

K812544 RELIANCE OPHTHALMIC INSTRUMENT CONTROLSep 21, 1981
17 days to decisionK812544 · 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

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Device record: <https://www.510kdatabase.net/k812544/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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