FDA 510(k) Premarket Notification

CLEARED

K150791 Healgen Secobarbital Test(Strip, Cassette, Cup, Dip Card), Healgen Burprenorphine Test (Strip, Cassette, Cup, Dip Card), Healgen Methadone Test (Strip, Cassette, Cup, Dip Card)

Apr 24, 2015
30 days to decision

K150791 · Product code: **DIS** · Toxicology
Source: https://www.510kdatabase.net/k150791/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Enzyme Immunoassay, Barbiturate (DIS)
Date received	Mar 25, 2015
Decision date	Apr 24, 2015
Days to decision	30 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Healgen Scientific, LLC**
Location	Houston, TX, US
Contact	JIANQIU FANG
Website	https://www.healgen.com
510(k) history	27 submissions · 27 cleared · 2012-2026

Healgen Scientific, LLC is a leading in-vitro diagnostics (IVD) developer and manufacturer based in Houston, Texas. Established in 2007, the company specializes in high-quality diagnostic testing technologies across multiple therapeutic areas. Healgen has achieved FDA 510(k) clearances from total submissions since 2012, with no denied submissions on record. The company's regulatory portfolio is dominated by toxicology devices, including drug screening and fentanyl detection products, alongside offerings in chemistry, microbiology, and infectious disease diagnostics. The l...

REGULATORY CONSULTANT

Consulting firm	**LSI International, Inc.**
Contact	JOE SHIA

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k150791/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 13, 2026