FDA 510(k) Premarket Notification

K012889 BANICIDE LIQUID CHEMICAL STERILANT/HIGH-LEVEL DISINFECTANT

CLEARED

May 27, 2003
637 days to decision

K012889 · Product code: **MED** · General Hospital
Source: https://www.510kdatabase.net/k012889/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sterilant, Medical Devices (MED)
Date received	Aug 28, 2001
Decision date	May 27, 2003
Days to decision	637 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Pascal Co., Inc.**
Location	Bellevue, WA, US
Contact	DAVID R JOY
510(k) history	8 submissions · 8 cleared · 1991-2010