FDA 510(k) Premarket Notification

K782148 HR UNIT

CLEARED

Jan 8, 1979
17 days to decision

K782148 · Product code: **IQP** · Physical Medicine
Source: https://www.510kdatabase.net/k782148/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Rotator, Transverse (IQP)
Date received	Dec 22, 1978
Decision date	Jan 8, 1979
Days to decision	17 days
Third-party review	No

APPLICANT

Company	**Parke-Davis Co.**
Location	Mchenry, IL, US
510(k) history	47 submissions · 47 cleared · 1976-1986