

K812683 NEONATE FLEXFLOATOct 13, 1981
21 days to decisionK812683 · 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

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Device record: <https://www.510kdatabase.net/k812683/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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