

**K802223 QUINTREX 0515 PROGRAMMABLE PULSE GEN.**

Sep 12, 1980

K802223 - Cardiovascular

Source: <https://www.510kdatabase.net/k802223/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent - SP
Submission type	Traditional
Date received	Sep 12, 1980
Decision date	Sep 12, 1980
Third-party review	No

**APPLICANT**

---

Company	<b>Medcor, Inc.</b>
Location	Mchenry, IL, US
510(k) history	8 submissions · 6 cleared · 1977-1980

---

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

Device record: <https://www.510kdatabase.net/k802223/> 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 July 9, 2026