1. **DISCUSSION:** Discuss the adequacy of the available data to support the use of minimal residual disease (MRD) as an accelerated approval endpoint in multiple myeloma (MM).

2. **DISCUSSION:** Discuss whether the available data supports the use of MRD as an endpoint in the different MM disease settings.
   - Newly diagnosed MM
   - Relapsed/Refractory MM

3. **DISCUSSION:** Discuss the acceptability of the timepoints for MRD assessment:
   - 9-months, 12-months, MRD negative complete response at any time
   - Requirement for assessment of durability

4. **VOTE:** Does the evidence support the use of MRD as an accelerated approval endpoint in MM clinical trials?