

**K123894 GE ECHOPAC**Jan 29, 2013  
42 days to decisionK123894 · Product code: **LLZ** · Radiology  
Source: <https://www.510kdatabase.net/k123894/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Dec 18, 2012
Decision date	Jan 29, 2013
Days to decision	42 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Ge Vingmed Ultrasound AS</b>
Location	Horten, NO
Contact	BRYAN BEHN
510(k) history	10 submissions · 10 cleared · 2009-2015

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k123894/> 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 16, 2026