

**K820082 POST-OP ELECTRODE (TENS ELECTRODE)**Jan 28, 1982  
16 days to decisionK820082 · Product code: **GXY** · Neurology  
Source: <https://www.510kdatabase.net/k820082/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	Jan 12, 1982
Decision date	Jan 28, 1982
Days to decision	16 days
Third-party review	No

**APPLICANT**

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Company	<b>Consolidated Medical Equipment, Inc.</b>
Location	Mchenry, IL, US
510(k) history	15 submissions · 15 cleared · 1980-1986

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Device record: <https://www.510kdatabase.net/k820082/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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