FDA 510(k) Premarket Notification

K936271 FRESH CELLS MULTI-WELL PLATE CULTURES AND SHELL VIAL CULTURE

CLEARED

Aug 26, 1994
239 days to decision

K936271 · Product code: **KIR** · Pathology
Source: https://www.510kdatabase.net/k936271/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cells, Animal And Human, Cultured (KIR)
Date received	Dec 30, 1993
Decision date	Aug 26, 1994
Days to decision	239 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Diagnostic Hybrids, Inc.**
Location	Athens, OH, US
Contact	J.L. BROWN
510(k) history	37 submissions · 36 cleared · 1988-2014

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k936271/ 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 May 14, 2026