

K770888 POMERANTZEFF EQUATOR PLUS CAMERAJun 3, 1977
18 days to decisionK770888 · Product code: **HKI** · Ophthalmic
Source: <https://www.510kdatabase.net/k770888/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Camera, Ophthalmic, Ac-powered (HKI)
Date received	May 16, 1977
Decision date	Jun 3, 1977
Days to decision	18 days
Third-party review	No

APPLICANT

Company	Medical Instrument Research Assoc., Inc.
Location	Walker, MI, US
510(k) history	4 submissions · 4 cleared · 1976-1979

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k770888/> 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 1, 2026