

US patent granted for SPL7013 eye drops for conjunctivitis

Melbourne, Australia; 12 December 2018: Starpharma (ASX: SPL, OTCQX: SPHRY) today announced that the US Patent and Trademark Office has granted a patent for SPL7013 ophthalmic drops for viral conjunctivitis. The patent has been granted with broad claims for treating and preventing microbial infections of the eye, including adenoviral conjunctivitis, bacterial conjunctivitis and other eye infections. The patent term is to 2033.

Viral conjunctivitis is the most common cause of infectious conjunctivitis, affecting ~6 million people in the US and ~4 million in Europe each year. The viral conjunctivitis market is estimated to be around US\$700M¹ annually. There are currently no approved therapies for viral conjunctivitis and it remains an area of significant unmet medical need globally.

SPL7013 ophthalmic drops have demonstrated compelling efficacy in animal models of viral conjunctivitis. The drops have shown potent anti-viral activity, decreased infectivity and importantly, they are non-irritating.

Starpharma also recently completed formal independent market research in the US with 18 ophthalmologists, payers and primary care physicians who expressed a high level of interest in Starpharma's SPL7013 ophthalmic drops. Clinicians interviewed confirmed that the product would address a major unmet need for a viral conjunctivitis therapeutic and described its novel mechanism of action as "compelling". These clinicians were also impressed by the product's ability to inhibit the spread of this highly contagious disease.

Dr Jackie Fairley, CEO of Starpharma, commented "We are very pleased with the independent market research findings, with positive responses in 87% of clinicians surveyed. This clinical feedback confirms a high level of interest in Starpharma's novel therapeutic for adenoviral conjunctivitis."

This recent progress has generated partnering interest for the product and Starpharma is currently evaluating the optimal development strategy for this novel therapy. SPL7013 is a proprietary dendrimer which is the active ingredient in Starpharma's VivaGel[®] products. Given the extensive preclinical and clinical data already available on SPL7013, there will be significant development advantages both in terms of cost and time.

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, astodimer sodium, a proprietary dendrimer. VivaGel[®] BV is approved for marketing in the EU and Australia for bacterial vaginosis (BV) and a new drug application is under Fast Track review by the US FDA. Starpharma has licensed the sales and marketing of VivaGel[®] BV to 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 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 two internal DEP[®] products – DEP[®] docetaxel and DEP[®] cabazitaxel - in clinical development in patients with solid tumours, and further DEP[®] products approaching clinical

¹ Roth Capital Partners



development. 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.

Starpharma.com | [Twitter](#) | [LinkedIn](#)

Media

WE Buchan Consulting

Rebecca Wilson

Mob: +61 417 382 391

rwilson@buchanwe.com.au

Arthur Chan

+61 2 9237 2805

achan@buchanwe.com.au

Starpharma

Dr Jackie Fairley, Chief Executive Officer

Nigel Baade, CFO and Company Secretary

+61 3 8532 2704

investor.relations@starpharma.com

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, financial results and business prospects. Should one or more of these risks or uncertainties materialize, or should underlying assumptions prove incorrect, actual results 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 this document as a result of new information, future events or developments or otherwise.