

K833829 TOXOPLASMA GOND II IGG DIA TEST KITApr 24, 1984
173 days to decisionK833829 · Product code: **LGD** · Microbiology
Source: <https://www.510kdatabase.net/k833829/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Enzyme Linked Immunoabsorbent Assay, Toxoplasma Gondii (LGD)
Date received	Nov 3, 1983
Decision date	Apr 24, 1984
Days to decision	173 days
Third-party review	No

APPLICANT

Company	Labsystems, Inc.
Location	Walker, MI, US
510(k) history	17 submissions · 17 cleared · 1984-1993

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Device record: <https://www.510kdatabase.net/k833829/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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