

K183262 EsoCheck CCD Cell Collection DeviceJun 21, 2019
210 days to decisionK183262 · Product code: **EOX** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k183262/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent - K
Submission type	Traditional
Device classification	Esophagoscope (flexible Or Rigid) (EOX)
Date received	Nov 23, 2018
Decision date	Jun 21, 2019
Days to decision	210 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Lucid Diagnostics, Inc.
Location	New York, NY, US
Contact	Lishan Aklog
510(k) history	4 submissions · 0 cleared · 2019-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k183262/> 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