FDA 510(k) Premarket Notification

K943921 RYDER CONTACT LENS CASE

CLEARED

Oct 11, 1994
60 days to decision

K943921 · Product code: **LRX** · Ophthalmic
Source: https://www.510kdatabase.net/k943921/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Case, Contact Lens (LRX)
Date received	Aug 12, 1994
Decision date	Oct 11, 1994
Days to decision	60 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Clinical Research Assoc.**
Location	Brunswick, NJ, US
Contact	PATRICIA W BRADSTREET
510(k) history	1 submissions · 1 cleared · 1994-1994