FDA 510(k) Premarket Notification

K830877 AP-CHECK

CLEARED

Jun 8, 1983
82 days to decision

K830877 · Product code: **GYA** · Neurology
Source: https://www.510kdatabase.net/k830877/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tester, Electrode/lead, Electroencephalograph (GYA)
Date received	Mar 18, 1983
Decision date	Jun 8, 1983
Days to decision	82 days
Third-party review	No

APPLICANT

Company	**General Devices**
Location	Bergenfield, NJ, US
510(k) history	13 submissions · 13 cleared · 1983-2002