

K081756 OCULAR RESPONSE ANALYZERAug 7, 2008
48 days to decisionK081756 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k081756/> 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 May 26, 2026