FDA 510(k) Premarket Notification

K894173 BUN-BLOOD UREA NITROGEN FOR MANUAL/AUTO APPLICAT

CLEARED

Sep 5, 1989
82 days to decision

K894173 · Product code: **CDQ** · Chemistry
Source: https://www.510kdatabase.net/k894173/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Urease And Glutamic Dehydrogenase, Urea Nitrogen (CDQ)
Date received	Jun 15, 1989
Decision date	Sep 5, 1989
Days to decision	82 days
Third-party review	No

APPLICANT

Company	**Catachem, Inc.**
Location	Port Chester, NY, US
Contact	LUIS P LEON
510(k) history	41 submissions · 41 cleared · 1985-2007

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k894173/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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