

**K241600 Dermablade**Jul 3, 2024  
29 days to decisionK241600 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k241600/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Jun 4, 2024
Decision date	Jul 3, 2024
Days to decision	29 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Asclepion Laser Technologies GmbH</b>
Location	Chelmsford, MA, US
Contact	Tom Gruender
510(k) history	29 submissions · 29 cleared · 2004-2026

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k241600/> 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 14, 2026