

K811423 VENTRE/SEPJun 9, 1981
19 days to decisionK811423 · Product code: **CFP** · Chemistry
Source: <https://www.510kdatabase.net/k811423/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioimmunoassay, Immunoreactive Insulin (CFP)
Date received	May 21, 1981
Decision date	Jun 9, 1981
Days to decision	19 days
Third-party review	No

APPLICANT

Company	Ventrex Laboratories, Inc.
Location	Mchenry, IL, US
510(k) history	82 submissions · 82 cleared · 1979-1992

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Device record: <https://www.510kdatabase.net/k811423/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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