FDA 510(k) Premarket Notification

CLEARED

K981246 BEVERLY REFERENTIAL VALVE (NL850-9710) WITH INTEGRAL PERITONEAL CATHETER (NL850-9720) VALVE KIT (NL850-9730)

May 26, 1999
415 days to decision

K981246 · Product code: **JXG** · Neurology
Source: https://www.510kdatabase.net/k981246/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Shunt, Central Nervous System And Components (JXG)
Date received	Apr 6, 1998
Decision date	May 26, 1999
Days to decision	415 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Integra Neurocare, LLC**
Location	San Diego, CA, US
Contact	LORI HAYS
510(k) history	12 submissions · 12 cleared · 1996-2000

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k981246/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 30, 2026