FDA 510(k) Premarket Notification

K882975 RE-VENT ARTERIAL BLOOD SAMPLER AND/OR BIOSAMPLER

CLEARED

Jan 25, 1989
194 days to decision

K882975 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k882975/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Jul 15, 1988
Decision date	Jan 25, 1989
Days to decision	194 days
Third-party review	No

APPLICANT

Company	**Richmann Laboratories**
Location	Lake Park, FL, US
Contact	REED
510(k) history	1 submissions · 1 cleared · 1989-1989