FDA 510(k) Premarket Notification

K780501 ELECTRODE TESTER

CLEARED

May 19, 1978
52 days to decision

K780501 · Product code: **KRC** · Cardiovascular
Source: https://www.510kdatabase.net/k780501/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tester, Electrode, Surface, Electrocardiographic (KRC)
Date received	Mar 28, 1978
Decision date	May 19, 1978
Days to decision	52 days
Third-party review	No

APPLICANT

Company	**Mg Medical Electronics**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1978-1978