

K003305 PEDIATRIC INGESTASCANDec 22, 2000
63 days to decisionK003305 · Product code: **HPM** · Ophthalmic
Source: <https://www.510kdatabase.net/k003305/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Locator, Metal, Electronic (HPM)
Date received	Oct 20, 2000
Decision date	Dec 22, 2000
Days to decision	63 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Adams Electronics, Inc.
Location	Enid, OK, US
Contact	ROBERT E ADAMS
510(k) history	1 submissions · 1 cleared · 2000-2000

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k003305/> 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 June 29, 2026