FDA 510(k) Premarket Notification

K820119 PODIOSCOPE BY ARROW

CLEARED

Feb 18, 1982
31 days to decision

K820119 · Product code: **KPW** · Radiology
Source: https://www.510kdatabase.net/k820119/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Beam Limiting, X-ray, Diagnostic (KPW)
Date received	Jan 18, 1982
Decision date	Feb 18, 1982
Days to decision	31 days
Third-party review	No

APPLICANT

Company	**Arrow X-Ray Corp.**
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1982-1982