

K780319 ORBITOME-UGMar 14, 1978
18 days to decisionK780319 · Product code: **IZF** · Radiology
Source: <https://www.510kdatabase.net/k780319/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Tomographic (IZF)
Date received	Feb 24, 1978
Decision date	Mar 14, 1978
Days to decision	18 days
Third-party review	No

APPLICANT

Company	Philips Medical Systems (Cleveland), Inc.
Location	Mchenry, IL, US
510(k) history	190 submissions · 190 cleared · 1977-2017

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Device record: <https://www.510kdatabase.net/k780319/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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