

K840118 RHINOTHERMApr 13, 1984
93 days to decisionK840118 · Product code: **KFZ** · AnesthesiologySource: <https://www.510kdatabase.net/k840118/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Humidifier, Non-direct Patient Interface (home-use) (KFZ)
Date received	Jan 11, 1984
Decision date	Apr 13, 1984
Days to decision	93 days
Third-party review	No

APPLICANT

Company	Ascot Pharmaceuticals, Inc.
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1984-1984

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Device record: <https://www.510kdatabase.net/k840118/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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