

CPG Sec. 100.350 FDA Jurisdiction on Indian Reservations

BACKGROUND and POLICY:

The FDA considers Indian Reservations to be possessions of the United States within the meaning of section 201(a)(2) of the Federal Food, Drug, and Cosmetic Act. Consequently, FDA has complete jurisdiction over products within the purview of the Act that are manufactured on an Indian reservation. The products are in interstate commerce within the meaning of section 201(b) of the Act at all times.

Under the Food, Drug and Cosmetic Act, the FDA has the same authority on a reservation as it does anywhere else, to inspect, to take official samples, and to initiate regulatory actions.

Material between asterisks is new or revised

Issued: 3/15/77

Revised: 10/1/80, 9/1/87

More in Compliance Policy Guides

(/ICECI/ComplianceManuals/CompliancePolicyGuidanceManual/default.htm)

Foreword: Compliance Policy Guides (CPGs)

(/ICECI/ComplianceManuals/CompliancePolicyGuidanceManual/ucm116271.htm)

Chapter 1 - General

(/ICECI/ComplianceManuals/CompliancePolicyGuidanceManual/ucm116280.htm)

Chapter 2 - Biologics

(/ICECI/ComplianceManuals/CompliancePolicyGuidanceManual/ucm116336.htm)

Chapter 3 - Devices

(/ICECI/ComplianceManuals/CompliancePolicyGuidanceManual/ucm116801.htm)

Chapter 4 - Human Drugs

(/ICECI/ComplianceManuals/CompliancePolicyGuidanceManual/ucm119572.htm)

Chapter 5 - Food, Colors, and Cosmetics

(/ICECI/ComplianceManuals/CompliancePolicyGuidanceManual/ucm119194.htm)

Chapter 6 - Veterinary Medicine

(/ICECI/ComplianceManuals/CompliancePolicyGuidanceManual/ucm117042.htm)