

K810206 DUAL TYPE ELECTRODEMar 20, 1981
53 days to decisionK810206 · Product code: **GXY** · Neurology
Source: <https://www.510kdatabase.net/k810206/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	Jan 26, 1981
Decision date	Mar 20, 1981
Days to decision	53 days
Third-party review	No

APPLICANT

Company	Professional Instruments Co.
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1981-1987

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Device record: <https://www.510kdatabase.net/k810206/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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