FDA 510(k) Premarket Notification

K802637 TECA MODEL-10 ELECTROMYOGRAPH

CLEARED

Nov 26, 1980
35 days to decision

K802637 · Product code: **IKN** · Neurology
Source: https://www.510kdatabase.net/k802637/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electromyograph, Diagnostic (IKN)
Date received	Oct 22, 1980
Decision date	Nov 26, 1980
Days to decision	35 days
Third-party review	No

APPLICANT

Company	**Teca, Inc.**
Location	Mchenry, IL, US
510(k) history	29 submissions · 29 cleared · 1976-1996