

INFORMATION PAPER

MRMC-MMP-I
3 October 2014

Subject: Medical Care Support Equipment (MEDCASE) / Super Capital Expense Equipment Program (SuperCEEP)

1. Purpose. To provide a MEDCASE / SuperCEEP Programs summary.

2. Facts:

a. The MEDCASE / SuperCEEP programs are regulated by the Department of the Army Supply Bulletin (SB) 8-75-MEDCASE. MEDCASE is defined as procurement for medical devices over \$250K purchase price based on OP funding requirements. SuperCEEP requirements are defined as medical device procurements that are \$100K to \$249.9K in purchase price based on O&M funding requirements.

b. Requirements can be generated by three different sources: TARA, Site (Medical Treatment Facility (MTF)), Special inserts (Congressional, BRAC, MILCON).

c. Prioritization is the process of validating, evaluating, ranking, and prioritizing the requirements. OTSG consultants and SMEs provide input and key facts for where requirements are ranked. During this prioritization requirements are validated and if needed deleted or deferred. After prioritization is complete the requirements are then vetted through the governing body, Strategic Technology Clinical Policies Council (STCPC). The STCPC body again validates, verifies, deletes or defers requirements as needed to support OTSG initiatives, mission requirements or special projects. Approved requirements lists (MEDCASE, SuperCEEP) are then forwarded to the Chief of Staff, MEDCOM whom upon signature will forward to TSG for signature. These lists then become part of Congressional appropriation funding requirements.

d. Procurement dollars are passed from TMA through MEDCOM through MRMC to USAMMA for medical device requirements. The approved requirements, MEDCASE/SuperCEEP, will be joined with the proper funding and forwarded to the correct contracting agency, DLA or regional contracting office.

3. The POC for this effort is Mr. Jimmy Bisenieks, MRMC-MMP-I, 301-619-7965, jimmy.l.bisenieks.civ@mail.mil

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