FDA 510(k) Premarket Notification

K770888 POMERANTZEFF EQUATOR PLUS CAMERA

CLEARED

Jun 3, 1977
18 days to decision

K770888 · 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