

K832296 S2 POSTERIOR-TIBAug 31, 1983
49 days to decisionK832296 · Product code: **GXY** · Neurology
Source: <https://www.510kdatabase.net/k832296/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	Jul 13, 1983
Decision date	Aug 31, 1983
Days to decision	49 days
Third-party review	No

APPLICANT

Company	Anstek Corp.
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1983-1983

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Device record: <https://www.510kdatabase.net/k832296/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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