

K831958 RADIAL IMMUNO-TEST FOR HUMAN C-REACTIJul 19, 1983
32 days to decisionK831958 · Product code: **DCK** · Chemistry
Source: <https://www.510kdatabase.net/k831958/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	C-reactive Protein, Antigen, Antiserum, And Control (DCK)
Date received	Jun 17, 1983
Decision date	Jul 19, 1983
Days to decision	32 days
Third-party review	No

APPLICANT

Company	Kent Laboratories, Inc.
Location	Mchenry, IL, US
510(k) history	40 submissions · 37 cleared · 1977-1995

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Device record: <https://www.510kdatabase.net/k831958/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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