

**K954780 KOWA PROFESSIONAL FUNDUS CAMERA MODELS  
FX-500, FX-500S, FX-500C**Dec 1, 1995  
45 days to decisionK954780 · Product code: **HKI** · Ophthalmic  
Source: <https://www.510kdatabase.net/k954780/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Camera, Ophthalmic, Ac-powered (HKI)
Date received	Oct 17, 1995
Decision date	Dec 1, 1995
Days to decision	45 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Kowa Optimed, Inc.</b>
Location	Torrance, CA, US
Contact	SAMIA N RODRIGUEZ
510(k) history	10 submissions · 10 cleared · 1988-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k954780/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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