

K950520 BURR, CORNEAL, BATTERY-POWEREDMay 2, 1995
85 days to decisionK950520 · Product code: **HOG** · Ophthalmic
Source: <https://www.510kdatabase.net/k950520/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Burr, Corneal, Battery-powered (HOG)
Date received	Feb 6, 1995
Decision date	May 2, 1995
Days to decision	85 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Aaron Medical Industries
Location	St. Petersburg, FL, US
Contact	J. R SARON
510(k) history	46 submissions · 46 cleared · 1990-2006

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k950520/> 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 June 29, 2026