

K840823 TREONIC C30Mar 23, 1984
28 days to decisionK840823 · 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

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Device record: <https://www.510kdatabase.net/k840823/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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