FDA 510(k) Premarket Notification

K790966 ULTRASONIC HUMIDIFIER

CLEARED

Jun 11, 1979
20 days to decision

K790966 · Product code: **KFZ** · Anesthesiology
Source: https://www.510kdatabase.net/k790966/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Humidifier, Non-direct Patient Interface (home-use) (KFZ)
Date received	May 22, 1979
Decision date	Jun 11, 1979
Days to decision	20 days
Third-party review	No

APPLICANT

Company	**Hankscraft Div. of Gerber Products Co.**
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1979-1985