

K830600 REFERENCE & MONITORING ELECTRODEMar 24, 1983
28 days to decisionK830600 · Product code: **GXY** · Neurology
Source: <https://www.510kdatabase.net/k830600/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	Feb 24, 1983
Decision date	Mar 24, 1983
Days to decision	28 days
Third-party review	No

APPLICANT

Company	Precision Cap Systems
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1983-1983

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k830600/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 1, 2026