FDA 510(k) Premarket Notification

CLEARED

K020354 BAYLIS PAIN MANAGEMENT GENERATOR (PMG), MODELS PMG-115 (DOMESTIC), PMG-230 (INTERNATIONAL)

May 3, 2002
88 days to decision

K020354 · Product code: **GXD** · Neurology
Source: https://www.510kdatabase.net/k020354/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Generator, Lesion, Radiofrequency (GXD)
Date received	Feb 4, 2002
Decision date	May 3, 2002
Days to decision	88 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k020354/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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