

K960203 G.E. STERILE AMNIOCENTESIS TRAYMar 14, 1996
58 days to decisionK960203 · Product code: **HIO** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k960203/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sampler, Amniotic Fluid (amniocentesis Tray) (HIO)
Date received	Jan 16, 1996
Decision date	Mar 14, 1996
Days to decision	58 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	GE Medical Systems
Location	Milwaukee, WI, US
Contact	PATRICK J LAMB
510(k) history	169 submissions · 166 cleared · 1989-2022

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k960203/> 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 May 27, 2026