FDA 510(k) Premarket Notification

K831363 TESTCORP ASO

CLEARED

May 16, 1983
20 days to decision

K831363 · 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