FDA 510(k) Premarket Notification

K081756 OCULAR RESPONSE ANALYZER

CLEARED

Aug 7, 2008
48 days to decision

K081756 · Product code: **HKX** · Ophthalmic
Source: https://www.510kdatabase.net/k081756/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tonometer, Ac-powered (HKX)
Date received	Jun 20, 2008
Decision date	Aug 7, 2008
Days to decision	48 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Reichert, Inc.**
Location	Depew, NY, US
Contact	SANDRA BROWN
510(k) history	9 submissions · 9 cleared · 2004-2024