

**K883609 ACCES. PACKAGE FOR USE W/CARDIOGEN-82  
GENERATOR**Apr 14, 1989  
235 days to decisionK883609 · Product code: **FRN** · Radiology  
Source: <https://www.510kdatabase.net/k883609/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Infusion (FRN)
Date received	Aug 22, 1988
Decision date	Apr 14, 1989
Days to decision	235 days
Third-party review	No

**APPLICANT**

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Company	<b>Squibb Diagnostics</b>
Location	Mchenry, IL, US
Contact	LEONARD M BAUM
510(k) history	3 submissions · 3 cleared · 1983-1989

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k883609/>; 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 May 21, 2026