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## **BIONXT SOLUTIONS ANNOUNCES ODF CLADRIBINE UPDATE AND FINANCING**

**Vancouver, Canada (April 25, 2024)** - BioNxt Solutions Inc. (“**BioNxt**” or the “**Company**”) (CSE: BNXT / OTC: BNXTF / FSE: BXT) is pleased to announce that its Cladribine program is advancing to human comparative bioavailability studies in Europe. The Company is reviewing proposals from several contract research organizations to carry out the studies in accordance with EU medical regulatory guidelines. The recent success of both the Company’s ODF Cladribine toxicity and comparative pharmacokinetic (“PK”) studies and industry interest have set high expectations for the upcoming comparative bioavailability study. Further details on the study will be released in due course.

BioNxt is developing a 100% owned and proprietary ODF Cladribine dosage form, directed at the multiple sclerosis (“MS”) market. Cladribine tablets are currently approved for use in over 75 countries, including by the United States Food and Drug Administration (“FDA”) and the European Medicines Agency (“EMA”), with annual sales in excess of one billion USD. Cladribine tablets are approved for several indications, namely highly active forms of relapsing-remitting MS and certain forms of leukemia. MS represents the largest market segment for the sale of Cladribine with approximately 2.3 million people living with MS worldwide, with the highest prevalence in North America and Europe, noted by Atlas of MS. The global Multiple Sclerosis drug market is expected to top USD 41 billion by 2033 according to Market.us.

The Company has filed Cladribine ODF-related provisional patent applications with three to four patent applications expected to be on file in major international jurisdictions by late 2024 to early 2025 with potential patent protection extending to 2044.

The Company is also pleased to announce a non-brokered private placement of up to 6,000,000 common shares of the Company (the “**Common Shares**”) at a price of \$0.27 per Common Share for gross proceeds of up to \$1,620,000 (the “**Offering**”). The Company intends to use the net proceeds from the Offering for research, development, and commercialization programs and general working capital purposes. The Offering may close on one or more dates as the Company may determine.

The Company may pay a finder’s fee in connection with the Offering to eligible arm’s length finders in accordance with the policies of the Canadian Securities Exchange. All securities issued in connection with the Offering will be subject to a statutory hold period of four months and one day following the date of issuance in accordance with applicable Canadian securities laws.

The securities issued pursuant to the Offering have not, nor will they be registered under the United States Securities Act of 1933, as amended, and may not be offered or sold within the United States or to, or for the account or benefit of, U.S. persons in the absence of U.S. registration or an applicable exemption from the U.S. registration requirements. This news release shall not constitute an offer to sell or the solicitation of an offer to buy nor shall there be any sale of the securities in the United States or in any other jurisdiction in which such offer, solicitation or sale would be unlawful.



### **About BioNxt Solutions Inc.**

BioNxt Solutions Inc. is a bioscience accelerator focused on next-generation drug formulations and delivery systems, diagnostic screening tests, and new active pharmaceutical production and evaluation, including: precision transdermal and oral dissolvable drug formulations; rapid, low-cost infectious disease and oral health screening tests; and standardization and clinical evaluation of emerging active pharmaceutical ingredients for neurological applications. The Company has research and development operations in North America and Europe, with an operational focus in Germany, and is currently focused on regulatory approval and commercialization of medical products for European markets.

### **BioNxt Solutions Inc.**

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*This news release includes certain statements that may be deemed “forward-looking statements”. All statements in this new release, other than statements of historical facts, that address events or developments that the Company expects to occur, are forward-looking statements. Forward-looking statements are statements that are not historical facts and are generally, but not always, identified by the words “expects”, “plans”, “anticipates”, “believes”, “intends”, “estimates”, “projects”, “potential” and similar expressions, or that events or conditions “will”, “would”, “may”, “could” or “should” occur. Although the Company believes the expectations expressed in such forward-looking statements are based on reasonable assumptions, such statements are not guarantees of future performance and actual results may differ materially from those in the forward-looking statements. Factors that could cause the actual results to differ materially from those in forward-looking statements include market prices, continued availability of capital and financing, and general economic, market or business conditions. Investors are cautioned that any such statements are not guarantees of future performance and actual results or developments may differ materially from those projected in the forward-looking statements. Forward-looking statements are based on the beliefs, estimates and opinions of the Company’s management on the date the statements are made. Except as required by applicable securities laws, the Company undertakes no obligation to update these forward-looking statements in the event that management’s beliefs, estimates or opinions, or other factors, should change.*

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