FDA 510(k) Premarket Notification

K884079 MODEL CBX EPILATOR

CLEARED

Oct 4, 1988
7 days to decision

K884079 · Product code: **KCW** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k884079/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Epilator, High Frequency, Needle-type (KCW)
Date received	Sep 27, 1988
Decision date	Oct 4, 1988
Days to decision	7 days
Third-party review	No

APPLICANT

Company	**R. A. Fischer Co. Corp.**
Location	Mchenry, IL, US
Contact	VAN ORDEN
510(k) history	13 submissions · 12 cleared · 1976-1997

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k884079/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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