

EPICS

Conference Coverage: EHA 2024 – Focus on Multiple Myeloma
Saturday, June 15, 2024; Madrid, Spain
7.30 PM – 10.30 PM CEST; Total time: 3 hours

Chair: Rafael Fonseca, MD

Confirmed Faculty

- Shaji Kumar, MD
- Bruno Paiva, PharmD, PhD
- Thierry Facon, MD
- Mohamad Mohty, MD, PhD
- Hermann Einsele, MD, FRCP
- Evangelos Terpos, MD, PhD
- Sagar Lonial, MD, FACP

AGENDA

Time (CEST)	Topic	Presenter
7.30 PM – 7.35 PM (5 min)	Welcome and Introductions	Rafael Fonseca, MD
7.35 PM – 7.45 PM (10 min)	Newly Diagnosed Multiple Myeloma: Transplant Eligible	Evangelos Terpos, MD, PhD
	<ul style="list-style-type: none"> • <u>S201: Daratumumab + Bortezomib/Lenalidomide/Dexamethasone in Transplant-Eligible Patients With Newly Diagnosed Multiple Myeloma: Analysis of Minimal Residual Disease in the PERSEUS Trial. Sonneveld P, et al</u> • <u>S202: Isatuximab, Lenalidomide, Bortezomib and Dexamethasone for Newly-Diagnosed, Transplant-Eligible Multiple Myeloma: Post Transplantation Interim Analysis of the Randomized Phase III GMMG-HD7 Trial. Raab MS, et al</u> • <u>S204: Daratumumab (Dara) + Bortezomib/Thalidomide/Dexamethasone (D-VTd) Followed by Dara Maintenance in Transplant-Eligible (TE) Newly Diagnosed Multiple Myeloma (NDMM): >6-Year Update of CASSIOPEIA. Moreau P, et al</u> • <u>S205: Ciltacabtagene Autoleucl ± Lenalidomide Maintenance in Newly Diagnosed Multiple Myeloma With Suboptimal Response to Frontline Autologous Stem Cell Transplant: CARTITUDE-2 Cohort D. Roeloffzen W, et al</u> • <u>P974: Daratumumab (Dara)/Bortezomib/Lenalidomide/Dexamethasone (D-VRd) With D-R Maintenance (maint) in Transplant-Eligible (TE) Newly Diagnosed Myeloma (NDMM): Analysis of PERSEUS Based on Cytogenetic Risk. Dimopoulos M, et al</u> 	

Time (CEST)	Topic	Presenter
	<p><i>Backup:</i></p> <ul style="list-style-type: none"> • <u>S200: Phase I Open-Label Single-Arm Study of Dual Targeting BCMA and CD19 FasTCAR-T (GC012F) as First-Line Therapy for Transplant-Eligible Newly Diagnosed High-Risk Multiple Myeloma. Du J, et al</u> 	
7.45 PM – 8.05 PM (20 min)	Discussion	Moderator: Rafael Fonseca, MD
	<ul style="list-style-type: none"> • Should patients still be categorized by transplant eligibility? • How do you currently treat NDMM transplant-eligible patients, and has this changed? • What changes to maintenance therapy do you foresee happening in the coming years? • Are the anti-CD38 antibodies interchangeable? How do you decide which to use up front? • What role does MRD play in the clinic currently, especially now that the FDA recommends MRD as a possible primary endpoint in clinical trials? What barriers are there for community-based physicians to adopt such practices? 	
8.05 PM – 8.10 PM (5 min)	Key Takeaways	Evangelos Terpos, MD, PhD
8.10 PM – 8.20 PM (10 min)	Newly Diagnosed Multiple Myeloma: Transplant Ineligible	Shaji Kumar, MD
	<ul style="list-style-type: none"> • <u>S100: Phase 3 Study Results of Isatuximab, Bortezomib, Lenalidomide, and Dexamethasone (Isa-VRd) Versus VRd for Transplant-Ineligible Patients With Newly Diagnosed Multiple Myeloma (IMROZ). Facon T, et al</u> • <u>S203: Randomized Phase 3 Study of Isatuximab (Isa) Plus Lenalidomide and Dexamethasone (Rd) With Bortezomib (V) Versus IsaRd in Patients With Newly Diagnosed Transplant Ineligible Multiple Myeloma (NDMM TI). Leleu X, et al</u> • <u>P920: Safety Results From the Phase 3 MajesTEC-7 Study in Patients With Transplant Ineligible/Not Intended Newly Diagnosed Multiple Myeloma (NDMM). Van de Donk WCJ, et al</u> • <u>P907: KTd or KRd Induction Followed by K Maintenance or Observation in Transplant Ineligible (TIE) Patients With Newly Diagnosed Multiple Myeloma (MM). Final Analysis of the AGMT MM02 Trial. Ludwig H, et al</u> • <u>P968: Final Survival Analysis of Daratumumab Plus Lenalidomide and Dexamethasone Versus Lenalidomide and Dexamethasone in Transplant-Ineligible Patients With Newly Diagnosed Multiple Myeloma: MAIA Study. Facon T, et al</u> 	
8.20 PM – 8.40 PM (20 min)	Discussion	Moderator: Rafael Fonseca, MD
	<ul style="list-style-type: none"> • What is your current standard of care in transplant-ineligible patients? • What combination regimens would you like to see explored in this population? • What changes to the landscape do you foresee for transplant-ineligible 	

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	patients? <ul style="list-style-type: none"> Which patients do you treat with carfilzomib in front line? What are your thoughts on the carfilzomib maintenance data? When would you select Isa-Rd? For which patients, and where in the treatment sequence? 	
8.40 PM – 8.45 PM (5 min)	Key Takeaways	Shaji Kumar, MD
8.45 PM – 8.55 PM (10 min)	Break	
8.55 PM – 9.05 PM (10 min)	Relapsed/Refractory Multiple Myeloma: Small Molecules	Thierry Facon, MD
	<ul style="list-style-type: none"> <u>P999: Novel Selinexor Triplet and Quadruplet Regimens (SNd, SPed, SBd, SDPd): Results From the Phase 1b/2 STOMP Multiple Myeloma Trial. Madan S, et al</u> <u>S214: Results From DREAMM-7 a Randomized Phase 3 Study of Belantamab Mafodotin + Bortezomib, and Dexamethasone vs Daratumumab + Bortezomib, and Dexamethasone in Relapsed/Refractory Multiple Myeloma. Mateos M-V, et al</u> <u>P912: Efficacy of Venetoclax-Dexamethasone v Pomalidomide-Dexamethasone in Patients With T(11;14)-Positive Relapsed/Refractory Multiple Myeloma [T(11;14)+ RRMM]:Phase 3 CANOVA Biomarker Subgroup Analysis. Bahlis NJ, et al</u> <u>P1982: A Phase 2 Study of Isatuximab in Combination With Pomalidomide and Dexamethasone in RRMM Patients With 1 Prior Line of Therapy. Terpos E, et al</u> <u>P1993: Outcomes With Selinexor, Bortezomib and Dexamethasone (SVd) in Patients With Relapsed Refractory Multiple Myeloma (RRMM): Regional Subgroup Analysis of the Phase 3 BOSTON Trial. Spencer A, et al</u> 	
9.05 PM – 9.25 PM (20 min)	Discussion	Moderator: Rafael Fonseca, MD
	<ul style="list-style-type: none"> What are your thoughts on the different selinexor regimens? Do you think belantamab mafodotin will be used in Europe for MM? Describe your experiences with or opinions on sequencing CD38 antibodies Would use venetoclax for T(11;14)-positive patients? Are there certain patients for whom an isatuximab-based regimen is preferred over a daratumumab-based regimen, such as 1q21 gain or other high-risk groups? How do you currently treat elderly, frail patients? What combination regimens would you like to see explored in this population? How do you decide which patients require carfilzomib? High risk, aggressive disease, etc? Which novel combination regimens with small molecules would you like to see explored, and in which population? 	

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9.25 PM – 9.30 PM (5 min)	Key Takeaways	Thierry Facon, MD
9.30 PM – 9.40 PM (10 min)	Relapsed/Refractory Multiple Myeloma: Antibodies and Bispecifics I	Mohamad Mohty, MD, PhD
	<ul style="list-style-type: none"> • <u>S210: Cevostamab in Patients With RRMM Who Are Triple-Class Refractory and Have Received a Prior BCMA-Targeted ADC or CAR T-Cell: Initial Results From the Phase I/II CAMMA 2 Study. Kumar S, et al</u> • <u>P906: Efficacy and Safety of Elranatamab Monotherapy in the Real-World Setting in Relapsed-Refractory Multiple Myeloma (RRMM): Results of the French Compassionate Use Program on Behalf of the IFM. Mohty M, et al</u> • <u>P923: Efficacy and Safety of Teclistamab in Patients With Relapsed/Refractory Multiple Myeloma With High-Risk Features: A Subgroup Analysis From the Phase 1/2 MajesTEC-1 Study. Costa LJ, et al</u> • <u>P915: Long-Term Efficacy and Safety Results From the Phase 1/2 MonumenTAL-1 Study of Talquetamab, a GPRC5D×CD3 Bispecific Antibody, in Patients With Relapsed/Refractory Multiple Myeloma. Rasche L, et al</u> • <u>P938: DREAMM-7 Update: Subgroup Analyses From a Phase 3 Trial of Belantamab Mafodotin + Bortezomib and Dexamethasone vs Daratumumab, Bortezomib, and Dexamethasone in Relapsed/Refractory Multiple Myeloma. Mateos M-V, et al</u> 	
9.40 PM – 9.50 PM (10 min)	Relapsed/Refractory Multiple Myeloma: Antibodies and Bispecifics II	Hermann Einsele, MD, FRCP
	<ul style="list-style-type: none"> • <u>S211: Efficacy, Safety, and Determination of RP2D of ABBV-383, a BCMA Bispecific Antibody, in Patients With Relapsed/Refractory Multiple Myeloma (RRMM). Weisel K, et al</u> • <u>P902: Real-World Schedule De-Escalation of Teclistamab in Patients With Relapsed/Refractory Multiple Myeloma. Usmani SZ, et al</u> • <u>P932: Long-Term Survival After Elranatamab Monotherapy in Patients With Relapsed or Refractory Multiple Myeloma (RRMM): MagnetisMM-3. Mohty M, et al</u> • <u>S212: Linvoseltamab in Patients With Relapsed/Refractory Multiple Myeloma in the LINKER-MM1 Study: Depth and Durability of Response at 14-Month Median Follow-Up. Lentzsch S, et al</u> • <u>P911: Talquetamab, a GPRC5D×CD3 Bispecific Antibody, in Combination With Pomalidomide in Patients With Relapsed/Refractory Multiple Myeloma: Safety and Efficacy Results From the Phase 1b MonumenTAL-2 Study. Searle E, et al</u> 	

Time (CEST)	Topic	Presenter
9.50 PM – 10.15 PM (25 min)	Discussion	Moderator: Rafael Fonseca, MD
	<ul style="list-style-type: none"> • What bispecific combination regimens would you like to see investigated? • How will bispecific antibodies with different targets be sequenced? • Has the optimal dosing schedule been identified for any bispecific? • Now that longer follow-up data for bispecifics are available, what are your thoughts on the potential of bispecifics in curing MM? • What barriers remain for community adoption of bispecifics? • So far this year, which bispecific antibody trial data were the most impactful? • Will CAR T or bispecifics have the greatest impact on the MM treatment landscape, and why? 	
10.15 PM – 10.25 PM (10 min)	Key Takeaways	Mohamad Mohty, MD, PhD, and Hermann Einsele, MD, FRCP
10.25 PM – 10.30 PM (5 min)	Summary and Closing Remarks	Rafael Fonseca, MD