FDA 510(k) Premarket Notification

K884233 QUADROJECT(TM)

CLEARED

Jan 30, 1989
115 days to decision

K884233 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k884233/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Oct 7, 1988
Decision date	Jan 30, 1989
Days to decision	115 days
Third-party review	No

APPLICANT

Company	**Ellis Pharmaceutical Consulting, Inc.**
Location	Walker, MI, US
Contact	LEVI ELLIS
510(k) history	5 submissions · 5 cleared · 1983-1991

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k884233/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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