

**K252306 Verrix EVA™ STEAM Biological Indicator (BI) (Model: 1200), Verrix EVA™ STEAM Process Challenge Device (PCD) (Model: 1300) combined with Verrix EVA™ Auto-Reader (Model: 1100)**

Apr 22, 2026  
272 days to decision

K252306 · Product code: **QVB** · General Hospital  
Source: <https://www.510kdatabase.net/k252306/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Biological Sterilization Indicator With Indirect Growth Detection (QVB)
Date received	Jul 24, 2025
Decision date	Apr 22, 2026
Days to decision	272 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Steritec Products</b>
Location	Engelwood, CO, US
Contact	Darshan Yeliyur Siddegowda
510(k) history	1 submissions · 1 cleared · 2026-2026

**REGULATORY CONSULTANT**

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Consulting firm	<b>Getinge</b>
Contact	Barb Smith

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k252306/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://accessdata.fda.gov)).

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