb, Genentech, a member of the Roche Group, GSK, Johnson & Johnson, Pfizer Inc, Sanofi
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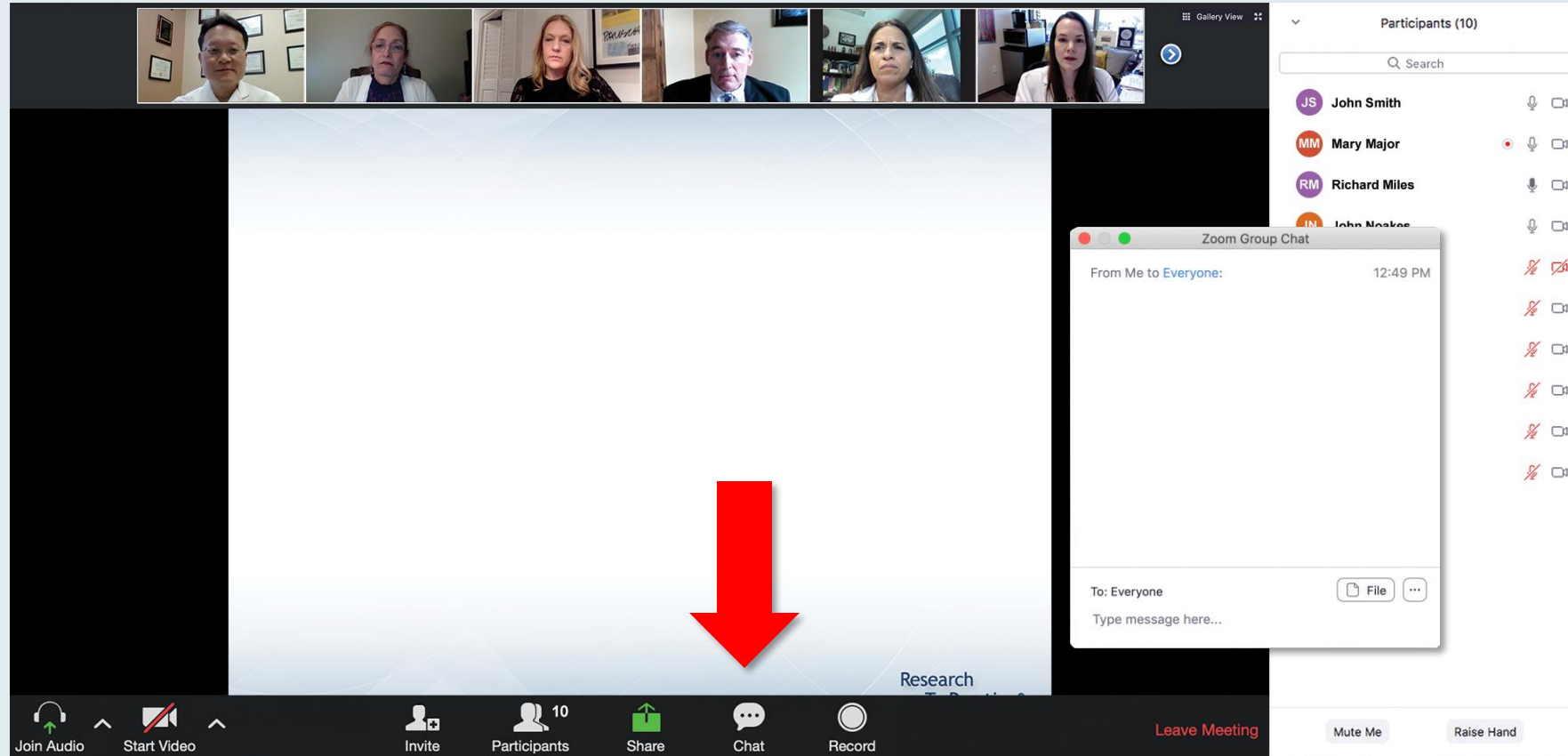
Dr Richardson — Disclosures

Survey Participant

Consulting Agreements	Bristol Myers Squibb, Celgene Corporation, GSK, Karyopharm Therapeutics, Oncopeptides, Regeneron Pharmaceuticals Inc, Sanofi
Contracted Research	Oncopeptides

This educational activity contains discussion of non-FDA-approved uses of agents and regimens. Please refer to official prescribing information for each product for approved indications.

We Encourage Patients to Submit Questions









Feel free to submit questions now before the program begins and throughout the program.

Familiarizing Yourself with the Zoom Interface

Expand chat submission box

The screenshot shows a Zoom meeting interface. At the top, there's a header bar with participant names: RTP Coordinat..., Kirsten Miller, RTP Mike Rivera, and Lisa Suarez. Below this, a slide titled "Meet The Professor Program Participating Faculty" is displayed. The slide lists six faculty members with their photos and titles. To the right of the slide, a chat window is open, showing messages from "Me to Panelists" and "Me to Panelists and Attendees". A red arrow points to the white line above the chat submission box, indicating where to drag to expand the box.

Meet The Professor Program Participating Faculty

 <p>Nancy L Bartlett, MD Professor of Medicine Koman Chair in Medical Oncology Washington University School of Medicine St Louis, Missouri</p>	 <p>Jonathan W Friedberg, MD, MMSc Samuel E Durand Professor of Medicine Director, James P Wilmot Cancer Institute University of Rochester Rochester, New York</p>
 <p>Carla Casulo, MD Associate Professor of Medicine Division of Hematology/Oncology Director, Hematology/Oncology Fellowship Program University of Rochester Wilmot Cancer Institute Rochester, New York</p>	 <p>Brian T Hill, MD, PhD Director, Lymphoid Malignancy Program Cleveland Clinic Taussig Cancer Institute Cleveland, Ohio</p>
 <p>Christopher R Flowers, MD, MS Chair, Professor Department of Lymphoma/Myeloma The University of Texas MD Anderson Cancer Center Houston, Texas</p>	 <p>Brad S Kahl, MD Professor of Medicine Washington University School of Medicine Director, Lymphoma Program Siteman Cancer Center St Louis, Missouri</p>

Chat

Me to Panelists 4:31 PM

Welcome and thank you for attending! To access the slides from today's session please use the link below.
http://images.researchtopractice.com/2021/Meetings/Slides/MTP_ToGo_CLL_2021_April1.pdf

Me to Panelists and Attendees 4:32 PM

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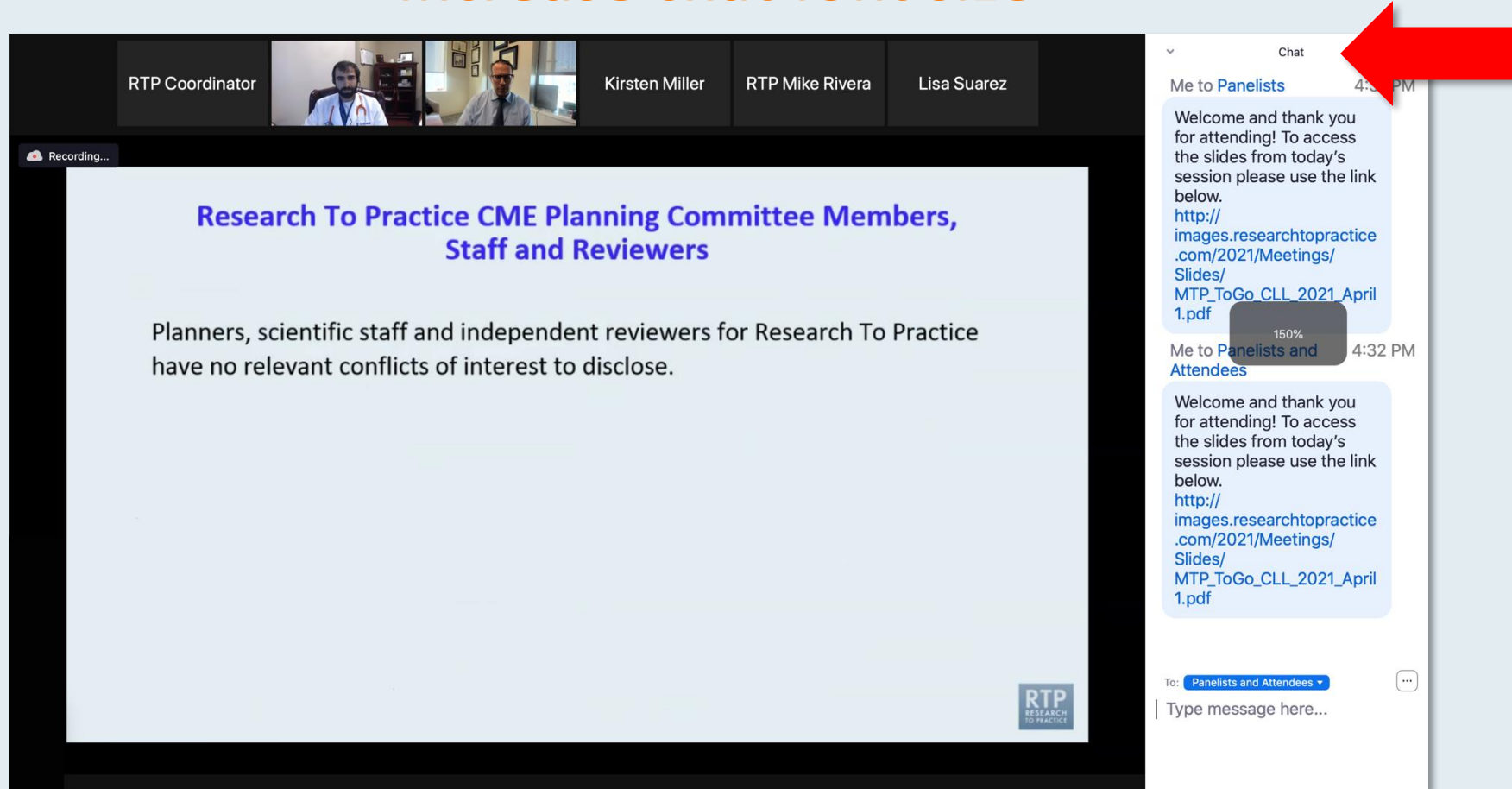
To: Panelists and Attendees

Type message here...

Drag the white line above the submission box up to create more space for your message.

Familiarizing Yourself with the Zoom Interface

Increase chat font size



The screenshot displays a Zoom meeting interface. At the top, a video bar shows participants: RTP Coordinator, Kirsten Miller, RTP Mike Rivera, and Lisa Suarez. The main area shows a presentation slide titled "Research To Practice CME Planning Committee Members, Staff and Reviewers" with the text: "Planners, scientific staff and independent reviewers for Research To Practice have no relevant conflicts of interest to disclose." The RTP logo is in the bottom right corner of the slide. On the right, the chat window is open, showing two messages from "Me to Panelists" and "Me to Panelists and Attendees". A red arrow points to the font size icon (a square with "150%") in the chat window's header.

**Press Command (for Mac) or Control (for PC) and the + symbol.
You may do this as many times as you need for readability.**

Patients in the Audience, Please Complete the Pre- and Postmeeting Surveys

The screenshot shows a Zoom meeting window. At the top, a gallery view of seven participants is visible. The main content area displays a presentation slide with the following text:

Meet The Professor
Optimizing the Selection and Sequencing of Therapy for Patients with Metastatic Gastrointestinal Cancer
Wednesday, August 25, 2022
5:00 PM – 6:00 PM EST
Faculty
Wells A Messersmith, MD
Moderator
Neil Love, MD

A "Quick Survey" pop-up window is overlaid on the slide, listing several treatment combinations with radio button options:

- ☐ Certizomab +/- dexamethasone
- ☐ Pomalidomide +/- dexamethasone
- ☐ Certizomab + pomalidomide +/- dexamethasone
- ☐ Elotuzumab + lenalidomide +/- dexamethasone
- ☐ Elotuzumab + pomalidomide +/- dexamethasone
- ☐ Daratumumab + lenalidomide +/- dexamethasone
- ☐ Daratumumab + pomalidomide +/- dexamethasone
- ☐ Daratumumab + bortezomib +/- dexamethasone
- ☐ Isaxomib + Rd
- ☐ Other

A "Submit" button is at the bottom of the survey. To the right of the main content, a "Participants (10)" list shows names and status icons. The bottom toolbar includes "Join Audio", "Start Video", "Invite", "Participants", "Share", "Chat", "Record", and "Leave Meeting".

The screenshot shows a Zoom meeting window. At the top, a gallery view of seven participants is visible. The main content area displays a presentation slide with the following text:

Regulatory and reimbursement issues aside, which treatment would you recommend for a 65-year-old patient with clear cell renal cell carcinoma (ccRCC) who has been on a tyrosine kinase inhibitor (TKI) for 3 years and follow-up 3 years later is found to have asymptomatic (PS 0)?

A "Quick Poll" pop-up window is overlaid on the slide, listing treatment options with radio button options:

- ☐ Nivolumab/ipilimumab
- ☐ Avelumab/axitinib
- ☐ Pembrolizumab/axitinib
- ☐ Pembrolizumab/lenvatinib
- ☐ Nivolumab/cabozantinib
- ☐ Tyrosine kinase inhibitor (TKI) monotherapy
- ☐ Anti-PD-1/PD-L1 monotherapy
- ☐ Other

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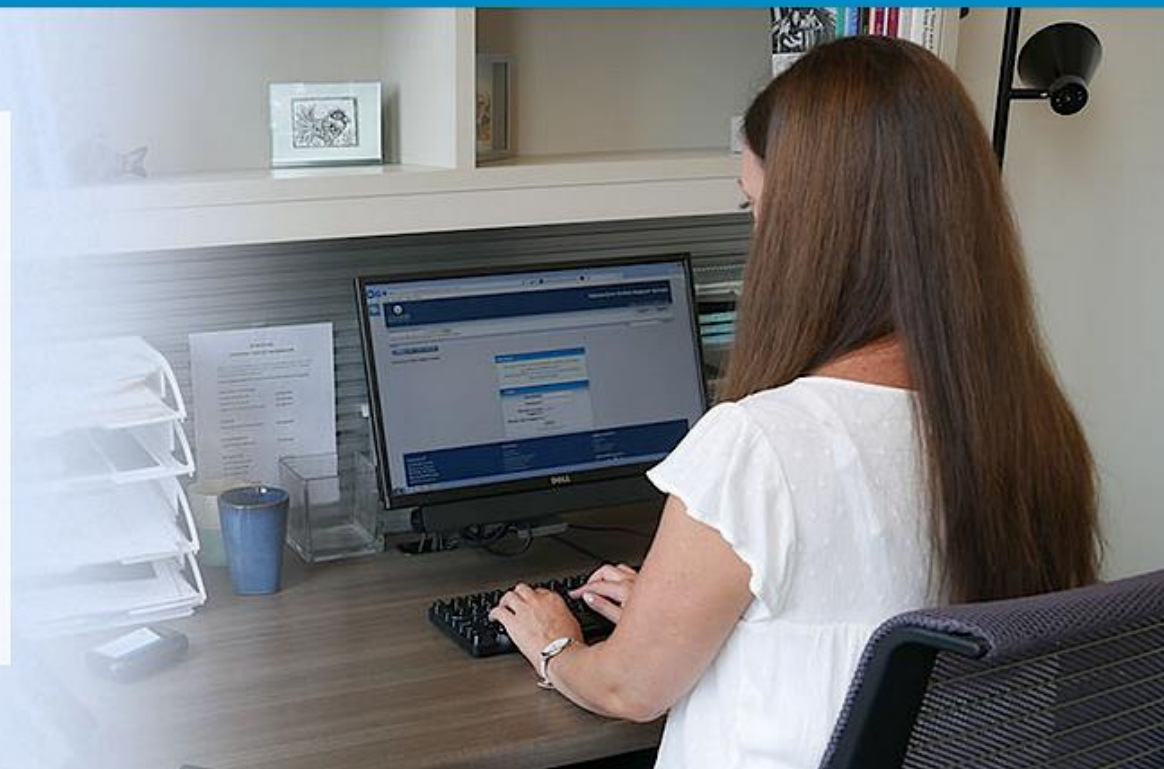
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Blood Cancers Patient Support Group

Online

Our blood cancers patient support group **provides a safe space to connect with others** coping with a blood cancer and is **led by an oncology social worker** who provides emotional and practical support.



Faculty



Natalie S Callander, MD

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Madison, Wisconsin



MODERATOR

Neil Love, MD

Research To Practice
Miami, Florida



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Distinguished Mayo Investigator
Mayo Clinic in Arizona
Phoenix, Arizona



Noopur Raje, MD

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Rita Kelley Chair in Oncology
Massachusetts General Hospital Cancer Center
Professor of Medicine
Harvard Medical School
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Robert Z Orlowski, MD, PhD

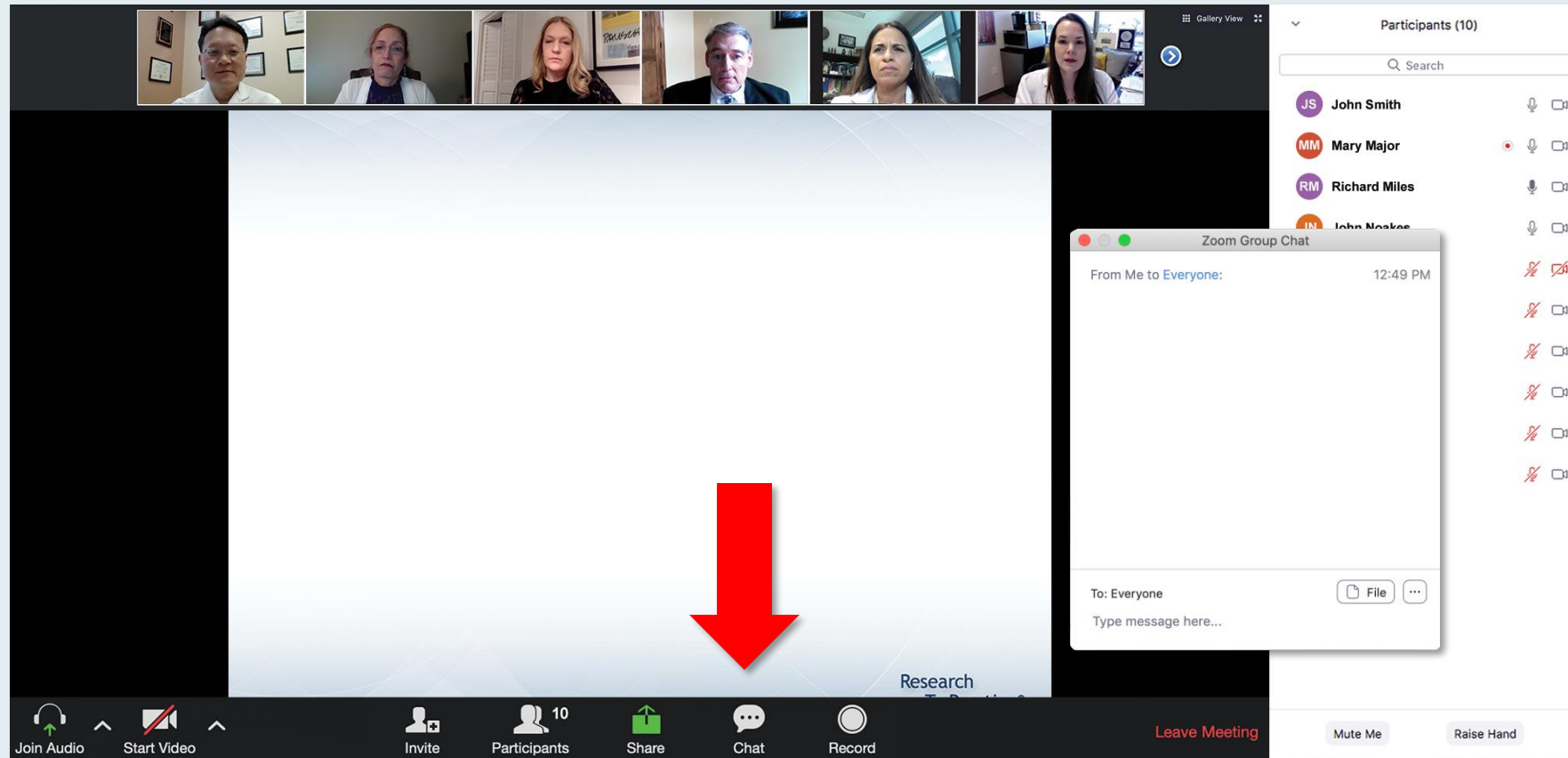
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Professor, Department of Experimental
Therapeutics
Vice Chair, Myeloma Translational Research
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Sagar Lonial, MD, FACP

Moderator

Neil Love, MD

Dr Callander — Disclosures Faculty

No relevant conflicts of interest to disclose.

Dr Lonial — Disclosures

Faculty

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Dr Fonseca — Disclosures

Survey Participant

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Dr Orlowski — Disclosures

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Dr Raje — Disclosures

Survey Participant

Advisory Committees	Advisor to AstraZeneca Pharmaceuticals LP, Bristol Myers Squibb, Genentech, a member of the Roche Group, GSK, Johnson & Johnson, Pfizer Inc, Sanofi
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Dr Richardson — Disclosures

Survey Participant

Consulting Agreements	Bristol Myers Squibb, Celgene Corporation, GSK, Karyopharm Therapeutics, Oncopeptides, Regeneron Pharmaceuticals Inc, Sanofi
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Dr Love — Disclosures

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Cancer Q&A: Addressing Common Questions Posed by Patients with Relapsed/Refractory Multiple Myeloma

A CME/MOC-Accredited Webinar Developed in Partnership with CancerCare®

Thursday, August 7, 2025

5:00 PM – 6:00 PM ET

Faculty

Natalie S Callander, MD

Sagar Lonial, MD, FACP

Moderator

Neil Love, MD

Oncology Q&A: Addressing Common Questions Posed by Patients with Metastatic Triple-Negative Breast Cancer

A Live Webinar for Patients, Developed in Partnership with the Triple Negative Breast Cancer Foundation

Wednesday, November 13, 2024

6:00 PM – 7:00 PM ET

Faculty

Lisa A Carey, MD, ScM, FASCO

Rita Nanda, MD

Moderator

Neil Love, MD

Oncology Q&A: Discussing Common Questions Posed by Patients with Metastatic Triple-Negative Breast Cancer

*A CME/MOC- and NCPD-Accredited Webinar Developed
in Partnership with the Triple Negative Breast Cancer Foundation*

Tuesday, January 7, 2025

5:00 PM – 6:00 PM ET

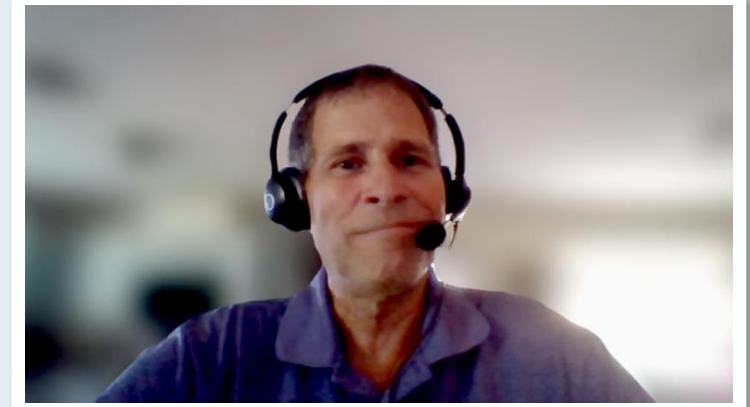
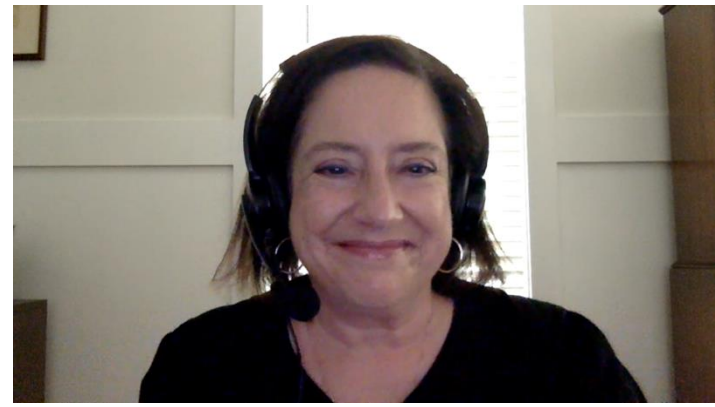
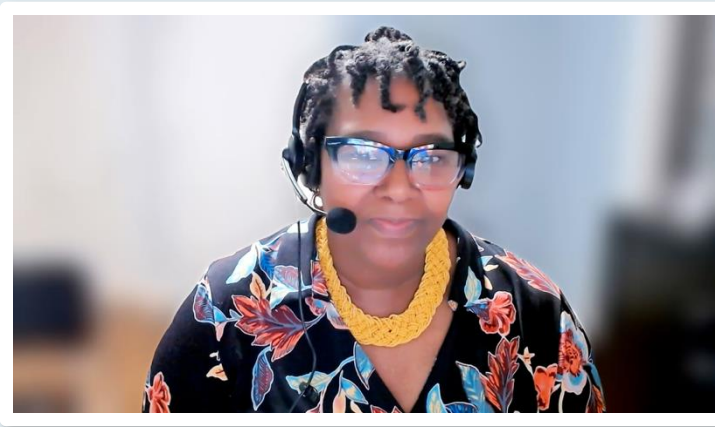
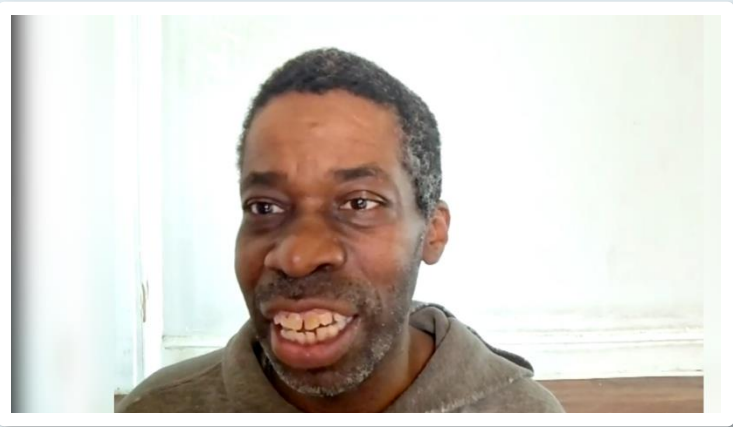
Faculty

Lisa A Carey, MD, ScM, FASCO

Rita Nanda, MD

Moderator

Neil Love, MD



Cancer Q&A

Relapsed/Refractory Multiple Myeloma

Introduction: Multiple myeloma — 2005 to 2025

Module 1: Questions from the beginning

Module 2: Choosing options

Module 3: Clinical trials

Module 4: Neuropathy

Module 5: Chimeric antigen receptor (CAR) T-cell therapy

Module 6: Bispecific antibodies

Module 7: Antibody-drug conjugates

Module 8: Interacting with the oncology team

Module 9: Other questions

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Questions from the beginning



For a patient with newly diagnosed standard-risk multiple myeloma (MM), what are the chances of cure?



Dr Callander

We still cannot cure MM for most patients, but we hope to be able to keep your myeloma controlled for many years



Dr Lonial

15%-20% can experience functional cure



Dr Fonseca

We have much better treatments so that patients can now live many years, and we are likely curing about 20%-30% of patients with today's best options



Dr Orlowski

About 10% of standard-risk patients will not experience relapse from their first line of therapy and could thus be considered cured



Dr Raje

Chance of cure for standard-risk MM is 50% to 60%



Dr Richardson

Estimated overall survival is now 10-15 years, but MM remains incurable in the longer term

For a patient who is receiving first-line treatment for standard-risk MM, on average how long will it be before a different therapy is needed?



Dr Callander

Today's therapies offer good control for many years, often 10 or more, before a treatment may have to be changed



Dr Lonial

Median time for patients receiving D-RVd induction and transplant is >10 years



Dr Fonseca

Greater than 5 years should be expected



Dr Orlowski

For a standard-risk patient, PFS is about 7 years so treatment with a second therapy will be needed in 7-8 years



Dr Raje

Close to 10 years



Dr Richardson

If quadruplet therapy is used as initial therapy, then a median of 7-8 years of PFS is expected

D-RVd = daratumumab with lenalidomide/bortezomib/dexamethasone; PFS = disease progression-free survival

Will I likely require continuous treatment for MM?



Dr Callander

This will depend on whether your MM is more aggressive or not



Dr Lonial

For now, yes



Dr Fonseca

**It is likely that after some time, 3 to 4 years,
you will be completely off therapy**



Dr Orlowski

**Continuous treatment is part of the standard for initial therapy, but emerging data suggest
a low relapse risk for patients who achieve and maintain MRD negativity 3-5 years**



Dr Raje

**Using MRD-adapted therapy, we should be able to discontinue therapy.
High-risk disease will likely require more continuous therapy**



Dr Richardson

Yes

MRD = minimal residual disease

Cancer Q&A

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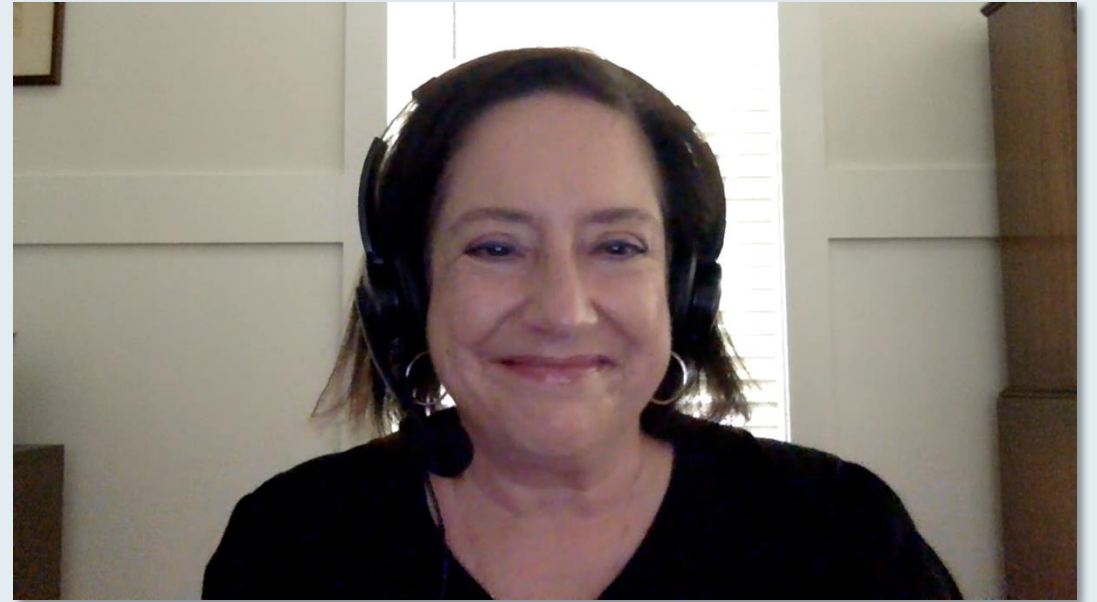
Module 6: Bispecific antibodies

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Module 9: Other questions

Choosing options



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Clinical trials



How likely is it that I will be able to participate in a clinical trial with a reasonable probability of benefitting?



Dr Callander

We will look for a clinical trial for you at every step, whether you have smoldering myeloma or have experienced relapse many times. We try to maintain a portfolio of many trials



Dr Lonial

80% or higher



Dr Fonseca

Most patients on trials benefit, and have a greater degree of scrutiny of their case



Dr Orlowski

It is likely that you will be eligible for a clinical trial at least once during your myeloma journey. The probability of benefitting will depend on several factors



Dr Raje

Very likely



Dr Richardson

Very high likelihood

What are some new therapies on the horizon that you consider to be promising for patients with R/R MM?



Dr Callander

Mezigdomide (CELMoD), trispecific antibodies, anitocabtagene autoleucel (anito-cel)



Dr Lonial

CELMoDs, p300 inhibitors



Dr Fonseca

We have so many developments including the trispecific antibodies



Dr Orlowski

CELMoDs, trispecific T-cell engagers, CAR T cells targeted against more than one antigen, and combinations of these drug classes with each other and with our current agents



Dr Raje

Cevostamab, anitocabtagene autoleucel (anito-cel), arlocabtagene autoleucel (arlo-cel), CELMoDs mezigdomide and iberdomide



Dr Richardson

CELMoDs

CELMoDs = cereblon E3 ligase modulatory drugs

Cancer Q&A

Relapsed/Refractory Multiple Myeloma

Introduction: Multiple myeloma — 2005 to 2025

Module 1: Questions from the beginning

Module 2: Choosing options

Module 3: Clinical trials

Module 4: Neuropathy

Module 5: Chimeric antigen receptor (CAR) T-cell therapy

Module 6: Bispecific antibodies

Module 7: Antibody-drug conjugates

Module 8: Interacting with the oncology team

Module 9: Other questions

Neuropathy



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Chimeric antigen receptor (CAR) T-cell therapy



What is the likelihood that my disease will respond to CAR T-cell therapy? What is the typical duration of response?



Dr Callander

Response rates, meaning obtaining a remission, are very high. Some patients may not need any other treatments for several years



Dr Lonial

90% of patients will respond to CAR T, and the average is 2-3 years of remission



Dr Fonseca

Most patients respond (>90%), and in general we think it can last for about 3 years



Dr Orlowski

Response rates are quite high and better than those seen with most other therapies available to us. The exact response rate and duration of response depends on several factors



Dr Raje

Response is 40%-80%, and the duration is 1 year to 3.6 years, depending on the type of CAR T construct used



Dr Richardson

80% chance of response with a duration of 1 to 2 years

What are the most likely side effects associated with CAR T-cell therapy?



Dr Callander

Increased risk of infections; most patients experience fever in the first week after infusion, and rarely more significant side effects called cytokine release syndrome



Dr Lonial

Infusion-related toxicity (CRS) and risk of longer-term side effects such as immune-related GI side effects and neurologic side effects



Dr Fonseca

The major and most likely toxicities include a heightened risk of infections. There are other serious toxicities, but fortunately they are rare



Dr Orlowski

Common side effects include CRS, ICANS, decreased blood counts and infections



Dr Raje

CRS, ICANS, low blood cell counts (cytopenias) and infections



Dr Richardson

CRS and ICANS

CRS = cytokine release syndrome; GI = gastrointestinal; ICANS = immune effector cell-associated neurotoxicity syndrome

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Bispecific antibodies



What is the likelihood that my disease will respond to bispecific antibody therapy?

What is the typical duration of response?



Dr Callander

About 7/10 pts respond. Those who get a deep response can be in remission for years. On average, the response is about 1 year



Dr Lonial

70%, the average is 1-2 years



Dr Fonseca

About 2 out of 3 patients will respond, but what matters more is the depth of response — those who respond very well (CR or better) can have results that are as good as CAR T cells



Dr Orlowski

Bispecific antibody response rates run from 60% to 75%, while the duration of response varies from about 12 to 24 months, depending on the bispecific used



Dr Raje

Likelihood of response is 60% with 18 to 24 months duration



Dr Richardson

70% chance of response with a duration of 12 months

CR = complete response

What are the most likely side effects associated with bispecific antibodies?



Dr Callander

Bispecifics that target BCMA are associated with a high risk of infection and patients need prophylactic antibiotics and antibodies (IVIg)



Dr Lonial

Similar to those of CAR T-cell therapy (CRS, neurologic effects, immune-related GI effects) but lower severity



Dr Fonseca

Mainly infections and the need to be very proactive with IgG replacement



Dr Orlowski

CRS, ICANS, decreased blood counts and infections



Dr Raje

CRS, ICANS, low blood cell counts (cytopenias) and infections



Dr Richardson

CRS and infections

IVIg = intravenous immunoglobulin; CRS = cytokine release syndrome; GI = gastrointestinal; IgG = immunoglobulin;
ICANS = immune effector cell-associated neurotoxicity syndrome

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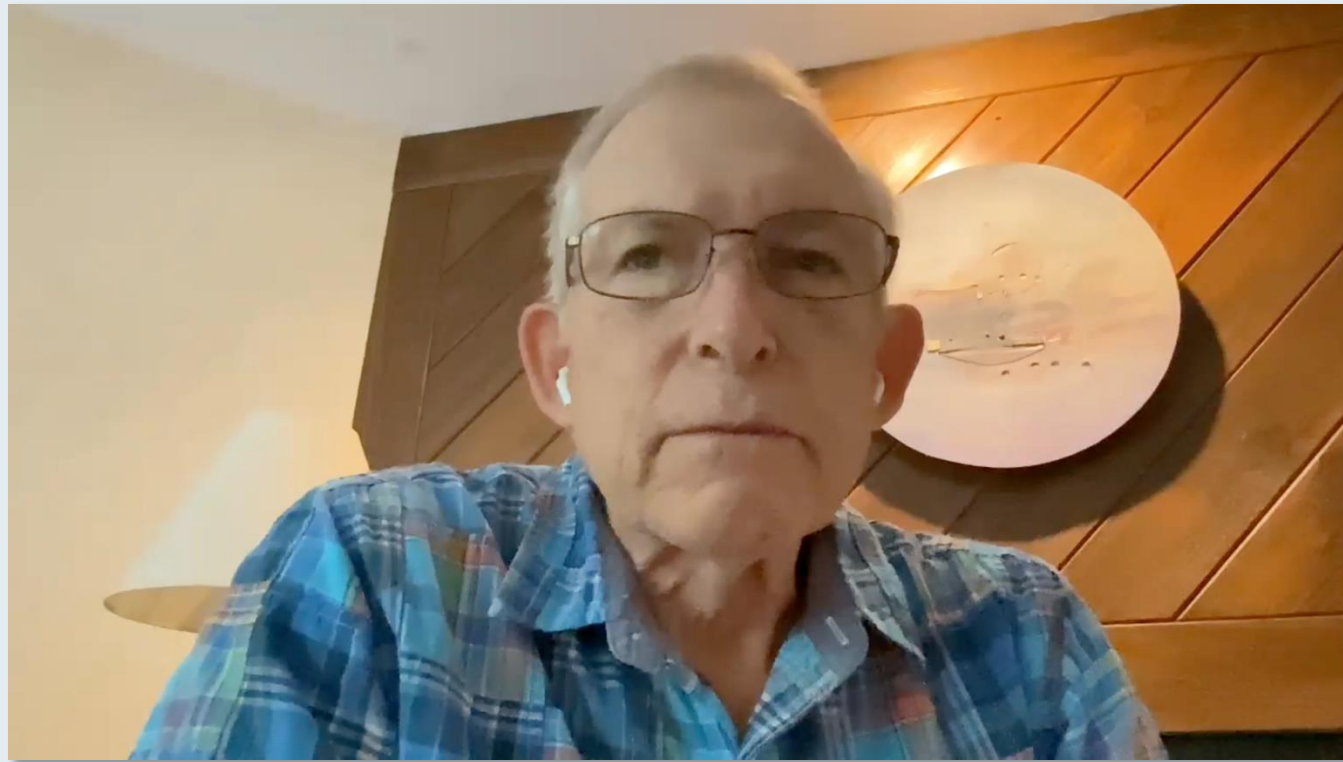
Module 6: Bispecific antibodies

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Antibody-drug conjugates



What is the likelihood that my disease will respond to belantamab mafodotin?

What is the typical duration of response?



Dr Callander

If given alone, about 1/3 of patients respond, and they tend to have long response durations, often about 1 year or more. If given in combination, most will have a response that can last several years



Dr Lonial

In combinations, chances of response are 85% or higher, and the duration is 2-3 years



Dr Fonseca

We can see responses that are very durable, sometimes close to 3 years



Dr Orlowski

The response rate for belantamab combined with bortezomib/dex was about 83%, while the average combined with pomalidomide/dex was 77%



Dr Raje

Likelihood of response is 30%, and response duration is 9 to 12 months



Dr Richardson

In combination, the chance of response is 70%-80%, and the median PFS is 3 years

dex = dexamethasone; PFS = disease progression-free survival

What are the ocular side effects associated with belantamab mafodotin? What signs or symptoms of ocular toxicity do I need to be aware of? Are the ocular side effects associated with belantamab mafodotin reversible?



Dr Callander

If belantamab is given in short intervals (eg, 3 weeks) dry eye is common, as well as blurry vision and rarely pain and inflammation



Dr Lonial

Impact on vision, changes in visual acuity, ability to read or drive, dry eyes, itchy eyes; these effects are reversible



Dr Fonseca

Symptoms range from scratchy eyes to blurred vision. With newer dosing and schedule approaches we can make the eye toxicities more manageable. They can be very uncomfortable, but know they are always reversible



Dr Orlowski

Dry eyes or excessive tearing, inflammation of the eyelids or cornea with eye pain and reduced sharpness of vision; these effects typically resolve over time



Dr Raje

Blurry vision — due to cysts in the conjunctiva; reversible with dose decrease and delays



Dr Richardson

Keratopathy, which is manageable and reversible

How are the ocular side effects associated with belantamab mafodotin prevented and managed? Will I need to see an optometrist or an ophthalmologist to monitor for ocular toxicities? How often?



Dr Callander

Side effects are managed by using a longer interval between doses, lubricating eye drops. Before every dose, we will have you see an ophthalmologist or optometrist familiar with the drug



Dr Lonial

Holding or reducing the dose or spreading out the dose to reduce future risk; monitoring by eye care professional is done once every 3-4 weeks depending on severity



Dr Fonseca

We will engage with the ophthalmology team, usually once a month, and monitor. We are able to manage these better now



Dr Orlowski

We currently do not have good prevention strategies for the ocular side effects. You would typically see an optometrist or ophthalmologist before starting belantamab and then before each next dose



Dr Raje

Side effects may be managed by decreasing belantamab dose and decreasing dose intensity



Dr Richardson

Side effects are manageable utilizing KVA monitoring, eye drops, dose reduction and schedule change; initially will need to consult eye specialist monthly x 4, then as needed

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Interacting with the oncology team



What do you recommend to your patients with R/R MM in terms of exercise, diet and nutrition and complementary treatments like acupuncture and meditation?



Dr Callander

I always advocate for exercise, even chair exercises may be helpful. I ask patients to be skeptical of supplements; acupuncture often not covered by insurance



Dr Lonial

Balanced diet, limited use of supplements, keep as fit as they are able



Dr Fonseca

Just follow your normal life



Dr Orlowski

For patients with lytic/erosive bony lesions, I recommend exercise after evaluation by orthopedics/physical therapy; I obtain a consult with a nutritionist and refer patients to our Integrative Medicine team



Dr Raje

Healthy diet with adequate hydration, daily exercise



Dr Richardson

A comprehensive and proactive approach with exercise, diet/nutrition and certain complementary therapies

At what point do you introduce discussion about palliative care with your patients with R/R MM? Do you discuss advanced directives and other issues related to end-of-life preparation?



Dr Callander

Yes, particularly before high-dose therapy, when we encourage patients to have an advanced directive



Dr Lonial

When symptoms are challenging to manage, or patients have exhausted many treatment options



Dr Fonseca

It is hard as patients have so many options. It is not unusual for MM patients to only have a few days or weeks when they enroll in hospice. I use palliative care support for symptom control as needed



Dr Orlowski

I start mentioning palliative care as an option at the first sign of refractory disease; our institutional policy is to discuss advanced directives no later than the third visit of the patient in our system



Dr Raje

Usually when performance status decreases and treatment options are narrowing



Dr Richardson

Later in the disease course, typically after 3 to 4 lines of therapy

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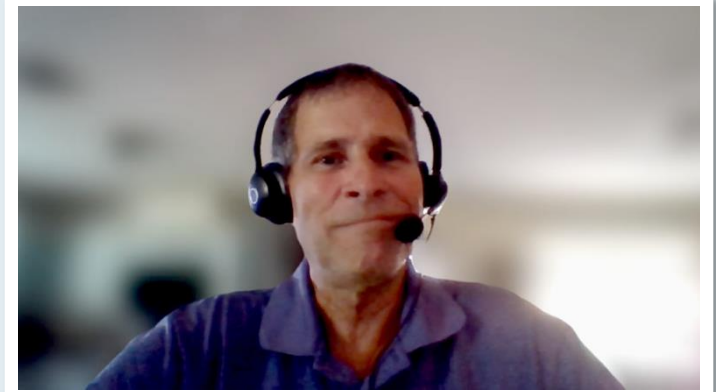
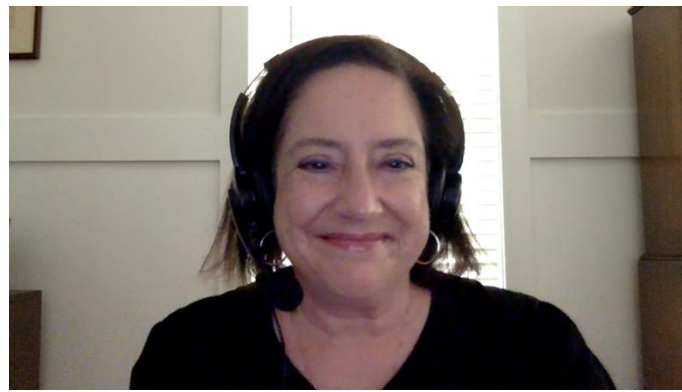
Module 8: Interacting with the oncology team

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Other questions



Thank you



Thank you for joining us!

Please take a moment to complete the survey currently up on Zoom. Your feedback is very important to us. The survey will remain open for 5 minutes after the meeting ends.