FDA 510(k) Premarket Notification

K130777 INBODY

CLEARED

Nov 25, 2013
249 days to decision

K130777 · Product code: **MNW** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k130777/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Body Composition (MNW)
Date received	Mar 21, 2013
Decision date	Nov 25, 2013
Days to decision	249 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k130777/ 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