



ASX ANNOUNCEMENT

Quarterly Cashflow Report

Melbourne Australia; 28 October 2013 - Starpharma Holdings Ltd (ASX: SPL; OTCQX: SPHRY) today released its Appendix 4C – Quarterly Cashflow report for the period ended 30 September 2013.

The cash balance at 30 September 2013 was \$31.5 million, a net cash burn of \$2.3 million for the quarter.

During the quarter Starpharma reported positive results of pre-clinical studies which examined the Company's dendrimer-enhanced version of the blockbuster cancer drug, oxaliplatin (ELOXATIN[®]) in a colon cancer model (xenograft). The results demonstrated that compared with the original version, ELOXATIN[®], Starpharma's Dendrimer-Enhanced Oxaliplatin was more effective in inhibiting tumour growth, and substantially reduced the serious bone marrow toxicities that are reported in a high proportion of patients receiving oxaliplatin.

Further positive results for Starpharma's Dendrimer-Enhanced Oxaliplatin nanoparticles were announced on 14 October, when Starpharma reported the results of a trial that showed the dendrimer formulation resulted in a significant reduction in neurotoxicity, the major (and dose-limiting) toxicity of oxaliplatin.

More recently Starpharma also reported that its lead oncology candidate, dendrimer docetaxel which will enter the clinic shortly, did not cause bone marrow toxicity or neutropenia, the dose limiting side effect commonly seen with the original formulation of docetaxel. This formulation has already been shown in preclinical studies to have several other advantages compared to Taxotere[®] including improved efficacy.

Planning is also well advanced for the Phase 3 trials of VivaGel[®] for the prevention of recurrence of bacterial vaginosis, while the VivaGel[®]-coated condom – licensed to Ansell and Okamoto – is currently under regulatory review ahead of market launch.

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 uses. Starpharma has three core development programs: VivaGel[®]

portfolio, drug delivery, and agrochemicals with the Company developing a number of products internally and others via commercial partnerships.

Starpharma's lead product is VivaGel® (SPL7013 Gel), a gel-based formulation of a proprietary dendrimer. VivaGel® is under clinical development for the treatment and prevention of bacterial vaginosis (BV) and also as a vaginal microbicide to prevent the transmission of sexually transmitted infections including HIV and genital herpes. Starpharma has also signed separate licence agreements with Ansell Limited (ASX:ANN) and Okamoto Industries Inc (Tokyo Stock Exchange) to market a value-added, VivaGel®-coated condom. Ansell manufactures and sells leading condom brands worldwide, including Lifestyles®, ZERO® and SKYN®. Okamoto is the market leader for condoms sold in Japan, the world's second largest condom market.

In the wider pharmaceutical and life science fields, Starpharma has both partnered and internal programs in Drug Delivery. Drug Delivery partners include GSK, Lilly and AstraZeneca. In its internal program Starpharma has announced significant tumour-targeting results in its docetaxel (Taxotere®) program, with animal studies showing its dendrimer-enhanced version of docetaxel to have significantly superior anti-cancer effects across a range of important cancer types including breast, prostate, lung and ovarian tumour, when compared to Taxotere® (docetaxel).

In agrochemicals Starpharma has a series of partnerships with leading industry players including Nufarm (ASX:NUF) and Makhteshim Agan as well as internal programs including an enhanced version of glyphosate (the active ingredient in Roundup®).

FOR FURTHER INFORMATION

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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 health authorities' requirements regarding any one or more product candidates nor can there be any assurance that such product candidates will be approved by any health 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 clinical trial results, including additional analysis of existing clinical data, and new clinical 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.

Name of entity

Starpharma Holdings Limited

ABN

20 078 532 180

Quarter ended ("current quarter")

30 September 2013

Consolidated statement of cash flows**Cash flows related to operating activities**

	Current Quarter \$A'000	Year to Date \$A'000
1.1 Receipts from customers and grants (including R&D Tax Incentive)	172	172
1.2 Payments for		
(a) staff costs	(1,157)	(1,157)
(b) advertising and marketing	-	-
(c) research and development	(1,836)	(1,836)
(d) other working capital	-	-
1.3 Dividends received	-	-
1.4 Interest and other items of a similar nature received	370	370
1.5 Interest and other costs of finance paid	(2)	(2)
1.6 Income taxes paid	-	-
1.7 Other	-	-
Net operating cash flows	(2,453)	(2,453)

Cash flows related to investing activities

1.9 Payment for acquisition of:		
(a) businesses (item 5)	-	-
(b) equity investments	-	-
(c) intellectual property	-	-
(d) physical non-current assets	(11)	(11)
(e) other non-current assets	-	-
1.10 Proceeds from disposal of:		
(a) businesses (item 5)	-	-
(b) equity investments	-	-
(c) intellectual property	-	-
(d) physical non-current assets	-	-
(e) other non-current assets	-	-
1.11 Loans to other entities	-	-
1.12 Loans repaid by other entities	-	-
1.13 Other	-	-
Net investing cash flows	(11)	(11)
1.14 Total operating and investing cash flows	(2,464)	(2,464)

Cash flows related to financing activities

1.15 Proceeds from issues of shares (net)	56	56
1.16 Proceeds from sale of forfeited shares	-	-
1.17 Proceeds from borrowings	-	-
1.18 Repayment of borrowings	-	-
1.19 Dividends paid	-	-
1.20 Other - lease repayments	(8)	(8)
Net financing cash flows	48	48
Net increase (decrease) in cash held	(2,416)	(2,416)
1.21 Cash at beginning of quarter/year to date	33,840	33,840
1.22 Exchange rate adjustments	53	53
1.23 Cash at end of quarter	31,477	31,477

Payments to directors of the entity and associates of the directors
Payments to related entities of the entity and associates of the related entities

	Current quarter \$A'000
1.24 Aggregate amount of payments to the parties included in item 1.2	(196)
1.25 Aggregate amount of loans to the parties included in item 1.11	-

1.26 Explanation necessary for an understanding of the transactions

Item 1.24 consists of the following:

(a) Remuneration paid to the Chief Executive Officer.

(b) Director's fees paid to non-executive directors.

Non-cash financing and investing activities

2.1 Details of financing and investing transactions which have had a material effect on consolidated assets and liabilities but did not involve cash flows

2.2 Details of outlays made by other entities to establish or increase their share in businesses in which the reporting entity has an interest

Nil

Financing facilities available

Add notes as necessary for an understanding of the position. (See AASB 1026 paragraph 12.2).

	Amount available \$A'000	Amount used \$A'000
3.1 Loan facilities - Finance facility for laboratory equipment	200	100
3.2 Credit standby arrangements - Credit card facility	160	24

Item 3.1 A \$200,000 master asset finance facility with National Australia Bank for laboratory equipment, guaranteed by term deposit.

Reconciliation of cash

Reconciliation of cash at the end of the quarter (as shown in the consolidated statement of cash flows) to the related items in the accounts is as follows.		Current quarter \$A'000	Previous quarter \$A'000
4.1	Cash on hand and at bank	1,665	1,503
4.2	Deposits at call	29,812	32,337
4.3	Bank overdraft	-	-
4.4	Other (provide details)	-	-
Total: cash at end of quarter (item 1.23)		31,477	33,840

Acquisitions and disposals of business entities

- 5.1 Name of entity
- 5.2 Place of incorporation or registration
- 5.3 Consideration for acquisition or disposal
- 5.4 Total net assets
- 5.5 Nature of business

Acquisitions (Item 1.9(a))	Disposals (Item 1.10(a))
-	-
-	-
-	-
-	-
-	-

Compliance statement

1. This statement has been prepared under accounting policies which comply with accounting standards as defined in the Corporations Law (except to the extent that information is not required because of note 2) or other standards acceptable to ASX.
2. This statement does give a true and fair view of the matters disclosed.



28 October 2013

B P Rogers
Company Secretary