FDA 510(k) Premarket Notification

K001923 REGENCY POWER WHEELCHAIR, MODEL 7200, 7500 & 7800

CLEARED

Aug 25, 2000
63 days to decision

K001923 · Product code: **ITI** · Physical Medicine
Source: https://www.510kdatabase.net/k001923/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wheelchair, Powered (ITI)
Date received	Jun 23, 2000
Decision date	Aug 25, 2000
Days to decision	63 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Gendron, Inc.**
Location	Archbold, OH, US
Contact	FREDERIC W STROBEL
510(k) history	1 submissions · 1 cleared · 2000-2000