FDA 510(k) Premarket Notification

K844244 DERATA MEDI-JECTOR PRO

CLEARED

Jul 5, 1985
246 days to decision

K844244 · Product code: **KZE** · General Hospital
Source: https://www.510kdatabase.net/k844244/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Injector, Fluid, Non-electrically Powered (KZE)
Date received	Nov 1, 1984
Decision date	Jul 5, 1985
Days to decision	246 days
Third-party review	No

APPLICANT

Company	**Derata Corp.**
Location	Mchenry, IL, US
Contact	KENNETH W DUNLAP
510(k) history	6 submissions · 6 cleared · 1977-1988