FDA 510(k) Premarket Notification

K811432 MILLIPORE DUAL RATE INTRAVENOUS CASSETTE

CLEARED

Jul 13, 1981
52 days to decision

K811432 · Product code: **LDR** · General Hospital
Source: https://www.510kdatabase.net/k811432/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Controller, Infusion, Intravascular, Electronic (LDR)
Date received	May 22, 1981
Decision date	Jul 13, 1981
Days to decision	52 days
Third-party review	No

APPLICANT

Company	**Millipore Corp.**
Location	Mchenry, IL, US
510(k) history	30 submissions · 30 cleared · 1977-2003

Millipore Corp. was a global life science company founded in 1954, based in McHenry, US. Now part of Merck Group, the brand operates under the MilliporeSigma name following Merck's 2010 acquisition and subsequent 2015 merger with Sigma-Aldrich. The company holds FDA 510(k) clearances from total submissions, spanning 1977 to 2003. Millipore specialized in chemistry devices and filtration technologies, with cleared devices across anesthesiology, toxicology, and gastroenterology applications. This regulatory record reflects the company's historical focus on laboratory and cl...