

K062314 HORUS LASER KERATOMEDec 22, 2006
136 days to decisionK062314 · Product code: **GEX** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k062314/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Aug 8, 2006
Decision date	Dec 22, 2006
Days to decision	136 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Carl Zeiss Meditec, AG
Location	Dublin, CA, US
Contact	KENT W JONES
Website	http://www.zeiss.com/meditec-ag/en_de/home.html
510(k) history	45 submissions · 44 cleared · 2004-2025

Carl Zeiss Meditec, AG is a global medical device manufacturer specializing in innovative solutions for ophthalmology and microsurgery. The company operates with a manufacturing facility in Dublin, US, and delivers diagnostic and surgical instruments to healthcare professionals worldwide. The company has received FDA 510(k) clearances from total submissions since 2004. Ophthalmic devices represent the dominant category, accounting for 71% of submissions. The latest clearance in 2025 reflects continued regulatory activity and product innovation in this specialized field. C...
