FDA 510(k) Premarket Notification

K840104 PETRUS IOL GUIDE

CLEARED

Apr 24, 1984
105 days to decision

K840104 · 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