FDA 510(k) Premarket Notification

K810220 POWERED PATIENT SYSTEM

CLEARED

Feb 4, 1981
8 days to decision

K810220 · Product code: **IKZ** · Physical Medicine
Source: https://www.510kdatabase.net/k810220/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bed, Patient Rotation, Powered (IKZ)
Date received	Jan 27, 1981
Decision date	Feb 4, 1981
Days to decision	8 days
Third-party review	No

APPLICANT

Company	**Tri W-G, Inc.**
Location	Mchenry, IL, US
510(k) history	35 submissions · 35 cleared · 1979-1996