

**K010070 ULTRAPAK NEHA**Mar 15, 2001  
66 days to decisionK010070 · Product code: **MVL** · Dental  
Source: <https://www.510kdatabase.net/k010070/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cord, Retraction (MVL)
Date received	Jan 8, 2001
Decision date	Mar 15, 2001
Days to decision	66 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Ultradent Products, Inc.</b>
Location	Salt Lake City, UT, US
Contact	MARKUS R GEE
Website	<a href="https://www.ultradent.com">https://www.ultradent.com</a>
510(k) history	103 submissions · 103 cleared · 1992-2026

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k010070/> 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 June 28, 2026