

K831761 SIMPLICITY PLUSAug 12, 1983
72 days to decisionK831761 · Product code: **FRN** · General Hospital
Source: <https://www.510kdatabase.net/k831761/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Infusion (FRN)
Date received	Jun 1, 1983
Decision date	Aug 12, 1983
Days to decision	72 days
Third-party review	No

APPLICANT

Company	Critikon Company, LLC
Location	Mchenry, IL, US
510(k) history	51 submissions · 51 cleared · 1979-2000

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Device record: <https://www.510kdatabase.net/k831761/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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