

K940670 EARCRAFT HEARING AIDAug 4, 1994
169 days to decisionK940670 · Product code: **ESD** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k940670/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Feb 16, 1994
Decision date	Aug 4, 1994
Days to decision	169 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Earcraft, Inc.
Location	Hallowell, ME, US
Contact	ERIC WEEKS
510(k) history	1 submissions · 1 cleared · 1994-1994

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k940670/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 5, 2026