

**K902889 A1-A,A1-B,A1-ABP,A1-MBP,A1-Z,A6-A,A6-B,PEH  
PACE.**Aug 23, 1990  
52 days to decisionK902889 · Product code: **DTD** · Cardiovascular  
Source: <https://www.510kdatabase.net/k902889/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pacemaker Lead Adaptor (DTD)
Date received	Jul 2, 1990
Decision date	Aug 23, 1990
Days to decision	52 days
Third-party review	No

**APPLICANT**

---

Company	<b>Biotronik, GmbH &amp; Co.</b>
Location	Lake Oswego, OR, US
Contact	STOUT, MD
510(k) history	18 submissions · 17 cleared · 1989-2000

---

**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k902889/> 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 July 4, 2026