

**K062222 UNITIP HIGH RESOLUTION CATHETER**Mar 15, 2007  
225 days to decisionK062222 · Product code: **FFX** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k062222/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Gastrointestinal Motility (electrical) (FFX)
Date received	Aug 2, 2006
Decision date	Mar 15, 2007
Days to decision	225 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Unisensor AG</b>
Location	Golden, CO, US
Contact	ROBERT N CLARK
510(k) history	2 submissions · 2 cleared · 2001-2007

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k062222/> 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