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Propanc Biopharma's Intellectual Property Portfolio Undergoes Rapid Growth

Over Sixty-Five Patents Either in Force or Pending Covering the Company's Technology

MELBOURNE, Australia--(BUSINESS WIRE)-- **Propanc Biopharma, Inc.** (OTC: PPCBD) ("Propanc"), a biopharmaceutical company developing new cancer treatments for patients suffering from recurring and metastatic cancer, announced today that Propanc's intellectual property ("IP") portfolio has undergone rapid growth recently, with sixty-five patents currently either in force or pending in most major countries and regions around the world. The IP covers Propanc's underlying anti-cancer technology in development. In the past year, three additional patent families entered national phase, where a patent application is filed in individual countries and regions, in order to achieve grant status.

"Our IP portfolio is a world leader in the field of administration of proenzymes for the treatment and prevention of metastatic cancer. IP is the cornerstone of any biotech company, and places us in a strong position to unlock significant value, as we plan to progress into a First-In-Human study and then hopefully towards proof of concept in a specific cancer indication, where we will then look at possible licensing opportunities," said Dr. Julian Kenyon, Propanc's Chief Scientific Officer. "We are firmly of the opinion that our current market value does not truly reflect the advancement of our lead product, PRP, or its true potential as a breakthrough, long term treatment for aggressive and fast spreading cancers from solid tumors by targeting and eradicating cancer stem cells."

Propanc has a total of four patent families covering the use of proenzymes for treating cancer via a combination of trypsinogen and/or chymotrypsinogen pancreatic proenzymes, which includes Propanc's lead product candidate, PRP. The IP portfolio includes composition of matter claims and a method of treatment to eradicate cancer stem cells. Further applications are also expected relating to the formulation and methods of use, as well as research programs with its joint research partners designed to further optimize the quality, safety and performance of PRP.

About Propanc Biopharma, Inc.

Propanc Biopharma, Inc. (the "Company") is developing a novel approach to prevent recurrence and metastasis of solid tumors by using pancreatic proenzymes that target and eradicate cancer stem cells in patients suffering from pancreatic, ovarian and colorectal cancers. For more information, please visit www.propanc.com.

The Company's novel proenzyme therapy is based on the science that enzymes stimulate biological reactions in the body, especially enzymes secreted by the pancreas. These pancreatic enzymes could represent the body's primary defense against cancer.

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

Forward-Looking Statements

All statements other than statements of historical facts contained in this press release are "forward-looking statements," which may often, but not always, be identified by the use of such words as "may," "might," "will," "will likely result," "would," "should," "estimate," "plan," "project," "forecast," "intend," "expect," "anticipate," "believe," "seek," "continue," "target" or the negative of such terms or other similar expressions. These statements involve known and unknown risks, uncertainties and other factors, which may cause actual results, performance or achievements to differ materially from those expressed or implied by such statements. These factors include uncertainties as to the Company's ability to continue as a going concern absent new debt or equity financings; the Company's current reliance on substantial debt financing that it is unable to repay in cash; the Company's ability to successfully remediate material weaknesses in its internal controls; the Company's ability to reach research and development milestones as planned and within proposed budgets; the Company's ability to control costs; the Company's ability to obtain adequate new financing on reasonable terms; the Company's ability to successfully initiate and complete clinical trials and its ability to successful develop PRP, its lead product candidate; the Company's ability to obtain and maintain patent protection; the Company's ability to recruit employees and directors with accounting and finance expertise; the Company's dependence on third parties for services; the Company's dependence on key executives; the impact of government regulations, including FDA regulations; the impact of any

future litigation; the availability of capital; changes in economic conditions, competition; and other risks, including, but not limited to, those described in the Company's Registration Statement on Form S-1, Amendment No. 1, filed with the U.S. Securities and Exchange Commission (the "SEC") on June 14, 2019, and in the Company's other filings and submissions with the SEC. These forward-looking statements speak only as of the date hereof and the Company disclaims any obligations to update these statements except as may be required by law.

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Investor Relations and Media:
Parker Mitchell
Propanc Biopharma, Inc.
Investor Relations
irteam@propanc.com
+61-03-9882-6723

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