

**K141018 ORAL-B TEST DRIVE POWER BRUSH TRIAL
PROGRAM KIT, ORAL-B TEST DRIVE POWER TOOTHBRUSH
KIT, ORAL-B TEST DRIVE REFILL KIT**Sep 15, 2014
147 days to decisionK141018 · Product code: JEQ · Dental
Source: <https://www.510kdatabase.net/k141018/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Toothbrush, Powered (JEQ)
Date received	Apr 21, 2014
Decision date	Sep 15, 2014
Days to decision	147 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Procter & Gamble Co.
Location	Cincinnati, OH, US
Contact	MICHAEL KAMINSKI, PH.D.
Website	http://www.pg.com/
510(k) history	23 submissions · 23 cleared · 1988-2014

Procter & Gamble Co. is a consumer health and personal care company headquartered in Cincinnati, US. The company develops and markets a broad range of health and wellness products globally. Procter & Gamble has received FDA 510(k) clearances from total submissions, with no denied submissions on record. The company's regulatory focus has centered on Obstetrics & Gynecology devices, which represent the dominant category of its submissions. FDA 510(k) clearances span from 1988 to 2014, establishing a historical regulatory record in feminine care and oral health device categories.