FDA 510(k) Premarket Notification

K802223 QUINTREX 0515 PROGRAMMABLE PULSE GEN.

CLEARED

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	**Medcor, Inc.**
Location	Mchenry, IL, US
510(k) history	8 submissions · 6 cleared · 1977-1980