

**K233020 TauTona Pneumoperitoneum Assist Device (TPAD)**Jan 12, 2024  
112 days to decisionK233020 · Product code: **HIF** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k233020/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Insufflator, Laparoscopic (HIF)
Date received	Sep 22, 2023
Decision date	Jan 12, 2024
Days to decision	112 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Tautona Group Research &amp; Development, Co., LLC</b>
Location	Redwood City, CA, US
Contact	Michael Blanchette
510(k) history	1 submissions · 1 cleared · 2024-2024

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k233020/> 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