Food and Drug Administration, HHS

§ 810.1 Scope.

810.10 Cease distribution and notification order.

810.11 Regulatory hearing.

810.12 Written request for review of cease distribution and notification order.

810.13 Mandatory recall order.

810.14 Cease distribution and notification or mandatory recall strategy.

810.15 Communications concerning a cease distribution and notification or mandatory recall order.

810.16 Cease distribution and notification or mandatory recall order status reports.

810.17 Termination of a cease distribution and notification or mandatory recall order.

810.18 Public notice.


SOURCE: 61 FR 58018, Nov. 20, 1996, unless otherwise noted.

Subpart A—General Provisions

§ 810.1 Scope.

Part 810 describes the procedures that the Food and Drug Administration will follow in exercising its medical device recall authority under section 518(e) of the Federal Food, Drug, and Cosmetic Act.