

**K192035 EvaQMax Smoke Evacuation System**Aug 29, 2019  
30 days to decisionK192035 · Product code: **FYD** · General Hospital  
Source: <https://www.510kdatabase.net/k192035/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Apparatus, Exhaust, Surgical (FYD)
Date received	Jul 30, 2019
Decision date	Aug 29, 2019
Days to decision	30 days
Third-party review	Yes
Combination product	No
PCCP authorized	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Bio Protech, Inc.</b>
Location	Sparks, NV, US
Contact	Namhyun Cho
510(k) history	17 submissions · 17 cleared · 2002-2019

**REGULATORY CONSULTANT**

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Consulting firm	<b>Regulatory Technology Services, LLC</b>
Contact	Prithul Bom

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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