

Isodiol International Inc. Announces Letter of Intent to Supply 99.5%+ Pharma-Grade Cannabidiol to Zenabis

With the MHRA approval for the manufacturing of the CBD Isolate, Isodiol is able to export its pure, natural CBD from the United Kingdom into global markets

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VANCOUVER, British Columbia, May 04, 2018 (GLOBE NEWSWIRE) -- Isodiol International Inc. (CSE:ISOL) (OTC:ISOL<u>F</u>) (FSE:LB6A.F) (the "Company" or "Isodiol"), a global innovator specializing in the development of pharmaceutical and wellness products, is pleased to announce that it has signed a Letter of Intent ("LOI") with Zenabis Ltd. ("Zenabis"), one of Canada's largest Licensed Producers, to import CBD isolate as an Active Pharmaceutical Ingredient, into Canada from Isodiol's GMP-certified production facility in the United Kingdom.

Zenabis is a Canadian biopharmaceutical company positioned to become one of the largest cannabis producers in Canada, with facilities totalling more than 400,000 square feet of cannabis production space. Zenabis intends to import a minimum of 3,000 g of such CBD isolate per month for the purposes of R&D and new product formulation.

Marcos Agramont, CEO of Isodiol, stated: "Isodiol is pleased to announce another supply agreement for its highest quality, pharma grade CBD isolate. Zenabis's team is working on various R&D projects focused on developing new products and applications, and we are pleased to be a key partner in enabling this R&D process."

"We are excited to move forward with this partnership," explained Kevin Coft, CEO of Zenabis. "To have a CBD isolate accompanied by an API status sets it apart from every other global source, and we look forward to be working with Isodiol in Canada on new drug development, approved pharmaceutical applications, research, clinical studies and trials."

Having met all requirements, Isodiol recently announced that the Company's wholly owned subsidiary BSPG Laboratories LTD. has received government approval from United Kingdom's Medicines and Healthcare Products Regulatory Agency (MHRA), in accordance with The

Human Medicines Regulations 2012 (SI 2012/1916), for the manufacturing of the active

substance Cannabidiol (CBD) under certificate number: UK API 48727.

Active ingredients are the substances in drugs that are responsible for the beneficial health

effects experienced by consumers. Regulating active ingredients helps to increase the quality

and safety of drugs for consumers and Isodiol believes it is the first company in the industry to

receive this approval for a CBD related product. To manufacture, import or distribute an active

substance, organizations must comply with good manufacturing and distribution practice, where

products must be of consistent high quality, appropriate to their intended use, and meet the

requirements of the marketing authorisation (MA) or product specification.

Upon successful initial import and product testing, Zenabis and Isodiol intend to execute a

Definitive Supply Agreement, which expects to expand the scope of the collaboration, in

addition to the supply of raw isolate.

About Isodiol International Inc.

Isodiol International Inc. is the market leader in pharmaceutical grade phytochemical

compounds and the industry leader in the manufacturing and development of consumer

products.

Isodiol is the pioneer of many firsts for the cannabis industry including commercialization of

99%+ pure, pharmaceutical grade cannabinoids, micro-encapsulations, and nanotechnology for

the highest quality consumable and topical skin care products and most recently received

approval as having the first CBD designated as an Active Pharmaceutical Ingredient as was

announced April 26, 2018.

Isodiol's growth strategy includes the development of over-the-counter and pharmaceutical

drugs, expanding its phytoceutical portfolio and will aggressively continue international

expansion into Latin America, Asia and Europe.

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ON BEHALF OF THE BOARD

Marcos Agramont, CEO & Director

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About Zenabis Ltd.

Zenabis is a privately-held, cannabis company with one of the largest, federally licensed indoor

medical cultivation footprints in Canada. Zenabis currently operates two licensed production

facilities with more than 400,000 square feet of fully certified growing space in British Columbia

and New Brunswick. Zenabis' s operations are strategically positioned on Canada's coasts

facilitating national distribution and access to international markets. Zenabis is currently working

towards globally recognized GMP certifications.

Zenabis has one of the most experienced management teams in the industry, with an extensive

background in retail consumer packaged goods, global pharmaceutical sales, and

commercialized cultivation. Our growing team has more than two decades of experience in

organic cultivation and distribution of herbs and nutraceutical products throughout the Americas,

North Africa, and the Middle East. Our sales team has more than two decades in product

development, commercialization, and retail and pharmaceutical sales including international

distribution.

For more information contact Gurdeep Bains at ir@zenabis.com

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