FDA 510(k) Premarket Notification

K872747 LASER CENTERING DEVICE

CLEARED

Sep 4, 1987
53 days to decision

K872747 · Product code: **HMR** · Ophthalmic
Source: https://www.510kdatabase.net/k872747/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Marker, Ocular (HMR)
Date received	Jul 13, 1987
Decision date	Sep 4, 1987
Days to decision	53 days
Third-party review	No

APPLICANT

Company	**Innomed Corp.**
Location	Valrico, FL, US
Contact	PETER K BROWN
510(k) history	3 submissions · 3 cleared · 1982-1987

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k872747/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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