

K831363 TESTCORP ASOMay 16, 1983
20 days to decisionK831363 · Product code: **GTQ** · Microbiology
Source: <https://www.510kdatabase.net/k831363/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Antistreptolysin - Titer/streptolysin O Reagent (GTQ)
Date received	Apr 26, 1983
Decision date	May 16, 1983
Days to decision	20 days
Third-party review	No

APPLICANT

Company	Catheter Technology Corp.
Location	Walker, MI, US
510(k) history	13 submissions · 13 cleared · 1983-1988

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Device record: <https://www.510kdatabase.net/k831363/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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