FDA 510(k) Premarket Notification

K830450 DELMED IV CONTAINER

CLEARED

Mar 1, 1983
14 days to decision

K830450 · Product code: **KPE** · General Hospital
Source: https://www.510kdatabase.net/k830450/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Container, I.v. (KPE)
Date received	Feb 15, 1983
Decision date	Mar 1, 1983
Days to decision	14 days
Third-party review	No

APPLICANT

Company	**Delmed, Inc.**
Location	US
510(k) history	27 submissions · 26 cleared · 1979-1990

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k830450/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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