

K201952 CrossGlide ETS PlusAug 5, 2020
22 days to decisionK201952 · Product code: **HHK** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k201952/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Curette, Suction, Endometrial (and Accessories) (HHK)
Date received	Jul 14, 2020
Decision date	Aug 5, 2020
Days to decision	22 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Crossbay Medical
Location	San Diego, CA, US
Contact	Piush Vidyarthi
510(k) history	3 submissions · 3 cleared · 2019-2020

REGULATORY CONSULTANT

Consulting firm	Domecus Consulting Services, LLC
Contact	Cindy Domecus

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov

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