FDA 510(k) Premarket Notification

K812683 NEONATE FLEXFLOAT

CLEARED

Oct 13, 1981
21 days to decision

K812683 · Product code: **FOH** · General Hospital
Source: https://www.510kdatabase.net/k812683/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mattress, Water, Temperature Regulated (FOH)
Date received	Sep 22, 1981
Decision date	Oct 13, 1981
Days to decision	21 days
Third-party review	No

APPLICANT

Company	**Pharmaquest Corp.**
Location	Walker, MI, US
510(k) history	14 submissions · 14 cleared · 1981-1995