

K792526 PULSE GENERATOR, MODEL NO. LIT 222Feb 5, 1980
60 days to decisionK792526 · Product code: **DXY** · Cardiovascular
Source: <https://www.510kdatabase.net/k792526/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Dec 7, 1979
Decision date	Feb 5, 1980
Days to decision	60 days
Third-party review	No

APPLICANT

Company	Bentley Laboratories, Inc.
Location	Mchenry, IL, US
Website	https://www.bentleyinstruments.com
510(k) history	55 submissions · 55 cleared · 1976-1993

Bentley Laboratories, Inc. is located in McHenry, US. The company has a historical record of FDA 510(k) device clearances spanning from 1976 to 1993. Bentley Laboratories received FDA 510(k) clearances from total submissions. The company specialized primarily in Cardiovascular devices, which represented approximately 80% of its regulatory submissions. Notable cleared devices included blood cardioplegia heat exchangers, venous reservoir bags, membrane oxygenators, and central venous catheters used in cardiac surgery and perfusion applications. The company is inactive and s...

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