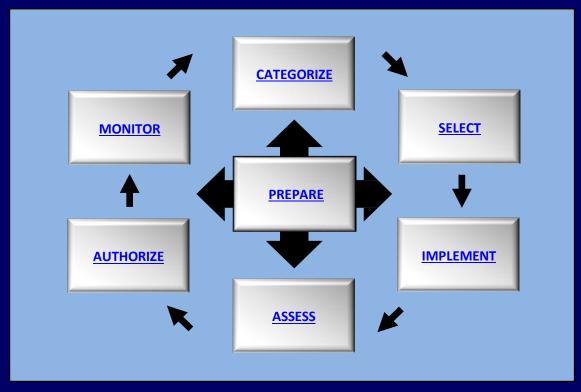
MEDICAL DEVICE ASSESSMENT AND AUTHORIZATION SERVICES





ASC will support your organization with applying the RMF Steps and activities including Assessment and Authorization and Continuous Monitoring activities. ASC will:

- Support your organization in identifying and assigning persons to RMF roles, establishing an organization-wide RM strategy, assessing organization-wide risk, developing organization-wide tailored security baselines, identifying Common Security Controls, and developing Organization-wide Information Security Continuous Monitoring (ISCM) strategy.
- Perform System Security Categorization activities (FIPS 199, NIST SPs 800-30, 800-39, 800-59, 800-60)
- Recommend Security Controls (FIPS Publications 199, 200; NIST SP 800-30, 800-53 Revision 4/5, Medical Device security control baseline)
- Perform Privacy Impact Assessment
- Perform E-Authentication Assessment
- Develop Medical Device and Equipment's System Contingency Plan
- Test Medical Device and Equipment's System Contingency Plan
- Develop Medical Device and Equipment's System Contingency Plan Test Results
- Develop Medical Device and Equipment's Incident Response Plan

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MEDICAL DEVICE ASSESSMENT AND AUTHORIZATION SERVICES

- Test Medical Device and Equipment's Incident Response Plan
- Develop Medical Device and Equipment's Incident Response Plan Test Results
- Develop Rules of Behavior
- Develop Organization/System/Device Security Policy
- Develop Medical Device and Equipment's Security Assessment Plan
- Perform Medical Device and Equipment's Risk Assessment (NIST SP 800-30, NIST SP 800-39)
- Develop Medical Device and Equipment's Risk Assessment Report
- Develop Medical Device and Equipment's System Security Plan (NIST SP 800-18)
- Conduct Medical Device and Equipment's Security Controls Assessment (NIST SP 800-53 R4/R5)
- Develop Medical Device and Equipment's Security Assessment Report
- Develop Medical Device and Equipment's Plan of Action and Milestones
- Monitor Security Controls

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