

**K203450 EsophaCap Swallowable Cellular Retrieval Device
(changed from Cell-Mata)**May 4, 2021
162 days to decisionK203450 · Product code: **EOX** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k203450/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Esophagoscope (flexible Or Rigid) (EOX)
Date received	Nov 23, 2020
Decision date	May 4, 2021
Days to decision	162 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Capnostics, LLC
Location	Concord, NC, US
Contact	Martin von Dyck
510(k) history	1 submissions · 1 cleared · 2021-2021

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k203450/> 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 26, 2026