

K260355 Webest Multi-Drug Urine CupMar 9, 2026
34 days to decisionK260355 · Product code: **NFT** · Toxicology
Source: <https://www.510kdatabase.net/k260355/>**SUBMISSION DETAILS**

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|-----------------------|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Test, Amphetamine, Over The Counter (NFT) |
| Date received | Feb 3, 2026 |
| Decision date | Mar 9, 2026 |
| Days to decision | 34 days |
| Third-party review | No |
| Combination product | No |
| PCCP authorized | No |
| Summary / Statement | Summary |
| Other names | Webest Home Multi-Drug Urine Cup |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | WEBEST Biotech,, LLC |
| Location | Irwindale, CA, US |
| Contact | Wei Midori |
| 510(k) history | 1 submissions · 1 cleared · 2026-2026 |

REGULATORY CONSULTANT

| | |
|-----------------|--------------------------------|
| Consulting firm | LSI International, Inc. |
| Contact | Jenny Xia |

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k260355/> 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 14, 2026