

**K883614 CYTOGUARD(R)**Sep 15, 1988  
22 days to decisionK883614 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k883614/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Aug 24, 1988
Decision date	Sep 15, 1988
Days to decision	22 days
Third-party review	No

**APPLICANT**

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Company	<b>Survival Technology, Inc.</b>
Location	Mchenry, IL, US
Contact	GARY W LEYLAND
510(k) history	10 submissions · 10 cleared · 1977-1993

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k883614/>; 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