FDA 510(k) Premarket Notification

K791896 CENTRIA PHENYTOIN RIA TEST SET

CLEARED

Oct 17, 1979
22 days to decision

K791896 · 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