

**K920469 MYOCARDINAL TEMPERATURE SENSOR WITH TYPE
400 SERIE**Sep 11, 1992
228 days to decisionK920469 · Product code: FLL · General Hospital
Source: <https://www.510kdatabase.net/k920469/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Continuous Measurement Thermometer (FLL)
Date received	Jan 27, 1992
Decision date	Sep 11, 1992
Days to decision	228 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Respiratory Support Products, Inc.
Location	Mchenry, IL, US
Contact	KERRY TOMIC
510(k) history	27 submissions · 26 cleared · 1979-1999

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k920469/> 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 May 25, 2026