

K945627 PIKOS LP 01, PIKOS LP E01Mar 4, 1996
474 days to decisionK945627 · Product code: **DXY** · Cardiovascular
Source: <https://www.510kdatabase.net/k945627/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent - SP
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Nov 16, 1994
Decision date	Mar 4, 1996
Days to decision	474 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Biotronik, Inc.
Location	Lake Oswego, OR, US
Contact	LAUREN L FOOTE CHRISTENSEN
Website	https://www.biotronik.com
510(k) history	85 submissions · 67 cleared · 1994-2026

Biotronik, Inc. designs and manufactures advanced active implants for cardiac rhythm management, monitoring, and electrophysiology. The company operates with a manufacturing facility in Lake Oswego, Oregon, and serves patients globally through innovative cardiovascular solutions. Biotronik has received FDA 510(k) clearances from total submissions since its first clearance in 1994. The company specializes exclusively in cardiovascular devices, including pacing systems, implantable cardioverter defibrillators, cardiac resynchronization therapies, and electrophysiology cathe...
