FDA 510(k) Premarket Notification

K802243 VAXIOM FAMILY OF RF LESION PROBES & ELEC

CLEARED

Nov 12, 1980
57 days to decision

K802243 · 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