MRB AGENDA:

1. Present documented evidence/facts of failure or non-conformance.
   a. What, where, why, when, how?
2. Overstress analysis or additional testing required?
4. Impact to inventory, WIP, supplier PO, already built product.
5. Corrective action to prevent recurrence.
   a. Design, test, procedure options, trades & recommendation.
   b. Recommended Was/Is change description.
   c. Impacts to other subsystems.
   d. Affected documentation. (dwg, ICD, existing analysis, supplier EIDP).
   e. Verification plan to validate effective corrective action.
   f. Estimated costs & impact to schedule.
6. Effectiveness of corrective action
   a. Any modification to performance capability.
   b. Impact on FMEA, reliability, risk assessment.
7. Recommended final disposition(s):
   a. Rework
   b. Repair
   c. Return to Vendor
   d. Reclassify
   e. Scrap/Purge
   f. Use As Is
8. Action items: who-what-ECD.