FDA 510(k) Premarket Notification

K913809 HY-PREP SYSTEM

CLEARED

Jan 15, 1992
142 days to decision

K913809 · Product code: **JQW** · Chemistry
Source: https://www.510kdatabase.net/k913809/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Station, Pipetting And Diluting, For Clinical Use (JQW)
Date received	Aug 26, 1991
Decision date	Jan 15, 1992
Days to decision	142 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Hyperion, Inc.**
Location	Mchenry, IL, US
Contact	GREGORY STAMATIS
510(k) history	8 submissions · 8 cleared · 1984-2002