FDA 510(k) Premarket Notification

CLEARED

K011155 INTERSTRAND 125 & INTERSTRAND 103, MODEL NUMBERS 1251S AND 1031S

Jul 12, 2001
87 days to decision

K011155 · Product code: **KXK** · Radiology
Source: https://www.510kdatabase.net/k011155/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Source, Brachytherapy, Radionuclide (KXK)
Date received	Apr 16, 2001
Decision date	Jul 12, 2001
Days to decision	87 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Ibt, Inc.**
Location	Norcross, GA, US
Contact	RUTH FEICHT
510(k) history	1 submissions · 1 cleared · 2001-2001