FDA 510(k) Premarket Notification

K903601 ESPI NON-INSULATED DOUBLE-ACTION GRASPER #8204

CLEARED

Sep 12, 1990
35 days to decision

K903601 · Product code: **HET** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k903601/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laparoscope, Gynecologic (and Accessories) (HET)
Date received	Aug 8, 1990
Decision date	Sep 12, 1990
Days to decision	35 days
Third-party review	No

APPLICANT

Company	**Dixon Medical, Inc.**
Location	Atlanta, GA, US
Contact	WILLIAM C DIXON
510(k) history	12 submissions · 12 cleared · 1990-1990