FDA 510(k) Premarket Notification

K821314 FEVER METER

CLEARED

May 28, 1982
24 days to decision

K821314 · 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