

K834300 HYBRITECH DATA STATIONFeb 21, 1984
74 days to decisionK834300 · Product code: **JQP** · Chemistry
Source: <https://www.510kdatabase.net/k834300/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Calculator/data Processing Module, For Clinical Use (JQP)
Date received	Dec 9, 1983
Decision date	Feb 21, 1984
Days to decision	74 days
Third-party review	No

APPLICANT

Company	Hybritech, Inc.
Location	Mchenry, IL, US
510(k) history	63 submissions · 63 cleared · 1981-1997

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k834300/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 30, 2026