

**K782161 PULSE GENERATOR, MODEL 171**Jul 24, 1979  
215 days to decisionK782161 · Product code: **DXY** · Cardiovascular  
Source: <https://www.510kdatabase.net/k782161/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Dec 21, 1978
Decision date	Jul 24, 1979
Days to decision	215 days
Third-party review	No

**APPLICANT**

---

Company	<b>Telectronics, Inc.</b>
Location	Mchenry, IL, US
510(k) history	107 submissions · 107 cleared · 1977-1990

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k782161/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 22, 2026