FDA 510(k) Premarket Notification

K900606 'CONDUCTOR' DISPOSABLE TENS ELECTRODE

CLEARED

Mar 13, 1990
33 days to decision

K900606 · Product code: **GXY** · Neurology
Source: https://www.510kdatabase.net/k900606/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	Feb 8, 1990
Decision date	Mar 13, 1990
Days to decision	33 days
Third-party review	No

APPLICANT

Company	**Medical Science Products, Inc.**
Location	Canal Fulton, OH, US
Contact	GARY SMITH
510(k) history	3 submissions · 3 cleared · 1990-1990

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k900606/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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