FDA 510(k) Premarket Notification

CLEARED

K041340 GUIDANT MICROWAVE ABLATION SYSTEM

Jul 28, 2004
69 days to decision

K041340 · Product code: **NEY** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k041340/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Ablation, Microwave And Accessories (NEY)
Date received	May 20, 2004
Decision date	Jul 28, 2004
Days to decision	69 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	**Guidant Corporation, Cardiac Surgery**
Location	S,Mta Clara, CA, US
Contact	NANCY GALLO
510(k) history	8 submissions · 8 cleared · 2002-2008