

**K924110 LEOCOR CORFLO PUMP -- MODIFICATION**Sep 1, 1992  
90 days to decisionK924110 · Product code: **DQI** · General Hospital  
Source: <https://www.510kdatabase.net/k924110/>**SUBMISSION DETAILS**

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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**

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Company	<b>Leacor, Inc.</b>
Location	Webster, TX, US
Contact	ROGER W SNYDER
510(k) history	3 submissions · 3 cleared · 1990-1992

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k924110/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated July 6, 2026