FDA 510(k) Premarket Notification

K961829 QUALITROL CRP CONTROL SET

CLEARED

Aug 19, 1996
98 days to decision

K961829 · Product code: **DCK** · Immunology
Source: https://www.510kdatabase.net/k961829/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	C-reactive Protein, Antigen, Antiserum, And Control (DCK)
Date received	May 13, 1996
Decision date	Aug 19, 1996
Days to decision	98 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Consolidated Technologies, Inc.**
Location	Austin, TX, US
Contact	WILLIAM CONE
510(k) history	20 submissions · 20 cleared · 1991-2000

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k961829/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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