

**K082855 MENICON PROGENT PROTEIN REMOVER FOR RIGID
GAS PERMEABLE CONTACT LENSES**Mar 9, 2010
526 days to decisionK082855 · Product code: **MRC** · Ophthalmic
Source: <https://www.510kdatabase.net/k082855/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Products, Contact Lens Care, Rigid Gas Permeable (MRC)
Date received	Sep 29, 2008
Decision date	Mar 9, 2010
Days to decision	526 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/k082855/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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