FDA 510(k) Premarket Notification

K950520 BURR, CORNEAL, BATTERY-POWERED

CLEARED

May 2, 1995
85 days to decision

K950520 · 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