



WPD Pharmaceuticals Secures Berubicin Sub-License Rights in Perpetuity

Commercially Reasonable Development Expenditures of at least US\$2,000,000 going towards the development, testing, regulatory approval or commercialization of the licensed product, Berubicin, during the applicable development period

Vancouver, British Columbia / February 1, 2022 – WPD Pharmaceuticals Inc. (the “Company” or “WPD”) (CSE: WBIO) (FSE: 8SV1), announced that in respect to its sublicense agreement with CNS Pharmaceuticals, Inc. (the “CNS Agreement”) on December 23, 2021, CNS confirmed that the Company has used “commercially reasonable development efforts” towards the development of Berubicin, defined as expenditures of at least USD\$2,000,000 on the development, testing, regulatory approval or commercialization of the licensed product during the applicable development period, and as such, the Company, through WPD Poland, is entitled to maintain its sublicense of Berubicin subject to the ongoing obligations under the CNS Agreement. The Company’s sub-license of Berubicin allows it geographic exclusivity for development and marketing in a region consisting of select countries in Eastern Europe and Central Asia.

Berubicin is an anthracycline, a class of anticancer agents that are among the most powerful chemotherapy drugs and effective against more types of cancer than any other class of chemotherapeutic agents. Anthracyclines are designed to utilize natural processes to induce deoxyribonucleic acid (DNA) damage in targeted cancer cells by interfering with the action of topoisomerase II, a critical enzyme enabling cell proliferation. Berubicin treatment of brain cancer patients appeared to demonstrate positive responses that include one durable complete response in a Phase 1 human clinical trial conducted by Reata Pharmaceuticals, Inc. Berubicin, was developed by Dr. Waldemar Priebe, Professor of Medicinal Chemistry at The University of Texas MD Anderson Cancer Center.

About WPD Pharmaceuticals

WPD is a biotechnology research and development company with a focus on oncology and virology, namely research and development of medicinal products involving biological compounds and small molecules. WPD has licensed in certain countries 10 novel drug candidates with 4 that are in clinical development stage. These drug candidates were researched at institutions, and WPD currently has ongoing collaborations with Wake Forest University and leading hospitals and academic centers in Poland.

WPD has entered into license agreements with Wake Forest University Health Sciences and sublicense agreements with Moleculin Biotech, Inc. and CNS Pharmaceuticals, Inc., respectively, each of which grant WPD an exclusive, royalty-bearing sublicense to certain technologies of the licensor. Such agreements provide WPD with certain research, development, manufacturing and sales rights, among other things. The sublicense territory from CNS Pharmaceuticals and Moleculin Biotech includes for most compounds 30 countries in Europe and Asia, including Russia.

On Behalf of the Board

'Mariusz Olejniczak'

Mariusz Olejniczak
CEO, WPD Pharmaceuticals

Contact

Investor Relations
Email: investors@wpdpharmaceuticals.com
Tel: 604-428-7050
Web: www.wpdpharmaceuticals.com

Neither the Canadian Securities Exchange nor the Investment Industry Regulatory Organization of Canada accepts responsibility for the adequacy or accuracy of this release.

This press release contains forward-looking statements. Forward-looking statements are statements that contemplate activities, events or developments that the Company can develop effective drugs against cancer and possibly viruses. Factors which may prevent the forward looking statement from being realized include that competitors or others may successfully challenge a granted patent and the patent could be rendered void; we may be unable to raise sufficient funding for our research; we may be unable to expend sufficient funds on research to keep our sublicense rights; our grant applications may not be successful or if successful, we may not meet the requirements to receive the grants awarded; that our drugs don't provide positive treatment, or if they do, the side effects are damaging; competitors may develop better or cheaper drugs; and we may be unable to obtain regulatory approval for any drugs we develop; the filing of the next annual general meeting information circular containing the executive compensation disclosure. Readers should refer to the risk disclosure included from time-to-time in the documents the Company files on SEDAR, available at www.sedar.com. Although the Company believes that the assumptions inherent in these forward-looking statements are reasonable, they are not guarantees of future performance and, accordingly, they should not be relied upon and there can be no assurance that any of them will prove to be accurate. Finally, these forward-looking statements are made as of the date of this press release and the Company assumes no obligation to update them except as required by applicable law.

