FDA 510(k) Premarket Notification

CLEARED

K072478 BAYLIS PAIN MANAGEMENT GENERATOR-TD, MODELS: PMG-115-TD AND PMG-230-TD

Dec 19, 2007
106 days to decision

K072478 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k072478/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Sep 4, 2007
Decision date	Dec 19, 2007
Days to decision	106 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Baylis Medical Co., Inc.**
Location	Mississauga, CA
Contact	MEGHAL KHAKHAR
510(k) history	28 submissions · 28 cleared · 1998-2013