FDA 510(k) Premarket Notification

K761339 RADIAL TRAVEL SPEC. PROCEDURES TABLE

CLEARED

Jan 5, 1977
9 days to decision

K761339 · Product code: **IZZ** · Radiology
Source: https://www.510kdatabase.net/k761339/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Table, Radiographic, Non-tilting, Powered (IZZ)
Date received	Dec 27, 1976
Decision date	Jan 5, 1977
Days to decision	9 days
Third-party review	No

APPLICANT

Company	**Xre Corp.**
Location	Walker, MI, US
510(k) history	20 submissions · 20 cleared · 1977-1997