FDA 510(k) Premarket Notification

K071745 BAYLIS PAIN MANAGEMENT SINGLE-USE PROBE

CLEARED

Jul 19, 2007
22 days to decision

K071745 · Product code: **GXI** · Neurology
Source: https://www.510kdatabase.net/k071745/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Probe, Radiofrequency Lesion (GXI)
Date received	Jun 27, 2007
Decision date	Jul 19, 2007
Days to decision	22 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