

K790269 CEPTOR HESIVEFeb 26, 1979
21 days to decisionK790269 · Product code: **GXY** · Neurology
Source: <https://www.510kdatabase.net/k790269/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	Feb 5, 1979
Decision date	Feb 26, 1979
Days to decision	21 days
Third-party review	No

APPLICANT

Company	Med General
Location	Mchenry, IL, US
510(k) history	12 submissions · 12 cleared · 1977-1979

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Device record: <https://www.510kdatabase.net/k790269/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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