FDA 510(k) Premarket Notification

K770202 CTM (CLINAC TREATMENT MONITOR)

CLEARED

Feb 7, 1977
7 days to decision

K770202 · Product code: **KPQ** · Radiology
Source: https://www.510kdatabase.net/k770202/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Simulation, Radiation Therapy (KPQ)
Date received	Jan 31, 1977
Decision date	Feb 7, 1977
Days to decision	7 days
Third-party review	No

APPLICANT

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