

K821314 FEVER METERMay 28, 1982
24 days to decisionK821314 · Product code: **KDP** · General Hospital
Source: <https://www.510kdatabase.net/k821314/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Regulator, Vacuum (KDP)
Date received	May 4, 1982
Decision date	May 28, 1982
Days to decision	24 days
Third-party review	No

APPLICANT

Company	Steridyne Corp.
Location	Mchenry, IL, US
510(k) history	6 submissions · 6 cleared · 1980-1993

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Device record: <https://www.510kdatabase.net/k821314/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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