

K130856 BIA / VITALITY ANALYZERNov 25, 2013
242 days to decisionK130856 · Product code: **MNW** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k130856/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Body Composition (MNW)
Date received	Mar 28, 2013
Decision date	Nov 25, 2013
Days to decision	242 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Ipgdx, LLC
Location	Littleton, CO, US
Contact	Kevin Walls
510(k) history	1 submissions · 1 cleared · 2013-2013

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k130856/> 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 8, 2026