

**K002565 BAYLIS PAIN MANAGEMENT GENERATOR MODEL:
PMG-115 (FOR DOMESTIC USE), PMG-230 (FOR
INTERNATIONAL USE)**

May 3, 2001
259 days to decision

K002565 · Product code: **GXD** · Neurology
Source: <https://www.510kdatabase.net/k002565/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Generator, Lesion, Radiofrequency (GXD)
Date received	Aug 17, 2000
Decision date	May 3, 2001
Days to decision	259 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/k002565/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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