FDA 510(k) Premarket Notification

CLEARED

K091153 SYNCHRON SYSTEMS ENZYMATIC CO2 (CO2E) REAGENT, MODEL A60291

Jul 13, 2009
83 days to decision

K091153 · Product code: **KHS** · Chemistry
Source: https://www.510kdatabase.net/k091153/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Enzymatic, Carbon-dioxide (KHS)
Date received	Apr 21, 2009
Decision date	Jul 13, 2009
Days to decision	83 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Beckman Coulter, Inc.**
Location	Chaska, MN, US
Contact	MARINE BOYAJIAN
Website	https://www.beckmancoulter.com
510(k) history	270 submissions · 270 cleared · 1993-2026

Beckman Coulter, Inc. is a diagnostic device manufacturer headquartered in Chaska, US. The company specializes in clinical laboratory and immunodiagnostic systems. Beckman Coulter has received FDA 510(k) clearances from total submissions since its first clearance in 1993. The company maintains active regulatory status, with the latest clearance in 2026. Its portfolio spans chemistry devices, microbiology testing systems, hematology analyzers, and immunoassay platforms. Recent cleared devices include chemistry assays for cardiac markers, microbiology susceptibility panels,...