FDA 510(k) Premarket Notification

K940842 UNIPOWER RECHARGEABLE BATTERY PACK

CLEARED

Sep 7, 1994
196 days to decision

K940842 · Product code: **LDD** · Cardiovascular
Source: https://www.510kdatabase.net/k940842/

SUBMISSION DETAILS

Decision	Substantially Equivalent - ST
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Feb 23, 1994
Decision date	Sep 7, 1994
Days to decision	196 days
Third-party review	No

APPLICANT

Company	**Unipower Corp.**
Location	Minneapolis, MN, US
Contact	L.E PERTL
510(k) history	3 submissions · 2 cleared · 1994-1994