FDA 510(k) Premarket Notification

K023525 RETISCAN, RETIPORT

CLEARED

Mar 26, 2004
522 days to decision

K023525 · Product code: **GWE** · Neurology
Source: https://www.510kdatabase.net/k023525/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Stimulator, Photic, Evoked Response (GWE)
Date received	Oct 21, 2002
Decision date	Mar 26, 2004
Days to decision	522 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Roland Consult**
Location	Brandenburg, DE
Contact	MATTHIAS MAI
510(k) history	1 submissions · 1 cleared · 2004-2004