

K833941 ORIONDec 29, 1983
44 days to decisionK833941 · Product code: **IYE** · Radiology
Source: <https://www.510kdatabase.net/k833941/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accelerator, Linear, Medical (IYE)
Date received	Nov 15, 1983
Decision date	Dec 29, 1983
Days to decision	44 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k833941/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 21, 2026