FDA 510(k) Premarket Notification

K921099 SYNTRON QUIKSTRIP/E.Z. STRIP PREGNANCY HOME TEST

CLEARED

May 29, 1992
81 days to decision

K921099 · Product code: **LCX** · Chemistry
Source: https://www.510kdatabase.net/k921099/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Kit, Test, Pregnancy, Hcg, Over The Counter (LCX)
Date received	Mar 9, 1992
Decision date	May 29, 1992
Days to decision	81 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Syntron Bioresearch, Inc.**
Location	San Diego, CA, US
Contact	BENEDICT ZIN
510(k) history	26 submissions · 26 cleared · 1987-1999

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k921099/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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