HealthManagement
Promoting Management and Leadership

Defibrillator Recalled Due to Defect

The product was recalled because the AED instructs the responder by voice prompts to press the shock button which is not visible because it is covered, thereby making the responder unable to provide shock therapy. The AED device should be removed from service, or the manufacturer-provided diagram should be consulted to remove and discard the shock button cover.

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