FDA 510(k) Premarket Notification

K935157 ARTICULATED, LAPAROSCOPIC, AND SIDE-VIEWING T-PROBES

CLEARED

Aug 9, 1994
285 days to decision

K935157 · Product code: **KPX** · Radiology
Source: https://www.510kdatabase.net/k935157/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Changer, Radiographic Film/cassette (KPX)
Date received	Oct 28, 1993
Decision date	Aug 9, 1994
Days to decision	285 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Regulatory Consultants TO the Medical Device Indus**
Location	Sandy, UT, US
Contact	WILLIAM E MCKAY
510(k) history	1 submissions · 1 cleared · 1994-1994