FDA 510(k) Premarket Notification

K874252 KREDA DISC

CLEARED

Jan 27, 1988
100 days to decision

K874252 · Product code: **HOQ** · Ophthalmic
Source: https://www.510kdatabase.net/k874252/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Grid, Amsler (HOQ)
Date received	Oct 19, 1987
Decision date	Jan 27, 1988
Days to decision	100 days
Third-party review	No

APPLICANT

Company	**Richmond Products, Inc.**
Location	Boca Raton, FL, US
Contact	LLOYD POWELL
510(k) history	15 submissions · 15 cleared · 1987-1988