FDA 510(k) Premarket Notification

K030517 BALL RECORDING ELECTRODE/STIMULATION PROBE

CLEARED

Aug 1, 2003
163 days to decision

K030517 · Product code: **GZK** · Neurology
Source: https://www.510kdatabase.net/k030517/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Electrode, Nasopharyngeal (GZK)
Date received	Feb 19, 2003
Decision date	Aug 1, 2003
Days to decision	163 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Viasys Healthcare, Inc.**
Location	Stoughton, WI, US
Contact	Gary Syring
510(k) history	3 submissions · 3 cleared · 2003-2006