

**K090713 ELEVATE PROLAPSE REPAIR SYSTEM WITH PC  
COATED INTEPRO LITE- APICAL AND POSTERIOR PROLAPSE  
REPAIR SYSTEM, ANTERIOR AND AP**Apr 2, 2009  
15 days to decisionK090713 · Product code: **OTP** · Obstetrics & Gynecology  
Source: <https://www.510kdatabase.net/k090713/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Mesh, Surgical, Synthetic, Urogynecologic, For Pelvic Organ Prolapse, Transvaginally Placed (OTP)
Date received	Mar 18, 2009
Decision date	Apr 2, 2009
Days to decision	15 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>American Medical Systems, Inc.</b>
Location	Mchenry, IL, US
Contact	DENISE THOMPSON
510(k) history	72 submissions · 72 cleared · 1979-2013

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k090713/> 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 14, 2026