

**K251216 0184 CO2 Sampling line**Jun 4, 2025  
47 days to decisionK251216 · Product code: **CCK** · Anesthesiology  
Source: <https://www.510kdatabase.net/k251216/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Gas, Carbon-dioxide, Gaseous-phase (CCK)
Date received	Apr 18, 2025
Decision date	Jun 4, 2025
Days to decision	47 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary
Other names	0184T CO2 Sampling line; 0139 CO2 Sampling line; 0182 CO2 Sampling line

**APPLICANT**

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Company	<b>Barbaras Development, Inc.</b>
Location	Newbury Park, CA, US
Contact	Matthew Baile
Website	<a href="https://www.bd.inc/">https://www.bd.inc/</a>
510(k) history	1 submissions · 1 cleared · 2025-2025

Barbaras Development, Inc. is a medical device manufacturer with a manufacturing facility in Newbury Park, US. The company specializes in product development, design, tooling, and manufacturing services across multiple industries. The company has received FDA 510(k) clearance from total submission. Barbaras Development focuses exclusively on Anesthesiology devices, with its first and latest clearance both occurring in 2025. The company remains active in FDA regulatory submissions. Explore the company's cleared device names, product codes, and clearance dates in the database above.

**REGULATORY CONSULTANT**

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Consulting firm	<b>ProMedic, LLC</b>
Contact	Paul Dryden

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/k251216/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://accessdata.fda.gov)).

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