FDA 510(k) Premarket Notification

K840118 RHINOTHERM

CLEARED

Apr 13, 1984
93 days to decision

K840118 · Product code: **KFZ** · Anesthesiology
Source: 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