

K840104 PETRUS IOL GUIDEApr 24, 1984
105 days to decisionK840104 · Product code: **KYB** · Ophthalmic
Source: <https://www.510kdatabase.net/k840104/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Guide, Intraocular (KYB)
Date received	Jan 10, 1984
Decision date	Apr 24, 1984
Days to decision	105 days
Third-party review	No

APPLICANT

Company	Precision-Cosmet Co., Inc.
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1984-1986

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k840104/>; 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 June 30, 2026