

For Immediate Release December 8, 2020

## SUNSHINE BIOPHARMA ORDERS A NEW BATCH OF Adva-27a FOR TESTING ON TOPII AMPLIFIED CANCERS

Montreal, Quebec, Canada -- (ACCESSWIRE) -- Sunshine Biopharma Inc. (OTC PINK: "SBFM"), a pharmaceutical company focused on the research, development and commercialization of oncology and antiviral drugs, today announced that it has ordered a new batch of Adva-27a from its manufacturer in China. The material will be delivered to the laboratories of Sunshine Biopharma's drug development partner in Montreal (Canada). Sunshine plans to use the material to conduct testing on various types of cancer cells that overproduce Topoisomerase II (TopII). Following these cell culture studies, Sunshine will proceed to performing studies on mice harboring tumors of human origin with TopII amplification. TopII is an essential cell cycle enzyme that is amplified in approximately 4% of all human cancers.

"All cancer types that have amplification of the Topoisomerase II gene are potential new targets for our Adva-27a," said Dr. Steve Slilaty, CEO of Sunshine Biopharma. "This is an additional application for our Adva-27a beyond the original multidrug resistance indication," he added.

## **About Sunshine Biopharma**

In addition, to working on the development of a treatment for COVID-19, Sunshine Biopharma is engaged in the development Adva-27a, a unique anticancer compound. Tests conducted to date have demonstrated the effectiveness of Adva-27a at destroying Multidrug Resistant Cancer Cells, including Pancreatic Cancer cells, Small-Cell Lung Cancer cells, Breast Cancer cells, and Uterine Sarcoma cells. Clinical trials for Pancreatic Cancer indication are planned to be conducted at McGill University's Jewish General Hospital in Montreal, Canada. Sunshine Biopharma is owner of all patents and intellectual property pertaining to Adva-27a.

## Safe Harbor Forward-Looking Statements

This press release may contain forward looking statements which are based on current expectations, forecasts, and assumptions that involve risks as well as uncertainties that could cause actual outcomes and results to differ materially from those anticipated or expected, including statements related to the amount and timing of expected revenues statements related to our financial performance, expected income, distributions, and future growth for upcoming quarterly and annual periods. These risks and uncertainties are further defined in filings and reports by the Company with the U.S. Securities and Exchange Commission (SEC). Actual results and the timing of certain

events could differ materially from those projected in or contemplated by the forward-looking statements due to a number of factors detailed from time to time in our filings with the SEC. Among other matters, the Company may not be able to sustain growth or achieve profitability based upon many factors including but not limited to general stock market conditions. Reference is hereby made to cautionary statements set forth in the Company's most recent SEC filings. We have incurred and will continue to incur significant expenses in our expansion of our existing as well as new service lines noting there is no assurance that we will generate enough revenues to offset those costs in both the near and long term. Additional service offerings may expose us to additional legal and regulatory costs and unknown exposure(s) based upon the various geopolitical locations we will be providing services in, the impact of which cannot be predicted at this time.

## For Additional Information Contact:

Camille Sebaaly, CFO Sunshine Biopharma Inc. Direct Line: 514-814-0464

camille.sebaaly@sunshinebiopharma.com

www.sunshinebiopharma.com