

**K121716 TBS INSIGHT**Oct 5, 2012  
116 days to decisionK121716 · Product code: **KGI** · Radiology  
Source: <https://www.510kdatabase.net/k121716/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Densitometer, Bone (KGI)
Date received	Jun 11, 2012
Decision date	Oct 5, 2012
Days to decision	116 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medimaps Group SA</b>
Location	Plan Les Ouates, CH
Contact	CHRISTOPHE LELONG
510(k) history	3 submissions · 3 cleared · 2012-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k121716/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 27, 2026