

K210613 HIT Clear AlignerJun 4, 2021
95 days to decisionK210613 · Product code: **NXC** · Dental
Source: <https://www.510kdatabase.net/k210613/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Aligner, Sequential (NXC)
Date received	Mar 1, 2021
Decision date	Jun 4, 2021
Days to decision	95 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Cdb Corporation
Location	Leland, NC, US
Contact	Leah M Lehman
510(k) history	6 submissions · 6 cleared · 2008-2022

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k210613/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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