

K191209 Aneroid Type Blood Pressure KitJun 18, 2019
43 days to decisionK191209 · Product code: **DXQ** · Cardiovascular
Source: <https://www.510kdatabase.net/k191209/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Blood Pressure Cuff (DXQ)
Date received	May 6, 2019
Decision date	Jun 18, 2019
Days to decision	43 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Dongguan Mars Medical Products Co., Ltd.
Location	Dani Village, Shatian Town, Dongguan Cit, CN
Contact	Lily Liao
510(k) history	1 submissions · 1 cleared · 2019-2019

REGULATORY CONSULTANT

Consulting firm	Guangzhou Junyi Information Technology Co., Ltd.
Contact	Jinghua Zhou

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

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