March 23, 2018

Quarterly Report to NCIMS Executive Board on Technical Committee - AMI Subcommittee Activities

* The subcommittee members participated on 3 conference calls
* FDA was allowed to provide the draft document M-I-14-8 Supplement 2 to the subcommittee for review, comment and suggested revisions
* Members split out unresolved issues from the draft M-I for further discussion and research
* FDA accepted removal of issues by group and draft M-I-14-8 Supplement 2 will be issued by FDA
* Outstanding issues have been assigned to specific group members with instructions to work with subject matter experts. Group members will then bring information back to subcommittee for review and discussion.
* The subcommittee is also discussing using the Regional Equipment Review Committees as a venue to evaluate AMI equipment components that may not be covered by conventional equipment design standards.
* Next call is scheduled for April 16, 2018

Respectfully submitted by Helen Piotter – Vice Chair NCIMS Technical Review Committee