

K060120 TECO CX3 REAGENT SET FOR SYNCHRON CX SYSTEMMar 27, 2006
69 days to decisionK060120 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k060120/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 31, 2026