FDA 510(k) Premarket Notification

K820414 UNIVERSAL TESTING & ANALYSIS SYSTEMS

CLEARED

Mar 4, 1982
20 days to decision

K820414 · Product code: **GWF** · Neurology
Source: https://www.510kdatabase.net/k820414/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Electrical, Evoked Response (GWF)
Date received	Feb 12, 1982
Decision date	Mar 4, 1982
Days to decision	20 days
Third-party review	No

APPLICANT

Company	**Lkc Technologies, Inc.**
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1976-2015