

**K022214 MSI I2010 DUAL PLACE HYPERBARIC CHAMBER**Feb 26, 2003  
233 days to decisionK022214 · Product code: **CBF** · Anesthesiology  
Source: <https://www.510kdatabase.net/k022214/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Chamber, Hyperbaric (CBF)
Date received	Jul 8, 2002
Decision date	Feb 26, 2003
Days to decision	233 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Mechidyne Systems, Inc.</b>
Location	Houston, TX, US
Contact	AUDREY L AARON
510(k) history	1 submissions · 1 cleared · 2003-2003

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k022214/> 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 25, 2026