

K791896 CENTRIA PHENYTOIN RIA TEST SETOct 17, 1979
22 days to decisionK791896 · Product code: **DLP** · Toxicology
Source: <https://www.510kdatabase.net/k791896/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioimmunoassay, Diphenylhydantoin (DLP)
Date received	Sep 25, 1979
Decision date	Oct 17, 1979
Days to decision	22 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/k791896/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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