

Quarterly Cashflow Report

Melbourne, Australia; 30 July 2012: Starpharma Holdings Ltd (ASX:SPL;OTCQX: SPHRY) today released its Appendix 4C – Quarterly Cashflow report for the period ended 30 June 2012.

The cash balance at 30 June 2012 was \$42.8 million. Total operating and investing cash outflow for the quarter was \$3.8 million, resulting in an annual cash burn of \$9.9 million. The increase in cash burn this year is in line with budgets and is driven primarily by the concurrent Phase 2 and two Phase 3 clinical trials underway for the VivaGel® Bacterial Vaginosis (BV) Program.

Recent progress in these programs includes:

- Full recruitment of the first of Starpharma's two Phase 3 studies for VivaGel® as a treatment for BV. The second Phase 3 trial is now more than 70% recruited.
 - This study is the subject of a formal agreement with FDA under the Special Protocol Assessment (SPA) program, which confirms that the trial design, clinical endpoints and statistical analyses are acceptable for FDA approval once complete.
- Full recruitment for the Phase 2 clinical trial to investigate the ability of VivaGel[®] to prevent the recurrence of BV. The study is being conducted under an investigational new drug application (IND) and has enrolled 205 patients with a prior history of recurrent BV.

The two Phase 3 treatment trials and the Phase 2 trial for prevention of recurrence are all expected to complete in 2H CY2012. Following successful completion Starpharma will file an NDA and enter into one or more licensing agreements.

In addition to the active clinical trial program for VivaGel[®] referred to above, Starpharma has two commercial collaborations for a VivaGel[®] coated condom, as well as an internal drug delivery program with a focus on improvement of the blockbuster cancer drug docetaxel (Taxotere[®]). The drug delivery program is advancing rapidly with plans underway for clinical trials to commence in 2013.

Starpharma's dendrimers are also being applied to enhancing the performance of important leading agrochemical agents such as Glyphosate (Roundup®).

ABOUT STARPHARMA

Starpharma Holdings Limited (ASX:SPL, OTCQX:SPHRY) 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. In addition, products for diagnostics and laboratory reagents are already on market through licence arrangements with partners including Siemens Healthcare and Merck KGaA.

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. Most recently Starpharma announced pre-clinical results in its Docetaxel (Taxotere®) program demonstrating significant improvements in that agent's anticancer efficacy and the enhancement of solubility offering potential safety benefits as well. The company is also exploring dendrimer opportunities in agrochemicals in a series of industry partnerships as well as with internal programs including an enhanced version of glyphosate (the active ingredient in Roundup®).

Starpharma's headquarters and research facilities are located in Melbourne, Australia

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.

Starpharma Holdings Limited

ABN	Quarter ended ("current quarter")	
20 078 532 180	30 June 2012	

Consolidated statement of cash flows

Cash flows related to operating activities			rrent r \$A'000	Year to Date \$A'000	
1.1	Receipts from cus	stomers		152	1,546
1.2	Payments for	(a) staff costs		(1,486)	(4,598)
		(b) advertising and marketing		-	-
		(c) research and development		(3,247)	(8,318)
		(d) leased assets		(22)	(80)
		(e) other working capital		-	-
1.3	Dividends receive			-	-
1.4		items of a similar nature received		879	1,608
1.5		costs of finance paid		(3)	(8)
1.6	Income taxes paid	d		-	-
1.7	Other			-	-
	Net operating cash	flows		(3,727)	(9,850)
	ws related to investin				
1.9	Payment for acqu				
	(a) businesses (it			-	-
	(b) equity investm			-	-
	(c) intellectual pro			-	(122)
	(d) physical non-			(96)	(133)
1.10	(e) other non-cur			-	-
1.10	Proceeds from dis (a) businesses (it	•		-	-
	(b) equity investr	·		-	-
	(c) intellectual pro			_	
	(d) physical non-			_	_
	(e) other non-cur				_
	(c) other non our	Terri deserte		_	_
1.11	Loans to other en	tities		_	-
1.12	Loans repaid by o	ther entities		_	-
1.13	Other			-	-
	Net investing cash	flows		(96)	(133)
1.14	Total operating an	nd investing cash flows		(3,823)	(9,983)
		-		, ,	` ' '
	ws related to financia		Γ		22 545
1.15 1.16		sues of shares (net) le of forfeited shares		-	33,745
1.16	Proceeds from bo			-	-
1.17	Repayment of bor	•		-	-
1.10	Dividends paid	Townigs		-	-
1.19	Other				_ [
1.20	Net financing cash	flows			33,745
	Tite imaneing cush	. HOW'S			33,743
Net incre	ease (decrease) in cas	sh held		(3,823)	23,762
1.21	Cash at beginning	of quarter/year to date		46,607	18,918
1.22	Exchange rate ad			28	132
1.23	Cash at end of qua			42,812	42,812

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

		\$A'000	
1.24	Aggregate amount of payments to the parties included in item 1.2	(180)	
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 corassets and liabilities but did not involve cash flows	nsolidated	
2.2	Details of outlays made by other entities to establish or increase their share in busines which the reporting entity has an interest	sses in	
	Nil		
_	facilities available 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	297	140
3.2	Credit standby arrangements - Credit card facility	160	17

Reconciliation of cash

Item 3.1

	liation of cash at the end of the quarter (as shown in the consolidated statement of ws) 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	1,455	1,720
4.2	Deposits at call	41,357	44,887
4.3	Bank overdraft	-	-
4.4	Other (provide details)	-	-
	Total: cash at end of quarter (item 1.23)	42,812	46,607

A \$97,000 lease facility and a \$200,000 master asset finance facility with National Australia Bank for laboratory equipment, guaranteed by term deposit.

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.

30 July 2012

B P Rogers Company Secretary