FDA 510(k) Premarket Notification

K924110 LEOCOR CORFLO PUMP -- MODIFICATION

CLEARED

Sep 1, 1992
90 days to decision

K924110 · Product code: **DQI** · General Hospital
Source: https://www.510kdatabase.net/k924110/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Withdrawal/infusion Pump (DQI)
Date received	Jun 3, 1992
Decision date	Sep 1, 1992
Days to decision	90 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Leocor, Inc.**
Location	Webster, TX, US
Contact	ROGER W SNYDER
510(k) history	3 submissions · 3 cleared · 1990-1992