FDA 510(k) Premarket Notification

K022214 MSI I2010 DUAL PLACE HYPERBARIC CHAMBER

CLEARED

Feb 26, 2003
233 days to decision

K022214 · 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	**Mechidyne Systems, Inc.**
Location	Houston, TX, US
Contact	AUDREY L AARON
510(k) history	1 submissions · 1 cleared · 2003-2003