FDA 510(k) Premarket Notification

K781872 CLINAC 2500

CLEARED

Nov 27, 1978
24 days to decision

K781872 · Product code: **IYE** · Radiology
Source: 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