

K802243 VAXIOM FAMILY OF RF LESION PROBES & ELECNov 12, 1980
57 days to decisionK802243 · Product code: **GXI** · Neurology
Source: <https://www.510kdatabase.net/k802243/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Probe, Radiofrequency Lesion (GXI)
Date received	Sep 16, 1980
Decision date	Nov 12, 1980
Days to decision	57 days
Third-party review	No

APPLICANT

Company	Progress Mankind Technology
Location	Hopkins, MN, US
Contact	Alfred A Iversen
510(k) history	17 submissions · 16 cleared · 1980-1988

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