FDA 510(k) Premarket Notification

K883614 CYTOGUARD(R)

CLEARED

Sep 15, 1988
22 days to decision

K883614 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k883614/

SUBMISSION DETAILS

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

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