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): March 14, 2018

PROPANC BIOPHARMA, INC.

(Exact name of registrant as specified in its charter)

Delaware	**000-54878**	**33-0662986**
(State or other Jurisdiction of Incorporation)	(Commission File Number)	(IRS Employer Identification No.)

302, 6 Butler Street Camberwell, VIC, 3124 Australia	
(Address of Principal Executive Offices)	(Zip Code)

Registrant's telephone number, including area code: **61 03 9882 6723**

(Former name or former address, if changed since last report.)

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions:

[ ] Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)

[ ] Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)

[ ] Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))

[ ] Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (§230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§240.12b-2 of this chapter).

Emerging growth company [ ]

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act. [ ]

Item 7.01 Regulation FD Disclosure

On March 14, 2018, Propanc Biopharma, Inc. (the "Company") announced the successful reproduction run of the manufacturing process for the Company's two drug substances typsinogen and chymotrypsinogen.

A press release relating to such event is attached as Exhibit 99.1 hereto.

The foregoing (including Exhibit 99.1) is being furnished pursuant to Item 7.01 and will not be deemed to be filed for purposes of Section 18 of the Securities and Exchange Act of 1934, as amended (the "Exchange Act"), or otherwise be subject to the liabilities of that section, nor will it be deemed to be incorporated by reference in any filing under the Securities Act of 1933, as amended, or the Exchange Act.

Item 9.01 Exhibits

(d) Exhibits.

Exhibit	Description
99.1	Press Release, dated as of March 14, 2018

SIGNATURE

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

Propanc Biopharma, Inc.

Date: March 14, 2018

By: */s/ James Nathanielsz*
James Nathanielsz
Chief Executive Officer, Chief Financial Officer and Chief Accounting Officer

Propanc Biopharma Completes Successful Reproduction Run for PRP

Company Requests Scientific Advice Meeting with the MHRA (UK) to Discuss Next Steps

MELBOURNE, AUSTRALIA, March 14, 2018 -- Propanc Biopharma Inc. (OTCQB: PPCB) ("Propanc Biopharma" or the "Company"), a clinical stage biopharmaceutical company focusing on development of new and proprietary treatments for cancer patients suffering from solid tumors such as pancreatic, ovarian and colorectal cancers, today announced the successful reproduction run of the manufacturing process for the Company's two drug substances trypsinogen and chymotrypsinogen. The successful reproduction run demonstrates scalability of Propanc´s proprietary manufacturing process to enable routine production of the two active substances for the Company's lead product candidate, PRP. The process was developed in collaboration with a European Contract Manufacturing Organization (CMO) experienced in the production of biopharmaceuticals.

As a result of the successful reproduction run, a scientific advice meeting was requested with the UK Medicines and Healthcare Products Regulatory Agency (MHRA) to inform them of the results from recent manufacturing activities and confirm requirements for the next stage of GMP manufacture for PRP. Given the unique potential of these active biological drug substances, management continues to engage in proactive communication with the regulators to advance the product candidate towards clinical and market approval.

"We are delighted with the results from the reproduction run, which shows that the proprietary process is scalable and robust so that we can produce our drug substances, routinely," said James Nathanielsz, Propanc Biopharma's Chief Executive Officer. "As a result, the Company requested a scientific advice meeting with the MHRA to update the agency about our progress."

"The Propanc management team continues to remain focused and diligent regarding the advancement of PRP into clinical trials," said Professor Klaus Kutz, Propanc Biopharma's Chief Medical Officer. "Our regulatory and development team are working hard to ensure we have good communication with the MHRA."

PRP is a proposed solution for once-daily intravenous administration of a combination of two pancreatic proenzymes trypsinogen and chymotrypsinogen. Currently progressing towards a First-In-Human study, PRP aims to prevent tumor recurrence and metastasis from solid tumors. Eighty percent of all cancers are solid tumors and metastasis is the main cause of patient death from cancer. According to the World Health Organization, 8.2 million people died from cancer in 2012. Consequently, a report by IMS Health states innovative therapies are driving the global oncology market to meet demand, which is expected to reach $150 billion by 2020. The Company's initial target patient populations are pancreatic, ovarian and colorectal cancers, representing an estimated combined market segment of $14 billion in 2020, according to GBI Research.

To view Propanc Biopharma's "Mechanism of Action" video on anti-cancer product candidate, PRP, please click on the following link: http://www.propanc.com/news-media/video

To be added to Propanc Biopharma's email distribution list, please click on the following link: http://ir.propanc.com/email-alerts and submit the online request form.

About Propanc Biopharma:

Propanc Biopharma is a clinical stage biopharmaceutical company developing new cancer treatments initially for patients suffering from pancreatic, ovarian and colorectal cancers. We have developed a formulation of anti-cancer compounds, which exert a number of effects designed to control or prevent tumors from recurring and spreading throughout the body. Our products involve or employ pancreatic proenzymes, which are inactive precursors of enzymes. In the near term, we intend to target patients with limited remaining therapeutic options for the treatment of solid tumors. In future, we intend to develop our lead product candidate to treat (i) early stage cancer and (ii) pre-cancerous diseases and (iii) as a preventative measure for patients at risk of developing cancer based on genetic screening. For more information, visit: www.propanc.com.

Forward-Looking Statements:

All statements other than statements of historical fact contained herein are "forward-looking statements" for purposes of federal and state securities laws. Forward-looking statements may include the words "may," "will," "estimate," "intend," "continue," "believe," "expect," "plan" or "anticipate" and other similar words. Although we believe that the expectations reflected in our forward-looking statements are reasonable, actual results could differ materially from those projected or assumed. Our future financial condition and results of operations, as well as any forward-looking statements, are subject to change and to inherent risks and uncertainties including those regarding our earnings, revenues and financial condition, our ability to implement our plans, strategies and objectives for future operations, our ability to execute on proposed new products, services or development thereof, our ability to establish and maintain the proprietary nature of our technology through the patent process, our ability to license from others patents and patent applications, if necessary, to develop certain products, our ability to implement our long range business plan for various applications of our technology, our ability to enter into agreements with any necessary manufacturing, marketing and/or distribution partners for purposes of commercialization, the results of our clinical research and development, competition in the industry in which we operate, overall market conditions, and any statements or assumptions underlying any of the foregoing. Other risks, uncertainties and factors that could cause actual results to differ materially from those projected may be described from time to time in reports we file with the Securities and Exchange Commission, including our reports on Forms 10-K, 10-Q and 8-K. We do not intend, and undertake no obligation, to update any forward-looking statement contained herein, except as required by law.

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