

K781872 CLINAC 2500Nov 27, 1978
24 days to decisionK781872 · Product code: **IYE** · RadiologySource: <https://www.510kdatabase.net/k781872/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accelerator, Linear, Medical (IYE)
Date received	Nov 3, 1978
Decision date	Nov 27, 1978
Days to decision	24 days
Third-party review	No

APPLICANT

Company	Varian Assoc., Inc.
Location	Mchenry, IL, US
510(k) history	86 submissions · 86 cleared · 1976-2000

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Device record: <https://www.510kdatabase.net/k781872/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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