

K770202 CTM (CLINAC TREATMENT MONITOR)Feb 7, 1977
7 days to decisionK770202 · 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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k770202/> 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 19, 2026