FDA 510(k) Premarket Notification

K030140 DUOMED, SERIES 500, (MODELS ID 500, FL 500)

CLEARED

Mar 18, 2003
63 days to decision

K030140 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k030140/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Jan 14, 2003
Decision date	Mar 18, 2003
Days to decision	63 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Duomed, Inc.**
Location	West Palm Beach, FL, US
Contact	CHARLES KOKINOS
510(k) history	1 submissions · 1 cleared · 2003-2003