FDA 510(k) Premarket Notification

K944603 CONMED VERSA STEM NEUROMUSCULAR STIMULATION/TENS

CLEARED

Dec 15, 1994
87 days to decision

K944603 · Product code: **GXY** · Neurology
Source: https://www.510kdatabase.net/k944603/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	Sep 19, 1994
Decision date	Dec 15, 1994
Days to decision	87 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Conmedcorp**
Location	Dayton, OH, US
Contact	IRA D DUESLER JR.
510(k) history	92 submissions · 92 cleared · 1981-2010