

K896717 PARTNER INFUSION SYSTEM:Aug 2, 1990
246 days to decisionK896717 · Product code: **FRN** · General Hospital
Source: <https://www.510kdatabase.net/k896717/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Infusion (FRN)
Date received	Nov 29, 1989
Decision date	Aug 2, 1990
Days to decision	246 days
Third-party review	No

APPLICANT

Company	Q-Life Systems, Inc.
Location	Kingston, Ontario, CA
Contact	TIMOTHY SMITH
510(k) history	2 submissions · 2 cleared · 1990-1992

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k896717/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 7, 2026