

K934585 PRO-FLO SYSTEM WITH INFRA-RED REMOTE CONTROLNov 3, 1993
41 days to decisionK934585 · Product code: **FCD** · Radiology
Source: <https://www.510kdatabase.net/k934585/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Barium Enema Kit (FCD)
Date received	Sep 23, 1993
Decision date	Nov 3, 1993
Days to decision	41 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Lafayette Pharmaceuticals, Inc.
Location	Lafayette, IN, US
Contact	ROBERT A SHARP
510(k) history	1 submissions · 1 cleared · 1993-1993

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k934585/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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