FDA 510(k) Premarket Notification

K861687 BIOJECTOR SYRINGE SYSTEM

CLEARED

Apr 23, 1987
356 days to decision

K861687 · Product code: **KZE** · General Hospital
Source: https://www.510kdatabase.net/k861687/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Injector, Fluid, Non-electrically Powered (KZE)
Date received	May 2, 1986
Decision date	Apr 23, 1987
Days to decision	356 days
Third-party review	No

APPLICANT

Company	**Bioject, Inc.**
Location	Portland, OR, US
Contact	ALFRED ARONSON
510(k) history	9 submissions · 9 cleared · 1987-2009