

UK launch of VivaGel® BV

- VivaGel® BV has been launched in the UK by Mundipharma
- VivaGel® BV is available in Europe (including the UK) Over-The-Counter
- Mundipharma has launched VivaGel® BV in multiple countries in Europe, with additional European countries expected soon
- Launches are also planned in other Mundipharma regions, including Asia, and regulatory activities continue in multiple regions

Melbourne, Australia; 13 November 2019: Starpharma (ASX: SPL, OTCQX: SPHRY) today announced that VivaGel® BV has been launched in the UK under the brand Betafem® BV Gel. This launch follows the first European launches in June 2019, including in Germany and other countries.

To be sold under the Betadine[®] umbrella brand, VivaGel[®] BV is available Over-The-Counter (OTC) in Europe, without the need to see a doctor or obtain a prescription. Europe represents a large commercial opportunity for VivaGel[®] BV with access to more than 260 million women and Mundipharma continues to roll-out the product in the region.

Dr Jackie Fairley, Starpharma CEO, commented: "We are delighted to see VivaGel® BV launched in the UK. Starpharma and Mundipharma are working closely together to bring this breakthrough product to European women suffering from BV. We look forward to working with Mundipharma on further registrations and launches across the globe".

Raman Singh, Mundipharma CEO, commented: "VivaGel® BV is an important addition to the Betadine® product portfolio, which continues to evolve to support women's everyday health and well-being around the world. Together with Starpharma we look forward to realising our joint ambition for Betafem® and women's health."

Starpharma's and Mundipharma's teams continue to work proactively together on regulatory and roll-out activities of VivaGel® BV across Mundipharma's regions.



About VivaGel® BV

VivaGel® BV is a novel, non-antibiotic therapy for the treatment of bacterial vaginosis (BV) and prevention of recurrent BV. BV is the most common vaginal condition worldwide and twice as common as thrush. One in three women will experience BV and half of these women will have recurrent BV. VivaGel® BV is an Australian innovation – invented, fully developed and taken through to commercialisation by Starpharma.

BV is a troublesome and often recurrent condition that causes unpleasant vaginal odour and discharge symptoms that have significant social impact for women. BV has also been associated with a range of other serious reproductive health-related medical problems.



About Starpharma

Starpharma Holdings Limited (ASX: SPL, OTCQX:SPHRY), located in Melbourne Australia, is an ASX 300 company and is a world leader in the development of dendrimer products for pharmaceutical, life science and other applications.

Starpharma's underlying technology is built around dendrimers – a type of synthetic nanoscale polymer that is highly regular in size and structure and well suited to pharmaceutical and medical uses. Starpharma has two core development programs: VivaGel® portfolio and DEP® drug delivery with the Company developing several products internally and others via commercial partnerships.

VivaGel®: Starpharma's women's health product - VivaGel® BV is based on SPL7013, astodrimer sodium, a proprietary dendrimer. VivaGel® BV for bacterial vaginosis (BV), is available for sale under the brand name Betadine BV™ (Europe) and Fleurstat BVgel (Australia) and a new drug application has been submitted to the US FDA. Starpharma has licensed the sales and marketing of VivaGel® BV to ITF Pharma for the US; Mundipharma for Europe, Russia, CIS, Asia, the Middle East, Africa and Latin America; and to Aspen Pharmacare for Australia and New Zealand. Starpharma also has licence agreements to market the VivaGel® condom (an antiviral condom which includes VivaGel® in the lubricant) in several regions, including Australia, Europe, Canada, China and Japan (Okamoto). The VivaGel® condom has been launched in Japan under Okamoto's 003 brand, and in Australia and Canada under the Lifestyles® Dual Protect™ brand.

DEP® - Dendrimer Enhanced Product®: Starpharma's DEP® drug delivery platform has demonstrated reproducible preclinical benefits across multiple internal and partnered DEP® programs, including improved efficacy, safety and survival. Starpharma has three internal DEP® products – DEP® docetaxel, DEP® cabazitaxel and DEP® irinotecan - in clinical development in patients with solid tumours. Starpharma's partnered DEP® programs include a multiproduct DEP® licence with AstraZeneca, which involves the development and commercialisation of two novel oncology compounds, with potential to add more. In June 2019 Starpharma signed a Development and Option agreement with AstraZeneca for a DEP® version of one of AstraZeneca's major marketed oncology medicines.

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About Mundipharma

Mundipharma's independent associated companies are privately owned entities covering the world's pharmaceutical markets. Mundipharma is a prime example of a company that consistently delivers high quality products while standing by the values that represent the company. Our mission is to alleviate the suffering of patients with cancer and non-cancer pain and to substantially improve their quality of life. Mundipharma is dedicated to bringing to patients with severe and debilitating diseases the benefit of novel treatment options in fields such as pain, oncology, oncology supportive care, ophthalmology, respiratory disease and consumer healthcare. For more information please visit: www.mundipharma.com.sg.

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- ®: VivaGel and DEP are Registered Trademarks of Starpharma.

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Forward Looking Statements

This document contains certain forward-looking statements, relating to Starpharma's business, which can be identified by the use of forward-looking terminology such as "promising", "plans", "anticipated", "will", "project", "believe", "forecast", "expected", "estimated", "targeting", "aiming", "set to", "potential", "seeking to", "goal", "could provide", "intends", "is being developed", "could be", "on track", or similar expressions, or by express or implied discussions regarding potential filings or marketing approvals, or potential future sales of product candidates. Such forward-looking statements involve known and unknown risks, uncertainties and other factors that may cause actual results to be materially different from any future results, performance or achievements expressed or implied by such statements. There can be no assurance that any existing or future regulatory filings will satisfy the FDA's and other authorities' requirements regarding any one or more product candidates nor can there be any assurance that such product candidates will be approved by any authorities for sale in any market or that they will reach any particular level of sales. In particular, management's expectations regarding the approval and commercialization of the product candidates could be affected by, among other things, unexpected trial results, including additional analysis of existing data, and new data; unexpected regulatory actions or delays, or government regulation generally; our ability to obtain or maintain patent or other proprietary intellectual property protection; competition in general; government, industry, and general public pricing pressures; and additional factors that involve significant risks and uncertainties about our products, product candidates, may vary materially from those described herein as anticipated, believed, estimated or expected. Starpharma is providing this information as of the date of this document and does not assume any obligation to update any forward-looking statements contained in t