

Interim report January – September 2014

Aerocrine Achieves Record Sales for the 3rd Quarter and First Nine Months: The Board of Directors has resolved to raise further capital to support the Company's continued growth through an upcoming financing in the amount of approximately SEK 443m of which more than 75% has been guaranteed.

January – September 2014

- Net sales increased by 15% to SEK 116.0m (100.5)*. Adjusted for exchange rates, net sales increased by 10%.
 - Net sales for clinical use of NIOX products were SEK 92.8m, an increase of 21%.
 - Strategic sales (sales to pharmaceutical companies and CROs for clinical trials) were SEK 19.5m, an increase of 3%.
- Total tests sold (repeat and initial) were 1.8m (1.5) tests, an increase of 22%. Total repeat test volume increased by 26%.
- The Gross Margin for the period was 69% (73%). The reduction vs prior year was driven primarily by a change in channel mix.
- The loss after tax was SEK 167.7m (166.5), corresponding to a loss per share before and after dilution of SEK 1.1 (1.1).
- The operating loss improved to SEK 130.1m (154.1).
- Cash flow from operations was negative in the amount of SEK 154.7m (-169.0).

July – September 2014

- Net sales increased by 24% to SEK 38.5m (31.1)*. Adjusted for exchange rates, net sales increased by 16%.
 - Net sales for clinical use of NIOX products were SEK 30.4m (22.6), an increase of 34%.
 - Strategic sales were SEK 6.9m (7.1), a decrease of 2%.
- Total tests sold (repeat and initial) were 0.6m (0.5) tests, an increase of 33%. Total repeat test volume increased by 33%.
- Gross Margin was 68% (74%).
- The loss after tax was SEK 60.2m (56.5), corresponding to a loss per share before and after dilution of SEK 0.4 (0.4).
- The operating loss improved to SEK 41.8m (47.9).
- Cash flow from operations was negative in the amount of SEK 49.5m (-50.8).

Significant Events July – September 2014

- On September 29, 2014, the Company announced a collaboration with Microsoft. The collaboration will utilize Microsoft's secure cloud services to transmit device telemetry data from physician and Company sites in Sweden, UK and the US back to Aerocrine for analyses. Aerocrine's goal is to use this information to deploy its field resources for customer service and sales support. Microsoft chose Aerocrine due to its technology leadership, global reach and commitment to progressing human health management.

Significant Events, After the Period

- On November 6, 2014, the United States Food and Drug Administration (FDA) granted market clearance for NIOX VERO, the Company's next generation device for measuring Fractional exhaled Nitric Oxide (FeNO) levels in the clinical setting. NIOX VERO will add significant functionalities such as new User Interface, extended capacity and mobility (battery operated) which has been very well received in Europe during the launch 2014.
- On November 8, 2014, A position paper from the French Speaking Respiratory Society was made available in "Revue des Maladies Respiratoires (2014)". The French guidelines for FeNO are an important step towards submitting for reimbursement in France. The guidelines are summarized as "Measuring FeNO is the only noninvasive pulmonary function test allowing (1) detecting, (2) quantifying and (3) monitoring changes in inflammatory processes during the course of various respiratory disorders, including corticosteroid sensitive asthma".
- On November 26, 2014, Thomas Eklund resigned from the Board of Directors.
- On November 26, 2014, the Board of Directors resolved to secure financing of approximately SEK 443m, before transaction costs. The financing will be structured as a rights offering, which will be

subject to approval by the shareholders at an EGM to be held on January 7, 2015. Details of the rights offering will be communicated via a press release on November 28, 2014. In excess of 75% of the financing has been guaranteed by inter alia Aerocrine's largest shareholder Novo A/S and the largest Danish public pension fund, Arbejdsmarkedets Tillægspension (ATP).

Summary of Financial Information

(SEK m)	Aerocrine Group					
	Jul 1, 2014 - Sep 30, 2014	Jul 1, 2013 - Sep 30, 2013	Jan 1, 2014 - Sep 30, 2014	Jan 1, 2013 - Sep 30, 2013	Rolling 12 Months	Jan 1, 2013 - Dec 31, 2013
Net Sales	38,5	31,1	116,0	100,5	151,7	136,2
Strategic Sales	6,9	7,1	19,5	18,9	27,9	27,3
Gross Profit/Loss	26,2	22,9	79,6	73,1	104,4	97,8
Gross Margin %	68%	74%	69%	73%	69%	72%
Operating Loss	-41,8	-47,9	-130,1	-154,1	-181,8	-205,7
Net Loss After Tax	-60,2	-56,5	-167,7	-166,5	-226,9	-225,6
Cash Flow, Current Operations	-49,5	-50,8	-154,7	-169,0	-197,8	-212,1
Total Cash Flow	-50,5	-62,2	-157,7	132,7	-200,7	89,7

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*Note all numbers in () are the corresponding period previous year and in the same unit.

This is information that Aerocrine AB (publ) is required to publish in accordance with the Swedish Securities Markets Act and/or the Swedish Financial Trading Act. This information was submitted for publication on November 27, 2014, at 8.00 a.m.

CEO's Comments on Q3 2014

"We have overcome significant obstacles in the first nine months of the year and our understanding of our markets and our customers continues to evolve. Similarly, we have increased our focus on growth as a key imperative and worked to make sure that we do this in a capital efficient manner. We have accomplished both of these through Q3 of this year and for that I am very proud of our organization.

In terms of topline results, Q3 2014 represents the best third quarter in the Company's history. Additionally, with strong consecutive quarters in Q2 and Q3, the YTD results represent the best first three quarters ever. We mentioned in previous reports that our US business would begin showing signs of improvement in the second half of 2014, and Q3 represented our strongest clinical sales performance in the US ever. The total US clinical sales were SEK 11.7m in Q3, an increase of 2% compared to Q3 2013 and 23% compared to Q2 2014.

We attribute the improved performance in the US to the changes we have made to the selling model, including improved targeting data and messaging, the device evaluation program, and offering customers multiple pricing options. The overall reimbursement landscape improved with better coverage in key states such as Illinois and Texas and we expanded our sales force there to take advantage of better reimbursement. We now have 28 sales territories and plan to keep that structure into 2015. Our US team demonstrated increased productivity as well as increased repeat test sales which is a key indicator of the underlying health of our business.

Our Ex-US business continues strong double digit growth with Europe/ROW contributing 15% growth for both Q3 and the first nine months, as compared to the corresponding periods in 2013. Asia Pacific has exhibited exceptional performance with growth of 328% and 152% for Q3 and the first nine months, as compared to the corresponding periods in 2013, with Japan as a key contributor.

We have also focused on using our capital more efficiently. We accomplished this by globalizing certain functions and becoming more efficient in our processes through automation. We have been able to manage headcount down by 13% from this point last year while still supporting a growing business. On a percent of sales basis we continue to leverage our fixed cost base and have driven our operating expenses down by SEK 17m while increasing sales by approximately SEK 16m (Q3 YTD 2014 vs. Q3 YTD 2013) for a resulting decrease of 45 percentage points (226% to 181%). Absent these initiatives our expense base

would have been SEK 52m higher. While we need to drive this number lower to attain our overall goal of profitability the results have been three consecutive quarters of reduced operating losses.

With regard to product and service development, we announced an innovative collaboration with Microsoft whereby we were chosen by their global cloud computing and data analytics group to pioneer the use of their secure cloud network to actively monitor several of our devices on different continents. The pilot project is in full swing and we can now monitor devices in near real time. This will improve our ability to monitor device performance as well as also provide valuable information regarding test utilization and upcoming sensor and device expirations. This will add increased efficiency to our selling efforts and deployment of customer support resources. This project will conclude in December with a summit in Brussels where the collaboration results will be showcased by Microsoft.

On the regulatory front, on November 6th the NIOX VERO was approved by the FDA for US commercialization. The NIOX MINO re-registration in China is on track for approval in Q1 of 2015 and additional NIOX VERO submissions are awaiting approval in Japan and China. Expectations for approval are Q2 2015 in Japan and the second half of 2015 in China, but are all subject to the timelines and processes governed by each country.

As we have renewed our short, mid and long term planning for Aerocrine, we took into account the funding that would be needed to grow our topline and enable continued progress toward long-term profitability and value creation for our shareholders. We believe that we have created a strong foundation for our continued growth and strategic direction that we intend to support through the upcoming financing in the amount of approximately SEK 443m, before transaction costs.

We will focus on 4 key areas:

- 1 | Establish FeNO as Standard of Care
- 2 | Drive Penetration in Defined U.S. Professional Segment
- 3 | Attain Profitability
- 4 | Finalize Home Device Product and Business Model

The Company will focus its time and resources to accomplish these key objectives and continue to make Aerocrine the leader in providing FeNO measurement capability to the global medical community. This is an ambitious plan that will take time and resources and we are committed to building Aerocrine and increasing shareholder value" says Aerocrine's President and CEO, Scott Myers.

For case-studies on how FeNO assists in the daily practice to set correct diagnosis and initiate correct medication please visit: <http://www.niox.com/en/>

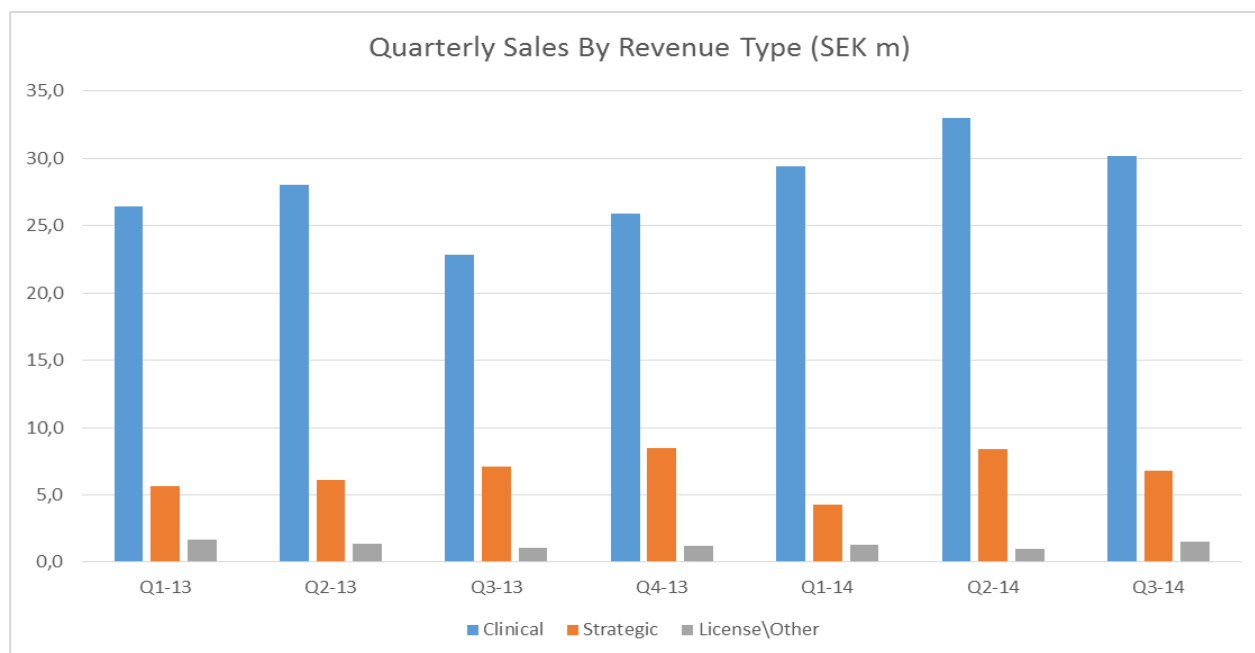
Overview, January – September 2014

Sales

Net sales for the period January – September 2014 reached SEK 116.0m (100.5), an increase of 15%. Adjusting for the change in currency during the year, the increase was 10%. The net sales for clinical use of NIOX products increased 21% to SEK 92.8m (76.5), driven mainly by strong sales in Japan following the market clearance received in the fourth quarter 2013 and solid performance in the EU.

The implementation of a new sales model in the US slowed growth in the first half of 2014, with clinical sales down by 9% in local currency, but sales rebounded in the last half of Q2 and showed solid growth in Q3. Changes were primarily related to additional focus and improved targeting of potential customers based on level of coverage, size of the clinic and number of patients with asthma symptoms. Additional

changes included new pricing options as well as a subscription plan and consignment plan and the introduction of the evaluation program to further stimulate sales, whereby potential customers are able to use the device for a period of time to further access the benefit of FeNO in their practice.

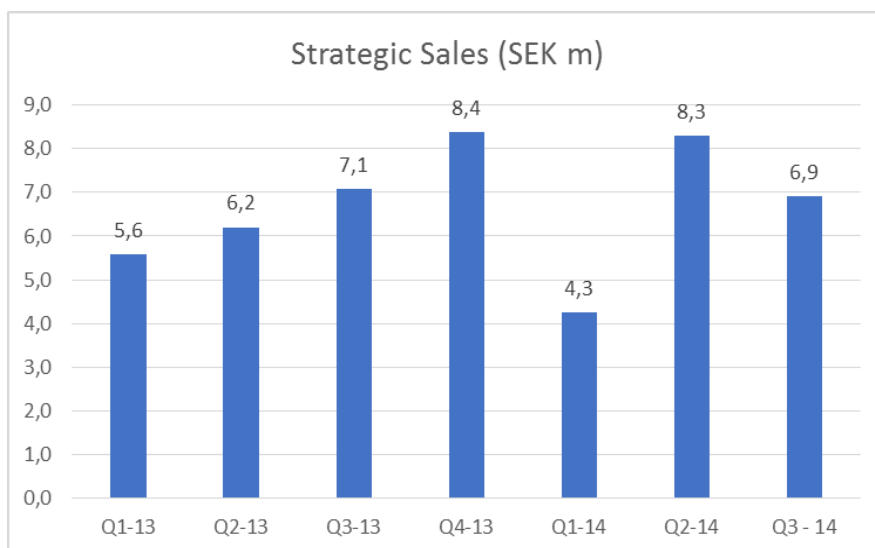


Sales Summary, Region/Other

	Aerocrine Group								
	Jul 1, 2014 - Sep 30, 2014	Jul 1, 2013 - Sep 30, 2013	% Change	Jan 1, 2014 - Sep 30, 2014	Jan 1, 2013 - Sep 30, 2013	% Change	Rolling 12 Months	Jan 1, 2013 - Dec 31, 2013	% Change
(SEK m, except for Tests)									
Clinical Sales									
US/North America	11,7	11,4	2%	29,4	32,2	-9%	39,5	42,3	-7%
EU/RoW	13,1	10,0	32%	44,3	37,1	19%	57,3	50,2	14%
Asia Pacific	5,6	1,2	354%	19,0	7,2	166%	21,8	10,0	119%
Total Clinical Sales	30,4	22,6	34%	92,8	76,5	21%	118,6	102,4	16%
Strategic Sales	6,9	7,1	-2%	19,5	18,9	3%	27,8	27,2	2%
Other Revenue	1,2	1,4	-19%	3,8	5,1	-26%	5,2	6,5	-21%
Total Tests (in 000s)	609	458	33%	1,827	1,498	22%	2,344	2,015	16%
Repeat Test (in 000s)	479	361	33%	1,480	1,175	26%	1,874	1,569	19%

Strategic Sales

Strategic sales for the period, which are included in Region results, increased by 3% compared to the corresponding period in 2013. Strategic sales represented 17% (19%) of total sales during the period. It is important to note that strategic sales fluctuate between quarters as they are impacted by the size and timing of shipment for clinical trials.



Test Volumes

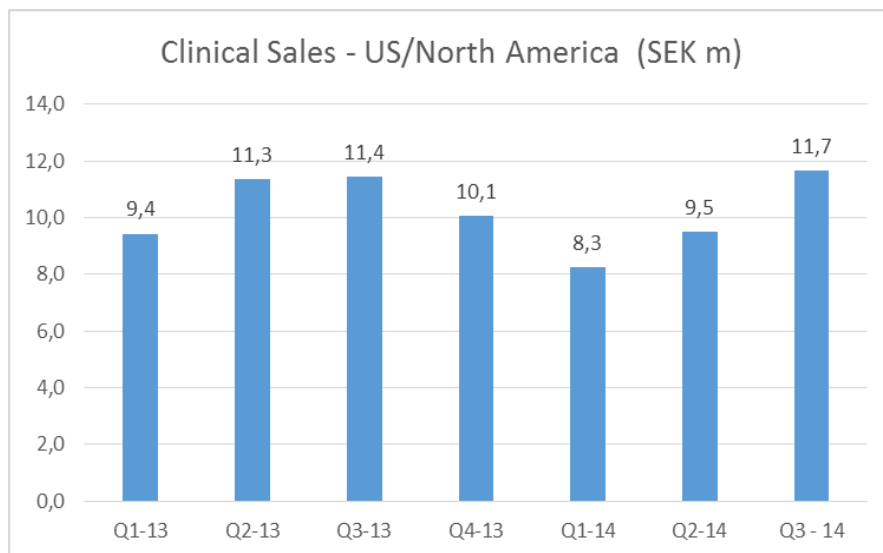
An important metric for the Company is repeat and initial

test sales volumes. A repeat test is defined as the second and subsequent purchases of test kits. During the period a total of approximately 1.5m (1.2) repeat tests were sold, an increase of approximately 26%. This increase is primarily due to an increase in the number of installed devices and also increased usage by physicians. Total tests sold for the period (repeat tests and initial test), were 1.8m (1.5) tests, an increase of approximately 22% compared with the corresponding period in 2013.

Segment Results

US/North America

Sales for the period January – September 2014 in the US/North America segment amounted to SEK 40.3m (43.8) a decrease of 8%. When adjusted for currency effects, sales in the segment decreased by 10%. The sales decrease is primarily due to clinical sales showing a decrease of 9% in local currency and by strategic sales decreasing by 10% in local currency. The changes in the clinical sales model vs. 2013 have resulted in a short-term decrease in sales in the first half of the year. The implementation of the new sales model is now finalized and we are seeing positive results in the third quarter.

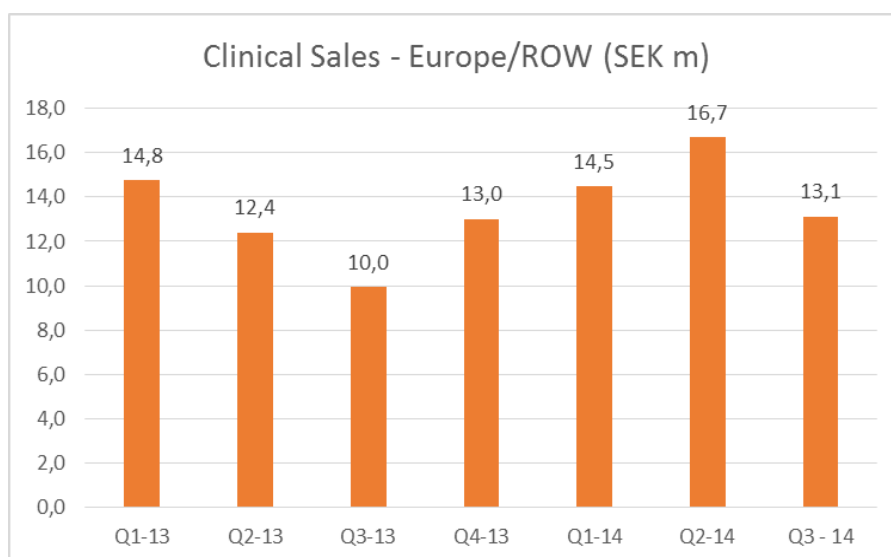


The number of tests sold for clinical use was essentially flat when compared to the same period in 2013 and amounted to approximately 455k (454k) tests sold. The sales of repeat tests for clinical use grew by 26% and the initial test sales decreased by 33% as a consequence of managing through the implementation of the new sales model. In the first nine months of 2014, 666 devices were placed as part of the US evaluation program, of which approximately 145 were still outstanding as of September 30th.

Sales to new and ongoing clinical studies are expected to continue to represent an important part of revenues in the US. Of the sales in the segment SEK 9.3m (10.1) are attributable to strategic sales.

Europe/ROW

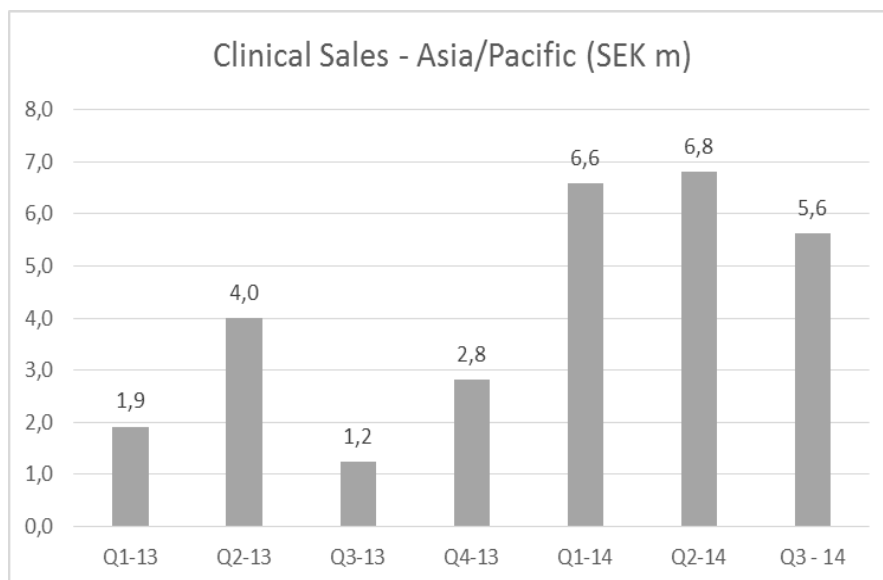
Sales for the segment in the period amounted to SEK 56.3m (49.0), an increase of 15%. When adjusted for currency effects, sales increased by approximately 10%. Clinical sales, excluding strategic sales and license revenues, increased by 19% and reached a record of SEK 44.3m (37.1). Germany and Spain were the main contributors to the clinical sales growth in the segment. During the period the NIOX VERO was introduced in selected markets in the segment. Strategic sales within the segment amounted to SEK 10.2m (8.8).



The sales of repeat tests increased by 8% in the segment compared to the corresponding period 2013 and amounted to 732k (679k) tests sold. The total number of tests sold for clinical use increased by 13% and amounted to 810k (718k).

Asia/Pacific

Sales in the segment amounted to SEK 19.5m (7.7), an increase of 152%. When adjusted for currency effects, sales in the segment increased by 139%. The main reason for the increased sales in the segment is the strong growth in the Japanese market, where the latest version of NIOX MINO received market clearance during the fourth quarter of 2013. A backlog of demand as well as the exchange of the previous version of NIOX MINO has been the main drivers behind the growth.



The sales of repeat tests for clinical use grew by 109% in the segment compared to the corresponding period in 2013 and amounted to 298k (143k) tests sold. The total number of sold tests for clinical use grew by 114% and amounted to approximately 396k (185k) tests.

Profit and Loss

The gross margin for the period was 69% (73%). The gross margin was mainly impacted by changes in the mix of direct/distributor markets.

The loss after tax for the period January – September 2014 amounted to SEK 167.7m (166.5). The loss per share amounted to SEK 1.1 (1.1). Adjusted for certain non-operating and non-cash items detailed in the table below, underlying on-going operations generated a loss of SEK 129.5m (149.0).

(SEK m)	Aerocrine Group					
	Jul 1, 2014 - Sep 30, 2014	Jul 1, 2013 - Sep 30, 2013	Jan 1, 2014 - Sep 30, 2014	Jan 1, 2013 - Sep 30, 2013	Rolling 12 Months	Jan 1, 2013 - Dec 31, 2013
Loss After Tax	-60,2	-56,5	-167,7	-166,5	-226,9	-225,6
Patent Litigation	0,0	-0,1	-0,1	-1,2	-0,3	-1,4
Stock Option Expense	0,2	-2,6	-1,0	-3,8	0,9	-1,9
Revaluation A/R, Liabilities, Cash	8,0	-9,6	15,2	-2,4	19,6	2,0
Effective Interest on Loan	-9,4	-8,7	-26,8	-14,6	-35,6	-23,4
Loan Revaluation	-17,4	9,9	-25,5	4,5	-28,2	1,8
Adjusted Loss	-41,6	-45,4	-129,5	-149,0	-183,3	-202,7

Adjusted earnings improved mainly due to increased sales and reduced expenses related to sales and marketing outside US and reduced administration and development expenses. Of the sales and marketing expenses SEK -0.2m (1.9) constitutes non-cash expenses related to the Group's personnel stock option program. The decrease in sales and marketing expenses outside the US is mainly attributable to a reduction in headcount as the UK and Swedish markets migrated to distributor model. Development costs have decreased primarily due to the decrease in litigation expenses and planned savings. Of the development expenses SEK -1.8m (+0.4) constitutes non-cash expenses related to the Group's personnel stock option program. The decrease in administration costs are primarily due to one-time expenses related to the financing in 2013. Of the administration expenses SEK +0.2m (2.2) are attributable to the Group's personnel stock option program.

The currency effect on the Group's consolidated sales was positive to the amount of SEK 5.3m, while the

effect on the Group's costs and purchasing was negative to the amount of SEK 5.9m. The total effect of exchange rates was slightly negative on the Group's net operating result compared to the corresponding period of 2013.

On 31 December 2013, the Group's consolidated tax loss was calculated at SEK 1,544.4m (1,315.3), of which SEK 1,488.7m (1,256.9) was attributable to the Parent Company. Of the total tax loss SEK 1,495.8m (1,264.6) is unlimited in terms of the period in which it can be offset against future taxable profits. The tax value of the tax-loss carry-forwards has not been capitalized.

Overview, July – September 2014

Sales

Net sales for the third quarter 2014 were SEK 38.5m (31.1), an increase of 24%. When adjusted for currency effects, net sales amounted to SEK 36.1m, an increase of 16%. The net sales for clinical use of NIOX products increased in the period by 34% and reached a record, for a third quarter, of SEK 30.4m (22.6) mainly due strong sales in Japan. Clinical sales in the US market increased in the third quarter by 2% primarily due to a stronger volume of repeat tests.

Test Volumes

During the period a total of approximately 0.5m (0.4) repeat tests were sold, an increase of 33%. Total tests sold for the period (repeat tests and initial test), reached nearly 0.6m (0.5) tests, an increase of 33% compared with the corresponding period 2013. Total and repeat tests sold as part of strategic sales represented 12% and 14% respectively, of total tests sold in the current year period. This compares to total and repeat tests sold as part of strategic sales of 13% and 14% respectively, of total tests sold in the prior year period.

Strategic Sales

Strategic sales, which are included in Region results, were SEK 6.9m (7.1), a decrease of 2%. Strategic sales represented 18% (23) of the total sales during the period. It is important to note that strategic sales fluctuate between quarters as they are impacted by the size and timing of shipment for clinical trials

Segment Results

US/North America

Sales for the third quarter 2014 in the US/North America segment amounted to SEK 16.0m (15.3) an increase of 5%. When adjusted for currency effects, sales in the segment showed no change. Clinical sales increased by 2% (less than 1% in local currency), driven primarily by increased repeat sales. Strategic sales experienced a slight decrease to SEK 3.4m (3.7).

The total number of tests sold for clinical use increased by 14% compared to the third quarter of 2013 and amounted to approximately 196k (173). Repeat tests sold for clinical use increased by 37% compared to the third quarter of 2013 and amounted to approximately 129K (94).

Europe/ROW

Sales in the third quarter amounted to SEK 16.8m (14.6), an increase of 15%. When adjusted for currency effects, sales increased by approximately 8%. The sales increase is mainly attributable to strong sales across EU with Germany and Spain being the main contributors. Strategic sales amounted to SEK 3.5m (3.7). Clinical sales increased by 22%.

The total number of tests sold for clinical use increased by 16% compared to the third quarter of 2013 and amounted to approximately 227k (196). Repeat tests sold for clinical use increased by 3% compared to the third quarter of 2013 and amounted to approximately 200k (195).

Asia/Pacific

Sales in the third quarter amounted to SEK 5.7m (1.3), an increase of 328%. When adjusted for currency effects, sales in the segment increased by approximately 300%. The main reason for the increased sales was the market clearance of NIOX MINO in Japan in Q4-2013.

The total number of tests sold for clinical use increased by 255% compared to the third quarter of 2013 and

amounted to approximately 111k (31). Repeat tests sold for clinical use increased by 250% compared to the third quarter of 2013 and amounted to approximately 83k (24).

Profit and Loss

The gross margin for the period was 68% (74%). The lower margin is mainly attributable to changes in the mix of direct/distributor markets.

The loss after tax for the period July - September 2014 amounted to SEK 60.2m (56.5) and the operating loss amounted to SEK 41.8m (47.9). The loss per share amounted to SEK 0.4m (0.4). The on-going operations generated a loss of SEK 41.6m (45.4) when adjusted for the non-operating/non-cash items detailed previously. Adjusted earnings improved mainly from increased sales and reduced operating expenses.

Market Development

US/North America

During 2014, the fifth largest insurance company Health Care Service Corporation decided to implement a positive coverage policy regarding FeNO testing for asthma diagnosis and management effective from April 1, 2014.

Of approximately 284.4m total covered lives in the US, approximately 65% or 184.1m are reimbursable with regards to Aerocrine products, as of September 30, 2014. Of approximately 178.7m private payor covered lives in the US, approximately 55% or 93.3m are reimbursable lives with regards to Aerocrine products as of September 30, 2014.

Europe/ROW

In the UK, NICE (National Institute of Health and Care Excellence) are due to publish new Asthma Management & Diagnosis Guidance in mid-2015 which we expect to further support the FeNO Guidance published by NICE in April 2014.

Asia/Pacific

The approval of the second generation of the NIOX MINO in Japan is an important step in the continued development of the Japanese market. Currently there is a reimbursement code for exhaled gas in-place of approximately SEK 70 per test, which also covers FeNO. The process of applying for increased reimbursement has been initiated by a number of Japanese specialist societies (Japanese Respiratory Society among others).

Significant Events, During the Period

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Clinical Data

In August 2014, the *Journal of Allergy and Clinical Immunology* published a study that examined the cost and clinical effectiveness of 3 different strategies to control asthma. Two of the strategies were based on the ACQ (Asthma Control Questionnaire) to guide control of asthma to either partially or more complete control. The third strategy utilized FeNO to drive asthma control. The study examined 611 patients. The results showed that the FeNO driven strategy reduced asthma medication use while sustaining asthma control and quality of life and was the preferred strategy for adult asthmatic patients in primary care. The FeNO driven strategy had the highest probability of cost-effectiveness.

In July 2014, the *Annals of Allergy Asthma and Immunology* published a study that examined whether the use of FeNO in specialist management of asthma could result in more effective and cost-effective treatment decisions. Fifty patients were evaluated by clinical examination, spirometry and symptom assessment using the Asthma Control Test (ACT). Physicians made a decision on the level of airway inflammation present along with treatment before and then after receiving results of the FeNO test. Without FeNO, airway inflammation was incorrectly classified in 50% of the patients. FeNO substantially altered treatment decisions in more than one third of the patients, adding medication in 20% and decreasing medication in 16%. The use of FeNO in addition to standard of care was estimated to save \$629 per patient per year.

Also, in July 2014, the results of a clinical trial were published in *the International Journal of COPD*. The study was established to characterize FeNO levels in patients with Chronic Obstructive Pulmonary Disease (COPD). Increases in FeNO were identified in a subset of patients with COPD, particularly in those previously diagnosed with both COPD and asthma. The authors of the study concluded that "since FeNO is useful for identifying patients with airway inflammation who will have a beneficial response to treatment with an inhaled corticosteroid, these data may have important implications for the management of COPD patients".

Investments and Cash flow

The Group's cash reserves amounted to SEK 146.3m (334.0) at the end of the period.

Cash flow for the period January - September 2014 was negative in the amount of SEK 157.7m (+132.7) and cash flow from current operations was negative in the amount of SEK 154.7m (-169.0). Cash flow for the third quarter 2014 was negative in the amount of SEK 50.5m (-62.2) and the cash flow from current operations was negative in the amount of SEK 49.5m (-50.8). The total cash flow in the prior year was positively impacted by a new share issue of SEK 91.4m and borrowings under a loan facility of SEK 223.4m in the second quarter in 2013.

The cash flow from current operations has been negatively impacted in the period by interest paid of SEK 22.0m (12.4) associated with borrowings under the 2013 loan facility as well as increased investments in the US and other changes in working capital primarily increased receivables associated with increased sales.

The Group's investments in tangible assets for the period amounted to SEK 1.0m (1.8). Investments in intangible assets for the period amounted to SEK 1.9m (0.7).

Parent Company

The Group's principal operations, including development, marketing and sales, are conducted by the Parent Company, Aerocrine AB. The Parent Company assumes the Group's market risk while the subsidiaries, Aerocrine Inc. and Aerocrine AG, are sales companies with the objective of conducting marketing and sales activities in the US and German markets respectively. In addition to its sales activities, Aerocrine Inc. also conducts service operations. In connection with the introduction of the Group's personnel stock options programme, Aerocrine ESOP AB was founded. During 2013 Aerocrine International GmbH, Switzerland, was formed with the purpose of supporting the European market.

The Parent Company's net sales for the year amounted to SEK 119.4m (97.9), of which sales to Group

companies amounted to SEK 58.6m (53.8). The loss after financial items for the period amounted to SEK 160.4m (156.5). The Parent Company's cash and equivalents amounted to SEK 130.6m (324.7) at the end of the period. Investments in machinery and equipment for the period amounted to SEK 1.0m (1.4) and investments in intangible assets amounted to SEK 1.9m (0.7). The earnings of the Parent Company were affected negatively by the Group's internal pricing model, whereby the Parent Company assumes all market risk and consequently makes marketing contributions to the subsidiaries to establish and develop their respective markets.

The internal pricing model with marketing support from the parent means that the equity of the parent company is consumed in approximately the same rate as for the Group. The board continually monitors the equity in relation to the shareholder's equity in regards to the potential need to establish a balance sheet for liquidation purposes. The board has made the assessment that there is additional value related to intellectual property rights which was confirmed by the balance sheet for liquidation purposes as per June 30, 2014. See further information below under Balance Sheet for Liquidation Purposes.

Ownership Status

As of September 30 2014, Aerocrine AB had approximately 4,706 shareholders, of whom the four largest represented approximately 61.4% of the votes and capital. On September 30, 2014, the total number of registered shares in the Group was 155,063,162. The largest owners in the Group on September 30, 2014 were Novo A/S (25%), Invifed AB (23%), HealthCap Holding KB (10%) and the Third AP-Fund (3%).

Employees

At the end of the period, the total number of employees in the Group amounted to 118 (136), of whom 39 (43) are employed in Sweden.

Financing

On a regular basis the Board reviews the Company's current and projected cash position in order to ensure that the Company has the means and resources necessary to carry out the operations and strategies as directed by the Board. The Company's long-term liquidity needs will largely be determined by the success of products already being commercialized, key development and regulatory events that might impact the ability to sell the Company's products or which could impact the reimbursement rates associated with use of the Company's products, and expenses associated with these same efforts.

Based on current forecasts, and absent a strategic collaboration or larger than expected growth in revenue, existing cash is sufficient to finance the current scope of operations until approximately the end of the second quarter 2015; and sufficient cash to meet the conditions of the credit agreement at least the end of December 2014. The Board of Directors, believing that additional value can be created for the Company's shareholders by continuing to fund the Company's operations and growth initiatives, and having reviewed alternatives for raising additional capital, has resolved to secure financing of approximately SEK 443m, before transaction costs. The financing will be structured as a rights offering, which will be subject to approval by the shareholders at an EGM to be held on January 7, 2015. Details of the rights offering will be communicated via a press release on November 28, 2014. In excess of 75% of the financing has been guaranteed by inter alia Aerocrine's largest shareholder Novo A/S and the largest Danish public pension fund, Arbejdsmarkedets Tillægspension (ATP). The objective for the upcoming financing will be to enhance the financial flexibility and facilitate the company's ability to realise its growth plans and reach profitability, and thereby create additional shareholder value. After a successful implementation of the upcoming rights offering, the Board of Directors and management believe that the Company will have sufficient liquidity to attain positive cashflow without additional financing.

There was no conversion of options during the quarter.

Balance Sheet for Liquidation Purposes

As the Company's equity as of 30 June 2014 was less than half of the registered share capital, the board decided to draw up a balance sheet for liquidation purposes as of 30 June 2014 for evaluation by its auditor. According to this balance sheet for liquidation purposes, the Company's share capital was intact.

Transactions with Related Parties

During the period interest amounting to SEK 6.3m has been paid to Novo A/S. In the balance sheet existing loans from Novo A/S amounts to SEK 70.3m. This relates to the combined equity and loan financing during

2013 in which debt was taken out with a consortium of OrbiMed and Novo A/S.

Accounting Principles

This interim report has been prepared in accordance with IAS 34 and the Swedish Financial Accounting Standards Council's guideline RFR 1 and, in relation to the Parent Company, RFR 2.

New Accounting Principles 2014

In May 2014, the IFRS issued International Financial Reporting Standard: IFRS 15 *Revenue from Contracts with Customers*. IFRS 15 establishes a comprehensive framework for determining *when* to recognize revenue and *how much* revenue to recognize. The core principle in that framework is that a company should recognize revenue to depict the transfer of promised goods or services to the customer in an amount that reflects the consideration to which the company expects to be entitled in exchange for those goods or services. The Company is currently evaluating the impact of the provisions of the new standard on its revenue recognition policies. The standard is effective 1 January 2017.

In other regards, the accounting principles and valuation methods remain unchanged compared with the description provided in the 2013 Annual Report.

Significant Risks and Uncertainty Factors

The principal risks and sources of uncertainty for Aerocrine include, albeit not exclusively, financial risks associated with the ability to raise sufficient capital in the upcoming financing to meet planned operational activities and debt covenants and other financial risks, such as the amount of future earnings, ability to secure additional financing – if and when needed and at a reasonable cost, and currency and credit risks. In addition to market risks, there are also risks associated with Aerocrine's operations, such as obtaining the necessary approval from authorities, product development, patents and intellectual property rights, product responsibility and forward looking information, which can affect the Company. Further information on the Company's risk exposure can be found on pages 25-27 of Aerocrine's 2013 Annual Report part 1.

Publication Dates 2014/2015

Year-end Report 2014	20 February 2015 08.00 a.m.
2014 Annual Report	13 April 2015
First Quarter Interim Report	12 May 2015 08.00 a.m.
AGM 2015	12 May 2015 05:00 p.m.
Second Quarter Interim Report	24 July 2015 08.00 a.m.
Third Quarter Interim Report	6 November 2015 08.00 a.m.

Solna, November 27, 2014

The Board of Directors and the President provide their assurance that this interim report provides an accurate overview of the operations, position and earnings of the Group and the Parent Company, and that it also describes the principal risks and sources of uncertainty faced by the Parent Company and its subsidiaries.

Rolf Classon

Chairman of the Board

Lars Gustafsson

Board Member

Staffan Lindstrand

Board Member

Scott Myers

President and CEO

Dennis Kane

Board Member

Michael Shalmi

Board Member

Maria Strömme

Board Member

Report of Review of Interim Financial Information

Introduction We have reviewed the condensed interim financial information (interim report) of Aerocrine AB as of 30 September 2014 and the nine-month period then ended. The Board of Directors and the CEO are responsible for the preparation and presentation of the interim financial information in accordance with IAS 34 and the Swedish Annual Accounts Act. Our responsibility is to express a conclusion on this interim report based on our review.

Scope of Review We conducted our review in accordance with the International Standard on Review Engagements ISRE 2410, *Review of Interim Report Performed by the Independent Auditor of the Entity*. A review consists of making inquiries, primarily of persons responsible for financial and accounting matters, and applying analytical and other review procedures. A review is substantially less in scope than an audit conducted in accordance with International Standards on Auditing, ISA, and other generally accepted auditing standards in Sweden. The procedures performed in a review do not enable us to obtain assurance that we would become aware of all significant matters that might be identified in an audit. Accordingly, we do not express an audit opinion.

Conclusion Based on our review, nothing has come to our attention that causes us to believe that the interim report is not prepared, in all material respects, in accordance with IAS 34 and the Swedish Annual Accounts Act, regarding the Group, and with the Swedish Annual Accounts Act, regarding the Parent Company.

Stockholm, 27 November 2014
Öhrlings PricewaterhouseCoopers
Mikael Winkvist
Authorized Public Accountant

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CONSOLIDATED INCOME STATEMENT

(SEK m)	Aerocrine Group				
	Jul 1, 2014 - Sep 30, 2014	Jul 1, 2013 - Sep 30, 2013	Jan 1, 2014 - Sep 30, 2014	Jan 1, 2013 - Sep 30, 2013	Jan 1, 2013 - Dec 31, 2013
Net sales	38,520	31,138	116,049	100,528	136,168
Cost of goods sold	-12,355	-8,226	-36,434	-27,462	-38,338
Gross Profit/Loss	26,165	22,912	79,615	73,066	97,830
Sales and marketing expenses	-41,418	-42,969	-123,261	-131,139	-170,082
Administration expenses	-12,584	-12,353	-40,007	-44,602	-58,997
Development expenses	-13,849	-15,825	-47,503	-52,095	-75,127
Other operating income	624	1,081	1,959	2,786	3,652
Other operating expenses	-707	-790	-931	-2,098	-2,986
Operation Profit/Loss	-41,770	-47,944	-130,128	-154,082	-205,710
Financial income	8,447	11,011	16,752	19,904	28,454
Financial expenses	-26,847	-19,524	-54,366	-32,196	-48,182
Profit/loss before taxes	-60,170	-56,457	-167,742	-166,374	-225,438
Income tax	-	-	-	-81	-160
Profit/loss for the period	-60,170	-56,457	-167,742	-166,455	-225,598
Net profit/loss attributable to:					
Parent company shareholders	-60,170	-56,457	-167,742	-166,455	-225,598
Earnings per share based on Net Profit attributable to parent company shareholders (in SEK per share)					
Profit/loss per share (before and after dilution)*	-0,4	-0,4	-1,1	-1,1	-1,5
* Profit/loss per share after dilution is not reported, since this would imply improved earnings per share.					
Other information:					
Average number of shares outstanding	155,063,162	154,581,405	154,896,644	150,302,696	151,381,295
Amortisation/depreciation included in operating expenses	2,878	2,689	9,196	8,773	11,880
- of which intangible assets	2,473	2,312	7,879	7,548	10,185
- of which tangible fixed assets	405	377	1,317	1,225	1,695

CONSOLIDATED STATEMENT OF COMPREHENSIVE INCOME

(SEK m)	Aerocrine Group				
	Jul 1, 2014 - Sep 30, 2014	Jul 1, 2013 - Sep 30, 2013	Jan 1, 2014 - Sep 30, 2014	Jan 1, 2013 - Sep 30, 2013	Jan 1, 2013 - Dec 31, 2013
Profit/loss for the period	-60,170	-56,457	-167,742	-166,455	-225,598
Other comprehensive income for the period:					
Items that will not be reclassified to profit or loss:	-	-	-	-	-
Reassessment of net pension obligation	-	-	-	-	-71
Items that have or may be reclassified to profit or loss:	-	-	-	-	-
Translation differences on foreign operations	1,035	-167	1,582	-411	293
Sum other comprehensive income for the period, net after taxes	1,035	-167	1,582	-411	222
Total comprehensive income for the period	-59,135	-56,624	-166,160	-166,866	-225,376
Total comprehensive income attributable to:					
Parent company shareholders	-59,135	-56,624	-166,160	-166,866	-225,376

CONSOLIDATED BALANCE SHEET

(SEK m)	Aerocrine Group		
	Sep 30, 2014	Sep 30, 2013	Dec 31, 2013
ASSETS			
Fixed Assets			
Intangible assets	22,844	31,047	28,830
Tangible fixed assets	6,251	6,339	6,226
Financial fixed assets	2,125	1,606	1,911
Total Tangible Assets	31,220	38,992	36,967
Current assets			
Inventories	25,998	21,917	19,513
Current receivables and prepaids	37,276	33,171	30,968
Cash equivalents	146,326	334,044	292,133
Total current assets	209,600	389,132	342,614
Total assets	240,820	428,124	379,581
SHAREHOLDERS' EQUITY			
Capital and reserves attributable to: Shareholders' equity attributable to parent company shareholders	-58,190	162,162	104,186
LIABILITIES			
Long term liabilities and Provisions			
Pension commitments	1,546	1,331	1,453
Provisions for payroll overheads, staff option schemes	2,321	6,642	4,117
Provisions, other	1,433	1,407	1,397
Loan	245,907	212,884	215,755
Long-term liabilities	245,907	212,884	215,755
Long term liabilities and Provisions	251,207	222,264	222,722
Current liabilities	47,803	43,698	52,673
Total shareholders' equity and liabilities	240,820	428,124	379,581

CONSOLIDATED STATEMENT OF CHANGES IN SHAREHOLDERS' EQUITY

(SEK m)	Attributable to Shareholders in the Parent Company					
	Share Capital	Other Capital Contributions	Cumulative Translation Differences	Revaluation of Net Pension Obligation	Accumulated Loss	Total Shareholders' Equity
Opening balance at January 1 2013	72,819	1,378,493	366	-348	-1,219,764	231,566
Comprehensive income						
Net earnings/Loss for the period	-	-	-	-	-166,455	-166,455
Other comprehensive income						
Reassessment of net pension obligation	-	-	-	-	-	-
Translation differences foreign operations	-	-	-411	-	-	-411
<i>Sum other comprehensive income</i>	-	-	-411	-	-	-411
Total comprehensive income	-	-	-411	-	-166,455	-166,866
Transactions with shareholders						
New share issue	4,472	90,563	-	-	-	95,035
Issue expenses	-	-3,665	-	-	-	-3,665
Convertible bond	-	-	-	-	-	-
<i>Staff option scheme:</i>						
-value of employee services	-	-	-	-	6,092	6,092
Total transactions with shareholders	4,472	86,898	-	-	6,092	97,462
Closing balance, September 30 2013	77,291	1,465,391	-45	-348	-1,380,127	162,162
Opening balance at January 1 2014	77,314	1,465,391	659	-419	-1,438,759	104,186
Comprehensive income						
Net earnings/Loss for the period	-	-	-	-	-167,742	-167,742
Other comprehensive income						
Reassessment of net pension obligation	-	-	-	-	-	-
Translation differences foreign operations	-	-	1,582	-	-	1,582
<i>Sum other comprehensive income</i>	-	-	1,582	-	-	1,582
Total comprehensive income	-	-	1,582	-	-167,742	-166,160
Transactions with shareholders						
New share issue	218	-	-	-	-	218
Issue expenses	-	-	-	-	-	-
Convertible bond	-	-	-	-	-	-
<i>Staff option scheme:</i>						
-value of employee services	-	-	-	-	3,566	3,566
Total transactions with shareholders	218	-	-	-	3,566	3,784
Closing balance, Sep 30 2014	77,532	1,465,391	2,241	-419	-1,602,935	-58,190

CONSOLIDATED CASHFLOW STATEMENT

(SEK m)	Aerocrine Group		
	Jan 1, 2014 Sep 30, 2014	Jan 1, 2013 Sep 30, 2013	Jan 1, 2013 Dec 31, 2013
Cash flow from operating activities before change in working capital	-137,937	-155,768	-212,374
Total change in working capital	-16,724	-13,265	295
Cash flow from operating activities	-154,661	-169,033	-212,079
Cash flow from investment activities	-3,080	-2,528	-3,605
Cash flow from financing activities	3	304,254	305,338
Cash flow for the period	-157,738	132,693	89,654
Increase/Decrease in cash equivalents			
Cash equivalents at start of the year	292,133	199,913	199,913
Exchange rate differences in cash equivalents	11,931	1,438	2,566
Cash equivalents at end of the period	146,326	334,044	292,133

KEY RATIOS	Aerocrine Group		
	Jan 1, 2014 -	Jan 1, 2013 -	Jan 1, 2013 -
	Sep 30, 2014	Sep 30, 2013	Dec 31, 2013
Net sales SEK ths	116,049	100,528	136,168
Gross margin %	69%	73%	72%
Return on average shareholders' equity %	neg	neg	neg
Equity/Asset ratio %	-24%	38%	27%
Net indebtness, multiple	-1,71	-0,75	-0,73
Liquid ratio %	384%	891%	613%
Average number of employees	117	136	133
Investments, SEK ths	2,866	2,528	3,320
Expenses related to development, SEK ths	47,503	52,095	75,127
Development expenses in % of total expenses	22%	23%	25%

Data per share	Aerocrine Group		
	Jan 1, 2014 -	Jan 1, 2013 -	Jan 1, 2013 -
	Sep 30, 2014	Sep 30, 2013	Dec 31, 2013
Number of shares at closing of period (before dilution)	155,063,162	154,581,405	154,628,698
Number of shares at closing of period (after dilution) ¹⁾	156,952,372	164,266,829	158,276,053
Average number of shares (before dilution)	154,896,644	150,302,696	151,381,295
Average number of shares (after dilution) ¹⁾	157,511,763	160,294,756	156,041,724
Shareholders' equity per share SEK, before full dilution	-0,38	1,05	0,67
Shareholders' equity per share SEK, after full dilution	-0,37	0,99	0,66
Earnings ¹⁾ per share, SEK (before dilution) ¹⁾	-1,1	-1,1	-1,5

¹⁾Profit/loss per share after dilution is not reported, since this would imply improved earnings per share.

Definitions

Gross margin

Gross profit as a percentage of net sales for the period

Return on average shareholders' equity %

Profit/loss as a percentage of average shareholders' equity

Average number of shares

Number of shares adjusted for share issues conducted during the year (before dilution) and option programmes outstanding (after dilution)

Net indebtness

Interest-bearing liabilities less current investments and cash and equivalents divided by shareholders' equity

Equity/Asset ratio

Shareholders' equity as a percentage of total assets

Earnings per share

Net profit/loss divided by average number of shares before and after full dilution

Shareholders' equity per share

Shareholders' equity (adjusted for dilution effects) divided by the number of shares at the close of the period before and after full dilution

Liquid ratio

Current assets, excluding inventories and work in progress, in relation to current liabilities

QUARTERLY FINANCIAL INFORMATION

(SEK m)	Aerocrine Group									
	Q3-2014	Q2-2014	Q1-2014	Q4-2013	Q3-2013	Q2-2013	Q1-2013	Q4-2012	Q3-2012	
Net sales for the period	38,520	42,501	35,028	35,640	31,138	35,651	33,739	36,180	34,838	
Gross profit/loss	26,165	28,960	24,490	24,764	22,912	25,002	25,152	26,381	25,996	
Gross margin %	68%	68%	70%	69%	74%	70%	75%	73%	75%	
Operating expenses for the period	-67,935	-72,706	-69,102	-76,392	-70,856	-77,608	-78,684	-89,897	-82,084	
Operating profit/loss for the period	-41,770	-43,746	-44,612	-51,628	-47,944	-52,606	-53,532	-63,516	-56,088	
Profit/loss from financial investments	-18,400	-10,928	-8,286	-7,436	-8,513	-3,109	-670	611	-1,985	
Profit/loss for the period, before taxes	-60,170	-54,674	-52,898	-59,064	-56,457	-55,715	-54,202	-62,905	-58,073	
Taxes	-	-	-	-79	-	-1	-80	-	-	
Profit/Loss after taxes	-60,170	-54,674	-52,898	-59,143	-56,457	-55,716	-54,282	-62,905	-58,073	

AEROCRINE GROUP SEGMENT FINANCIAL INFORMATION

Segment - Net sales	Jul 1, 2014 - Sep 30, 2014				Jul 1, 2013 - Sep 30, 2013			
	US/NA*	A/P*	EJ/ROW	Total	US/NA*	A/P*	EJ/ROW	Total
Net sales from external customers	16,048	5,672	16,800	38,520	15,251	1,325	14,562	31,138
Total net sales	16,048	5,672	16,800	38,520	15,251	1,325	14,562	31,138

Segment - Net sales	Jan 1, 2014 - Sep 30, 2014				Jan 1, 2013 - Sep 30, 2013			
	US/NA*	A/P*	EJ/ROW	Total	US/NA*	A/P*	EJ/ROW	Total
Net sales from external customers	40,255	19,467	56,327	116,049	43,814	7,727	48,987	100,528
Total net sales	40,255	19,467	56,327	116,049	43,814	7,727	48,987	100,528

Segment - Profitability measure	Jul 1, 2014 - Sep 30, 2014					Jul 1, 2013 - Sep 30, 2013				
	US/NA*	A/P*	EJ/ROW	Unallocated	Total	US/NA*	A/P*	EJ/ROW	Unallocated	Total
Total EBIT for reportable segment	-33,885	1,464	-9,349	-	-41,770	-31,416	-2,625	-13,903	-	-47,944
Financial income	-	-	-	8,447	8,447	-	-	-	11,011	11,011
Financial expenses	-	-	-	-26,847	-26,847	-	-	-	-19,524	-19,524
Group - earnings before tax	-33,885	1,464	-9,349	-18,400	-60,170	-31,416	-2,625	-13,903	-8,513	-56,457
<i>Negative impact on earnings:</i>										
- cost personnel option plan	-1,001	-	175	-	-826	-1,313	-	-1,267	-	-2,580
- depreciation	-150	-	-2,728	-	-2,878	-141	-	-2,548	-	-2,689

Segment - Profitability measure	Jan 1, 2014 - Sep 30, 2014					Jan 1, 2013 - Sep 30, 2013				
	US/NA*	A/P*	EJ/ROW	Unallocated	Total	US/NA*	A/P*	EJ/ROW	Unallocated	Total
Total EBIT for reportable segment	-104,281	5,919	-31,766	-	-130,128	-95,657	-3,085	-55,340	-	-154,082
Financial income	-	-	-	16,752	16,752	-	-	-	19,904	19,904
Financial expenses	-	-	-	-54,366	-54,366	-	-	-	-32,196	-32,196
Group - earnings before tax	-104,281	5,919	-31,766	-37,614	-167,742	-95,657	-3,085	-55,340	-12,292	-166,374
<i>Negative impact on earnings:</i>										
- cost personnel option plan	-2,614	-	-952	-	-3,566	-3,307	-	-501	-	-3,808
- depreciation	-430	-	-8,766	-	-9,196	-427	-	-8,346	-	-8,773

* US/NA = USA and North America, A/P = Asia/Pacific region

PARENT COMPANY INCOME STATEMENT

(SEK m)	Jan 1, 2014 - Sep 30, 2014	Jan 1, 2013 - Sep 30, 2013	Jan 1, 2013 - Dec 31, 2013
Net sales	119,378	97,878	139,369
Cost of goods sold	-38,339	-28,656	-41,702
Gross Profit/loss	81,039	69,222	97,667
Sales and marketing expenses	-139,863	-141,452	-206,223
Administration expenses	-27,438	-29,854	-37,722
Development expenses	-37,639	-43,208	-63,400
Other operating income	1,565	2,535	3,148
Other operating expenses	-917	-1,993	-2,140
Operation Profit/loss	-123,253	-144,750	-208,670
Earnings from shares in Group companies	-	20,409	-1,187
Financial income	17,199	-32,196	29,448
Financial expenses	-54,358	-	-48,179
Profit/ Loss from financial items	-37,159	-11,787	-19,918
Loss after financial items	-160,412	-156,537	-228,588
Taxes	-	-	-
Loss for the period	-160,412	-156,537	-228,588

PARENT COMPANY STATEMENT OF COMPREHENSIVE INCOME

(SEK m)	Jan 1, 2014 - Sep 30, 2014	Jan 1, 2013 - Sep 30, 2013	Jan 1, 2013 - Dec 31, 2013
Loss for the period	-160,412	-156,537	-228,588
Other comprehensive income	-	-	-
Total comprehensive income	-160,412	-156,537	-228,588

PARENT COMPANY BALANCE SHEET

(SEK m)	Sep 30, 2014	Sep 30, 2013	Dec 31, 2013
ASSETS			
Fixed Assets			
Intangible Assets	22,844	31,047	28,830
Tangible Assets	2,369	2,060	2,205
Financial Assets	47,063	37,827	37,825
Total fixed assets	72,276	70,934	68,860
Current assets			
Inventory	19,536	16,996	13,810
Current receivables and prepaids	23,798	20,418	19,256
Cash equivalents	130,598	324,741	283,686
Total current assets	173,932	362,155	316,752
Total assets	246,208	433,089	385,612
SHAREHOLDERS EQUITY			
Shareholders' equity	-40,015	188,130	116,613
LIABILITIES			
Long term liabilities and provisions			
Provisions			
Provisions for guarantees	2,283	600	1,800
Provisions for payroll overheads, staff option schemes	2,321	6,642	4,117
Loan	245,907	212,884	215,755
Long term liabilities and provision	250,511	220,126	221,672
Current liabilities	35,712	24,833	47,327
Total shareholders' equity and liabilities	246,208	433,089	385,612
Pledged assets	408,058	594,939	547,462
Contingent liabilities	none	none	none