FDA 510(k) Premarket Notification

K840823 TREONIC C30

CLEARED

Mar 23, 1984
28 days to decision

K840823 · Product code: **LDR** · General Hospital
Source: https://www.510kdatabase.net/k840823/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Controller, Infusion, Intravascular, Electronic (LDR)
Date received	Feb 24, 1984
Decision date	Mar 23, 1984
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**Vickers America Medical Corp.**
Location	Mchenry, IL, US
510(k) history	22 submissions · 22 cleared · 1978-1984