

# **SECURITIES & EXCHANGE COMMISSION EDGAR FILING**

# **BIOVIE INC.**

Form: 8-K

Date Filed: 2019-06-28

Corporate Issuer CIK: 1580149

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## **UNITED STATES SECURITIES AND EXCHANGE COMMISSION**

WASHINGTON, D.C. 20549

FORM 8-K

### **CURRENT REPORT PURSUANT** TO SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934

Date of Report (Date of earliest event reported) June 26, 2019

## **BioVie Inc.**

(Exact Name of Registrant as Specified in Its Charter)

Nevada	000-55292	46-2510769
(State or Other Jurisdiction of Incorporation)	(Commission File Number)	(I.R.S. Employer Identification No.)
2120 Colorado Avenue, #230 Santa Monica, California		90404
(Address of Principal Executive Offices)		(Zip Code)
(Re	(310) 444-4300 gistrant's Telephone Number, Including Area	Code)
,	Name or Former Address, if Changed Since L	, ,
Title of each class	Trading Symbol(s)	Name of each exchange on which registered
Check the appropriate box below if the Form 8-h following provisions (see General Instruction A.2. below)  Written communications pursuant to Rule 425 under  Soliciting material pursuant to Rule 14a-12 under the Pre-commencement communications pursuant to Rule Pre-commencement pursuant	the Securities Act (17 CFR 230.425) Exchange Act (17 CFR 240.14a-12) ule 14d-2(b) under the Exchange Act (17 CFR	` ''
Indicate by check mark whether the registrant is (§230.405 of this chapter) or Rule 12b-2 of the Securities		as defined in Rule 405 of the Securities Act of 1933 apter).
Emerging growth company $oxtimes$		
If an emerging growth company, indicate by chec new or revised financial accounting standards provided p	S .	e the extended transition period for complying with any . $\Box$

#### Item 8.01 Other Events.

On June 18, 2019, BioVie Inc., a Nevada corporation (the "Company") met with representatives of the US Food and Drug Administration ("FDA") for a Type C Guidance Meeting to plan the Company's next clinical study following the recently completed Phase 2a clinical trial. Company attendees discussed the Company's clinical development efforts with the FDA and proposed trial endpoints. While the FDA has not provided final guidance nor does the Company have certainty as to what that guidance would entail, the Company's goal remains to proceed into a Phase 2b/3 or Phase 3 clinical trial in a manner consistent with what was reviewed with the FDA. The Company may still need to address certain risks associated with unvalidated quality of life measures. The FDA is expected to provide its final meeting minutes, which the Company expects will include guidance on several issues discussed at the meeting, within 30 days of the meeting.

### **SIGNATURES**

Pursuant to the requirements of the Securities Exchange Act of 1934, the Registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

Dated: June 28, 2019

#### BIOVIE INC.

By: <u>/s/ Wendy Kim</u> Name: Wendy Kim Title: Chief Financial Officer