

K810232 LEECO P.A.P. - QUANT DIAGNOSTIC KITFeb 10, 1981
14 days to decisionK810232 · Product code: **JFH** · Chemistry
Source: <https://www.510kdatabase.net/k810232/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Acid Phosphatase (prostatic), Tartrate Inhibited (JFH)
Date received	Jan 27, 1981
Decision date	Feb 10, 1981
Days to decision	14 days
Third-party review	No

APPLICANT

Company	Leeco Diagnostics, Inc.
Location	Mchenry, IL, US
510(k) history	49 submissions · 49 cleared · 1979-1989

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Device record: <https://www.510kdatabase.net/k810232/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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