

K873740 ROTASCREEN EIANov 27, 1987
73 days to decisionK873740 · Product code: **LIQ** · Microbiology
Source: <https://www.510kdatabase.net/k873740/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Enzyme Linked Immunoabsorbent Assay, Rotavirus (LIQ)
Date received	Sep 15, 1987
Decision date	Nov 27, 1987
Days to decision	73 days
Third-party review	No

APPLICANT

Company	Mercia Diagnostics , Ltd.
Location	Guildford, Surrey England, GB
Contact	LEWIS (PHD)
510(k) history	17 submissions · 17 cleared · 1987-1992

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k873740/>, 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 30, 2026