FDA 510(k) Premarket Notification

K863429 MODIFIED DEVICE FOR SELF EXAMINATION OF EYES

CLEARED

Jan 20, 1987
138 days to decision

K863429 · Product code: **HOQ** · Ophthalmic
Source: https://www.510kdatabase.net/k863429/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Grid, Amsler (HOQ)
Date received	Sep 4, 1986
Decision date	Jan 20, 1987
Days to decision	138 days
Third-party review	No

APPLICANT

Company	**Wheel Checkers**
Location	Denver, CO, US
Contact	GIESKIENG
510(k) history	2 submissions · 2 cleared · 1985-1987