

**K913809 HY-PREP SYSTEM**

Jan 15, 1992  
 142 days to decision

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

**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k913809/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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