

K761339 RADIAL TRAVEL SPEC. PROCEDURES TABLEJan 5, 1977
9 days to decisionK761339 · 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

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Device record: <https://www.510kdatabase.net/k761339/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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