

# EPICS

**Conference Coverage: EHA 2023 – Focus on AML and MDS**  
**Saturday, June 10, 2023; 18.30 – 21.30 CEST**  
**Live Meeting**

**Chair:** Elias Jabbour, MD

**US Faculty**

- Naval Daver, MD (University of Texas MD Anderson Cancer Center)
- Guillermo Garcia-Manero, MD (University of Texas MD Anderson Cancer Center)
- Alexander Perl, MD (Abramson Cancer Center, University of Pennsylvania)
- Rami Komrokji, MD (Moffitt Cancer Center)
- Jessica K. Altman, MD (Robert H. Lurie Comprehensive Cancer Center of Northwestern University)

**AGENDA**

Time (CEST)	Topic	Presenter
18.30 – 18.35 (5 min)	<b>Welcome and Introductions</b>	Elias Jabbour, MD
18.35 – 18.45 (10 min)	<p><b>New Developments in First-Line Treatment of Myelodysplastic Syndromes (MDS)</b></p> <p><b><u>Low-risk MDS</u></b></p> <ul style="list-style-type: none"> <li>• KER-050 Treatment Improved Markers of Erythropoietic Activity and Hematopoiesis Over Six Months Which Resulted in Hematological Responses Across a Broad, Lower-Risk MDS Population. Aristoteles Giagounidis, S166</li> <li>• Luspatercept Versus Epoetin Alfa for Treatment of Anemia in ESA-Naive Lower-Risk Myelodysplastic Syndromes (LR-MDS) Patients (pts) Requiring RBC Transfusions: Data From the Phase 3 COMMANDS Study. Matteo Giovanni Della Porta, S102</li> <li>• Luspatercept Restores Effective Erythropoiesis and Provides Superior and Sustained Clinical Benefit vs Epoetin Alfa: Biomarker Analysis From the Phase 3 COMMANDS Study. Uwe Platzbecker, P693</li> </ul> <p><b><u>High-risk MDS</u></b></p> <ul style="list-style-type: none"> <li>• Phase 1/2 Study of Oral Decitabine/Cedazuridine in Combination With Venetoclax in Treatment-Naïve Higher-Risk Myelodysplastic Syndromes or Chronic Myelomonocytic Leukemia. Alex Bataller, S172</li> </ul>	Guillermo Garcia-Manero, MD

<p>18.45 – 19.05 (20 min)</p>	<p><b>Discussion</b></p> <ul style="list-style-type: none"> <li>In your opinion, what are the most impactful data in frontline MDS presented at EHA 2023? <ul style="list-style-type: none"> <li>How will you incorporate these new data into your current treatment approach for MDS?</li> </ul> </li> <li>Are there any investigational agents of particular interest, and why?</li> <li>What remains an unmet clinical need in first-line treatment of MDS?</li> </ul>	<p>All</p>
<p>19.05 – 19.15 (10 min)</p>	<p><b>New Developments in Treatment of Relapsed/Refractory (R/R) MDS</b></p> <p><b><u>MDS with mutations</u></b></p> <ul style="list-style-type: none"> <li>Higher <i>MDMX</i> Expression Was Associated With Hypomethylating Agent Resistance and Worse Survival in Myelodysplastic Syndrome Patients, Inferring It a Potential Therapeutic Target. Yu-Hung Wang, S171</li> </ul> <p><b><u>R/R MDS</u></b></p> <ul style="list-style-type: none"> <li>Continuous Transfusion Independence With Imetelstat in Heavily Transfused Non-del(5q) Lower-Risk Myelodysplastic Syndromes Relapsed/Refractory to Erythropoiesis Stimulating Agents in IMERGE Phase 3. Uwe Platzbecker, S165</li> <li>Disease Modifying Activity of Imetelstat in Patients With Heavily Transfused Non-del(5q) Lower-Risk Myelodysplastic Syndromes Relapsed/Refractory to Erythropoiesis Stimulating Agents in IMERGE Phase 3. Valeria Santini, S164</li> </ul> <p><b><u>Other</u></b></p> <ul style="list-style-type: none"> <li>Myelodysplastic Neoplasms (MDS) Classification From WHO 2017 to WHO 2022 and ICC 2022: An Expanded Analysis of 7017 Patients on Behalf of the International Consortium for MDS (ICMDS). Rami S. Komrokji, S170</li> </ul>	<p>Rami Komrokji, MD</p>
<p>19.15 – 19.35 (20 min)</p>	<p><b>Discussion</b></p> <ul style="list-style-type: none"> <li>In your opinion, what are the most impactful data in R/R MDS presented at EHA 2023? <ul style="list-style-type: none"> <li>How will you incorporate these new data into your current treatment approach for MDS?</li> </ul> </li> <li>What is your assessment of novel and emerging agents in the treatment of R/R MDS?</li> <li>What will be practice changing, and what will open new avenues of scientific investigation?</li> <li>What remains an unmet clinical need in R/R MDS?</li> </ul>	<p>All</p>
<p>19.35 – 19.40 (5 min)</p>	<p><b>Key Takeaways for MDS</b></p>	<p>Rami Komrokji, MD, and Guillermo Garcia-Manero, MD</p>

<p>19.40 – 19.50 (10 min)</p>	<p><b>Advances in Acute Myeloid Leukemia (AML): Newly Diagnosed</b></p> <p><b><u>AML with <i>FLT3</i>-wt</u></b></p> <ul style="list-style-type: none"> <li>• FLAG-IDA Combined With Gemtuzumab Ozogamicin (GO) Reduced MRD Levels and Improved Overall Survival in <i>NPM1</i>mut AML Independent of <i>FLT3</i> and MRD Status, Results From the AML19 Trial. Nigel Russell, S134</li> <li>• Preliminary Results of QUIWI: A Double Blinded, Randomized Clinical Trial Comparing Standard Chemotherapy Plus Quizartinib Versus Placebo in Adult Patients With Newly Diagnosed <i>FLT3</i>-ITD Wild-Type AML. Pau Montesinos, S130</li> </ul> <p><b><u>AML with <i>FLT3</i>-mut</u></b></p> <ul style="list-style-type: none"> <li>• Impact of Allogeneic Hematopoietic Cell Transplantation in First Complete Remission Plus <i>FLT3</i> Inhibition With Quizartinib in Acute Myeloid Leukemia With <i>FLT3</i>-ITD: Results From QuANTUM-First. Richard Schlenk, S137</li> <li>• Gemtuzumab-Based Induction Chemotherapy Combined With Midostaurin for <i>FLT3</i> Mutated AML. Updated Toxicity and Interim Survival Analysis From the NCR1 AML19V2 “Midotarg” Pilot Trial. Nigel Russell, P484</li> <li>• Next-Generation Sequencing-Based Measurable Residual Disease Monitoring in Acute Myeloid Leukemia With <i>FLT3</i> Internal Tandem Duplication Treated With Intensive Chemotherapy Plus Midostaurin. Frank Rucker, S135</li> </ul>	<p>Naval Daver, MD</p>
<p>19.50 – 20.15 (25 min)</p>	<p><b>Discussion</b></p> <ul style="list-style-type: none"> <li>• In your opinion, what are the most impactful data in newly diagnosed AML presented at EHA 2023?</li> <li>• How do you view the current and emerging data in newly diagnosed AML patients with <i>FLT3</i>-mutated or <i>NPM1</i>-mutated AML?</li> <li>• Will any of the presented data change your practice? How?</li> <li>• Can you comment on the sequencing of TKIs in <i>FLT3</i>-mutated AML patients, and which of these patients do you consider candidates for the different TKI regimens?</li> </ul>	<p>All</p>
<p>20.15 – 20.20 (5 min)</p>	<p><b>Break</b></p>	
<p>20.20 – 20.30 (10 min)</p>	<p><b>Advances in AML: Newly Diagnosed Elderly and/or Unfit</b></p> <ul style="list-style-type: none"> <li>• Phase II Study on Venetoclax Plus Decitabine for Elderly (<math>\geq 60 &lt; 75</math> years) Patients With Newly Diagnosed High-Intermediate Risk AML Eligible for Allo-SCT: Midterm Update of Ven-Dec GITMO Study. Domenico Russo, P502</li> </ul>	<p>Alexander Perl, MD</p>

	<ul style="list-style-type: none"> <li>Updated Results of VEN-A-QUI Study: A Phase 1-2 Trial to Assess the Safety and Efficacy of Triplets for Newly Diagnosed Unfit AML Patients: Azacitidine or Low-Dose Cytarabine With Venetoclax and Quizartinib. Juan Miguel Bergua Burgues, S132</li> <li>A Randomised Assessment of the Sequential Addition of the Kinase Inhibitor Quizartinib to Intensive Chemotherapy in Older Acute Myeloid Leukaemia (AML) Patients: Results From the NCRI AML18 Trial. Steven Knapper, S131</li> </ul>	
20.30 – 20.50 (20 min)	<p><b>Discussion</b></p> <ul style="list-style-type: none"> <li>What are your thoughts on the data presented regarding unfit and/or older adults with newly diagnosed AML?</li> <li>Will any of the presented data change your practice? How?</li> <li>What are unmet needs in AML treatment?</li> </ul>	All
20.50 – 20.55 (5 min)	<p><b>Advances in AML: R/R AML</b></p> <ul style="list-style-type: none"> <li>Olutasidenib in Post-venetoclax Patients With Mutant IDH1 AML. Jorge Cortes, P555</li> <li>Venetoclax (Ven) Combined With FLAG-IDA Is an Effective Regimen for Patients (pts) With Newly Diagnosed (ND) and Relapsed/Refractory (R/R) Acute Myeloid Leukemia (AML). Madelyn Burkart, P545</li> <li>Updated Data for Ziftomenib in Patients With NPM1-Mutated Relapsed or Refractory Acute Myeloid Leukemia. Amir Fathi, P504</li> </ul>	Jessica K. Altman, MD
20.55 – 21.20 (25 min)	<p><b>Discussion</b></p> <ul style="list-style-type: none"> <li>What is your preferred treatment approach, and how do you view the presented data in the real-life setting for R/R AML?</li> <li>What is your assessment of new and emerging targets in R/R AML?</li> <li>Will any of the presented data change your practice? How?</li> </ul>	All
21.20 – 21.25 (5 min)	<p><b>Key Takeaways</b></p>	Naval Daver, MD; Alexander Perl, MD; Jessica K. Altman, MD
21.25 – 21.30 (5 min)	<p><b>Summary and Closing Remarks</b></p>	Elias Jabbour, MD

**Total time: 3 hours**