

K790966 ULTRASONIC HUMIDIFIERJun 11, 1979
20 days to decisionK790966 · Product code: **KFZ** · AnesthesiologySource: <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

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Device record: <https://www.510kdatabase.net/k790966/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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