FDA 510(k) Premarket Notification

K900308 SPERM SELECT SYSTEM

CLEARED

Apr 13, 1990
81 days to decision

K900308 · Product code: **HDR** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k900308/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cap, Cervical (HDR)
Date received	Jan 22, 1990
Decision date	Apr 13, 1990
Days to decision	81 days
Third-party review	No

APPLICANT

Company	**Pharmacia, Inc.**
Location	Mchenry, IL, US
Contact	ALBERT P MAYO
510(k) history	129 submissions · 126 cleared · 1976-1998