

**K944603 CONMED VERSA STEM NEUROMUSCULAR STIMULATION/TENS**

Dec 15, 1994  
87 days to decision

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

**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k944603/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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