FDA 510(k) Premarket Notification

K812544 RELIANCE OPHTHALMIC INSTRUMENT CONTROL

CLEARED

Sep 21, 1981
17 days to decision

K812544 · Product code: **HMF** · Ophthalmic
Source: https://www.510kdatabase.net/k812544/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stand, Instrument, Ac-powered, Ophthalmic (HMF)
Date received	Sep 4, 1981
Decision date	Sep 21, 1981
Days to decision	17 days
Third-party review	No

APPLICANT

Company	**F. & F. Koenigkramer**
Location	Mchenry, IL, US
510(k) history	9 submissions · 9 cleared · 1977-1986