

**K092337 PAIN MANAGEMENT OPTIMA, MODEL PMO21-100-05,
PMO20-100-10CS, PMO20-145-10CS**Oct 16, 2009
73 days to decisionK092337 · Product code: **GXI** · Neurology
Source: <https://www.510kdatabase.net/k092337/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Probe, Radiofrequency Lesion (GXI)
Date received	Aug 4, 2009
Decision date	Oct 16, 2009
Days to decision	73 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k092337/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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