

**K150650 neon3**Sep 24, 2015  
196 days to decisionK150650 · Product code: **NKG** · Orthopedic  
Source: <https://www.510kdatabase.net/k150650/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Posterior Cervical Screw System (NKG)
Date received	Mar 12, 2015
Decision date	Sep 24, 2015
Days to decision	196 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Ulrich GmbH &amp; Co. KG</b>
Location	Ulm, DE
Contact	Christoph Ulrich
510(k) history	24 submissions · 23 cleared · 2005-2025

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k150650/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 16, 2026