FDA 510(k) Premarket Notification

K890695 HI-RIDER WHEELCHAIR

CLEARED

Apr 10, 1989
59 days to decision

K890695 · Product code: **IPL** · Physical Medicine
Source: https://www.510kdatabase.net/k890695/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wheelchair, Standup (IPL)
Date received	Feb 10, 1989
Decision date	Apr 10, 1989
Days to decision	59 days
Third-party review	No

APPLICANT

Company	**Retec USA, Inc.**
Location	Orchard Park, NY, US
Contact	FRANK J KOPFER
510(k) history	1 submissions · 1 cleared · 1989-1989