

K820119 PODIOSCOPE BY ARROWFeb 18, 1982
31 days to decisionK820119 · 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

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Device record: <https://www.510kdatabase.net/k820119/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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