FDA 510(k) Premarket Notification

K020930 MENICON PROGENT REMOVER FOR RIGID GAS PERMEABLE CONTACT LENSES

CLEARED

May 2, 2002
41 days to decision

K020930 · Product code: **MRC** · Ophthalmic
Source: https://www.510kdatabase.net/k020930/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Products, Contact Lens Care, Rigid Gas Permeable (MRC)
Date received	Mar 22, 2002
Decision date	May 2, 2002
Days to decision	41 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Menicon Co, Ltd.**
Location	Washington, DC, US
Contact	BEVERLEY D VENUTI
510(k) history	17 submissions · 17 cleared · 1994-2020