

K202760 Iridex 810 LaserOct 21, 2020
30 days to decisionK202760 · Product code: **HQF** · Ophthalmic
Source: <https://www.510kdatabase.net/k202760/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Laser, Ophthalmic (HQF)
Date received	Sep 21, 2020
Decision date	Oct 21, 2020
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Iridex Corporation
Location	Mountain View, CA, US
Contact	Bill Hyatt
Website	http://www.iridex.com/
510(k) history	9 submissions · 9 cleared · 2004-2023

Iridex Corporation is a worldwide leader in developing, manufacturing, and marketing innovative laser-based medical systems for the ophthalmology market. Founded in 1989, the company specializes in ophthalmic laser devices, delivery systems, and surgical instrumentation. Iridex maintains a manufacturing facility in Mountain View, California and serves customers globally through direct sales and approximately 60 independent distributors across over 100 countries. The company has received FDA 510(k) clearances from total submissions between 2004 and 2023. Iridex's regulator...
