FDA 510(k) Premarket Notification

K123228 INBODY 770, INBODY 570, INBODY S10, INBODY H20/INBODY H20(B)

CLEARED

Mar 8, 2013
144 days to decision

K123228 · 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