FDA 510(k) Premarket Notification

K183262 EsoCheck CCD Cell Collection Device

CLEARED

Jun 21, 2019
210 days to decision

K183262 · Product code: **EOX** · Gastroenterology & Urology
Source: 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