FDA 510(k) Premarket Notification

K930074 IN-VITRO DIAGNOSTIC REAGENT SET

CLEARED

May 11, 1993
124 days to decision

K930074 · Product code: **JGJ** · Chemistry
Source: https://www.510kdatabase.net/k930074/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Photometric Method, Magnesium (JGJ)
Date received	Jan 7, 1993
Decision date	May 11, 1993
Days to decision	124 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Horizon Diagnostics, Inc.**
Location	Jackson, MI, US
Contact	DAVID K GARCIA
510(k) history	7 submissions · 7 cleared · 1993-1997