FDA 510(k) Premarket Notification

K931839 ROTA SYSTEM XL

CLEARED

Nov 22, 1993
223 days to decision

K931839 · Product code: **HMF** · Ophthalmic
Source: https://www.510kdatabase.net/k931839/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stand, Instrument, Ac-powered, Ophthalmic (HMF)
Date received	Apr 13, 1993
Decision date	Nov 22, 1993
Days to decision	223 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Rota Systems Medfurniture, Inc.**
Location	Derby, KS, US
Contact	SAM R MYERS
510(k) history	2 submissions · 2 cleared · 1993-1994