

**K060923 NAVIAID BGE DEVICE, MODEL 21-001**Aug 15, 2006  
133 days to decisionK060923 · Product code: **FDA** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k060923/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Enteroscope And Accessories (FDA)
Date received	Apr 4, 2006
Decision date	Aug 15, 2006
Days to decision	133 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Smart Medical Systems , Ltd.</b>
Location	Kfar Saba, IL
Contact	AHAVA STEIN
510(k) history	9 submissions · 9 cleared · 2006-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k060923/> 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