FDA 510(k) Premarket Notification

K812180 LIFE-TECH #5102 VISULAB

CLEARED

Aug 21, 1981
18 days to decision

K812180 · Product code: **GWN** · Neurology
Source: https://www.510kdatabase.net/k812180/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Nystagmograph (GWN)
Date received	Aug 3, 1981
Decision date	Aug 21, 1981
Days to decision	18 days
Third-party review	No

APPLICANT

Company	**Life-Tech Instruments, Inc.**
Location	Mchenry, IL, US
510(k) history	19 submissions · 19 cleared · 1977-1981

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k812180/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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