

K002140 MENICON PROGENT REMOVEROct 10, 2000
85 days to decisionK002140 · Product code: **MRC** · Ophthalmic
Source: <https://www.510kdatabase.net/k002140/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Products, Contact Lens Care, Rigid Gas Permeable (MRC)
Date received	Jul 17, 2000
Decision date	Oct 10, 2000
Days to decision	85 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k002140/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 18, 2026