

**K935157 ARTICULATED, LAPAROSCOPIC, AND SIDE-VIEWING
T-PROBES**Aug 9, 1994
285 days to decisionK935157 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k935157/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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