

K812963 IBIS IINov 24, 1981
33 days to decisionK812963 · Product code: **JAA** · Radiology
Source: <https://www.510kdatabase.net/k812963/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Fluoroscopic, Image-intensified (JAA)
Date received	Oct 22, 1981
Decision date	Nov 24, 1981
Days to decision	33 days
Third-party review	No

APPLICANT

Company	Cgr Medical Corp.
Location	Mchenry, IL, US
510(k) history	20 submissions · 20 cleared · 1978-1984

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Device record: <https://www.510kdatabase.net/k812963/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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