

K123228 INBODY 770, INBODY 570, INBODY S10, INBODY H20/INBODY H20(B)Mar 8, 2013
144 days to decisionK123228 · Product code: **MNW** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k123228/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Body Composition (MNW)
Date received	Oct 15, 2012
Decision date	Mar 8, 2013
Days to decision	144 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Biospace Corporation Limited
Location	Deer Field, IL, US
Contact	DANIEL KAMM
510(k) history	9 submissions · 9 cleared · 2005-2014

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k123228/> Data sourced from FDA 510(k) public records (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. - Generated June 6, 2026