

**K041550 HOSPIRA PHOENIX INFUSION SYSTEM WITH
HOSPIRA MEDNET SOFTWARE**Jun 30, 2004
21 days to decisionK041550 · Product code: **FRN** · General Hospital
Source: <https://www.510kdatabase.net/k041550/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Infusion (FRN)
Date received	Jun 9, 2004
Decision date	Jun 30, 2004
Days to decision	21 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	Abbott Laboratories
Location	Abbott Park, IL, US
Contact	PATRICIA MELERSKI
Website	http://www.abbott.com
510(k) history	883 submissions · 868 cleared · 1976-2026

Abbott Laboratories is an American multinational medical devices and health care company headquartered in Abbott Park, Illinois. The company operates in over 160 countries and produces pharmaceuticals, diagnostics, nutritional products, and medical devices. Abbott maintains a substantial FDA 510(k) regulatory record with FDA 510(k) cleared devices from total submissions since 1976. The company's cleared devices span chemistry, microbiology, hematology, immunology, and toxicology categories. The latest clearance in 2025 reflects continued regulatory activity and product de...

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Device record: <https://www.510kdatabase.net/k041550/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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