FDA 510(k) Premarket Notification

K060120 TECO CX3 REAGENT SET FOR SYNCHRON CX SYSTEM

CLEARED

Mar 27, 2006
69 days to decision

K060120 · Product code: **CDQ** · Chemistry
Source: https://www.510kdatabase.net/k060120/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Urease And Glutamic Dehydrogenase, Urea Nitrogen (CDQ)
Date received	Jan 17, 2006
Decision date	Mar 27, 2006
Days to decision	69 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Teco Diagnostics**
Location	Placentia, CA, US
Contact	JIAN VAECHES
510(k) history	51 submissions · 51 cleared · 1993-2017