FDA 510(k) Premarket Notification

K843389 CLINITRON AIR FLUIDIZED SUPPORT J-101UD

CLEARED

Sep 5, 1984
7 days to decision

K843389 · Product code: **INX** · Physical Medicine
Source: https://www.510kdatabase.net/k843389/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bed, Air Fluidized (INX)
Date received	Aug 29, 1984
Decision date	Sep 5, 1984
Days to decision	7 days
Third-party review	No

APPLICANT

Company	**Uhi Corp.**
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1982-1984