

INTERIM REPORT

SECOND QUARTER 2017



Improving the lives of patients with serious diseases by being a science driven company with a long-term commitment to commercialize differentiated next generation medicines



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Affibody Medical AB (publ) (556714-5601)
Gunnar Asplunds Allé 24
SE-171 69 Solna, Sweden

E-mail: reception@affibody.com
Phone: +46 (0) 8 59 88 38 00

www.affibody.com

Key Events during the Second Quarter 2017

Affibody Medical AB (publ) (“Affibody” or “the Company”), a Swedish biotech company focused on developing next generation biopharmaceuticals based on its unique proprietary technology platforms: Affibody® molecules and Albumod™, today issued its Interim Report for the second quarter 2017.

Financial Highlights

- » Revenue for the 2nd Quarter 2017 amounted to SEK 26.2 (33.6) m, and for the six month period to 57.4 (54.6) m
- » Operating result for the quarter amounted to SEK -19.7 (9.2) m, and for the six month period to -17.4 (-2.3) m
- » EBITDA for the quarter amounted to SEK -19.4 (9.3) m, and for the six month period to -16.8 (-2.0) m
- » Net result for the quarter amounted to SEK -19.6 (9.4) m, and for the six month period to -17.3 (-2.3) m
- » Cash flow for the quarter amounted to SEK -54.2 (105.0) m, and for the six month period to -36.2 (125.2) m
- » Cash and cash equivalents at the end of the period amounted to SEK 90.9 (132.3) m.

Significant Events during the Reporting Period

- » In May it was announced that we have decided to initiate a Phase II development with ABY-035, our proprietary psoriasis program.
- » During the quarter MedImmune terminated the agreement primarily as a result of changing R&D priorities.

Significant Events during the rest of the Year

- » During the first quarter, the first patient was dosed and evaluated in the clinical trial with ABY-029, to guide cancer surgery.

Significant Events after the Close of the Reporting Period

- » In August we filed the clinical trial application (CTA) to progress our psoriasis compound ABY-035 to Phase II.

SEKk	2017 (3m)	2016 (3m)	2017 (6m)	2016 (6m)	2016 (12m)
Revenue	26 245	33 645	57 362	54 582	104 607
Operating result	-19 683	9 154	-17 392	-2 259	-7 515
Operating margin	-75%	27%	-30%	-4%	-7%
Net result	-19 640	9 357	-17 300	-2 300	-7 494

CEO Statement

The second quarter and the period up until now have been eventful and exciting. In this report we are very pleased to announce that we recently, in August, filed a clinical trial application (CTA) to progress our psoriasis compound ABY-035 to Phase II. The filing was made to the German regulatory agency BfArM. The Phase II study will include ~100 moderate-to-severe psoriasis patients from multiple sites in Germany. This study will provide valuable results to validate both the technology platform and ABY-035 and will, if positive, bring Affibody further along the path to becoming a leading biopharmaceutical company. It should be noted that the design has been carefully matched to the uniqueness of ABY-035. We have also continued to dose patients in the multi-dosing arm of our ongoing Phase I/II study and expect to collect additional efficacy results during the year. ABY-035 has been specifically developed to utilize the strengths of Affibody's technology.

The period up until August has also been eventful in our other projects. In the autoimmune disease project ABY-039 a scientific advice meeting with UK's regulatory agency MHRA was held in preparation for a CTA. The actual filing is planned for Q4 and we expect to be able to start dosing before the end of the year. The ABY-039 program has clear differentiating features that are directly related to the core of Affibody's technology. The Phase I study will provide important safety as well as biomarker efficacy data that will help us fully capitalize on the value of this substance.


Our metastatic breast cancer imaging program ABY-025 is also moving forward in development and a ~100 patient Phase II/III study is planned to be initiated in the latter part of the year.

As the results from our development programs are emerging the transformation of Affibody continues. We look forward to further develop the company and our technology to create value to our shareholders.

Solna, August 2017

David Bejker
President and CEO





Our mission is to advance healthcare by improving the lives of patients with serious diseases by being a science driven company with a long-term commitment to commercialize differentiated next generation medicines.

About Affibody

Affibody is a Swedish biotech company focused on developing next generation biopharmaceuticals based on its unique proprietary technology platforms: Affibody[®] molecules and Albumod[™].

Affibody is developing a portfolio of innovative drug projects and, in addition, offers the half-life extension technology, Albumod[™], for outlicensing.

The company is currently developing four proprietary programs. The first three are therapeutic programs that targets liver diseases, autoimmune diseases, and psoriasis respectively. The fourth program is a diagnostic imaging program that is directed primarily towards metastatic breast cancer.

Affibody also has ongoing commercial relationships with several companies such as AbClon, Biotest, Daewoong, Daiichi Sankyo, GE Healthcare, Nordic Nanovector, and Swedish Orphan Biovitrum.

In addition, Affibody is working in collaboration with other companies and academic institutions in a number of grant funded projects. Affibody was founded in 1998 by researchers from the Royal Institute of Technology and the Karolinska Institute and is based in Solna, Sweden. The major shareholder in the company is Investor AB. Further information can be found at: www.affibody.com

Mission

Our mission is to advance healthcare by improving the lives of patients with serious diseases by being a science driven company with a long-term commitment to commercialize differentiated next generation medicines.

Business Model

Affibody shall operate a long-term business that develops and commercializes innovative products based on the company's technology platforms independently and with partners.

Strategy

We develop and commercialize differentiated therapies by having a product vision focusing on unmet needs. We do so by identifying projects where the strengths of our proprietary technology platforms can be leveraged to transform the lives of patients with serious diseases. We aim to independently commercialize our products and will selectively complement this with partner-based development and commercialization. Operations are conducted by highly qualified resources in research and development which are supported by an extensive network of renowned researchers and clinicians.

Operational Review

Proprietary Programs

ABY-035 - Psoriasis

ABY-035 addresses the substantial non-TNF market segment in psoriasis. A first-in-human study is ongoing to establish clinical safety and first signs of efficacy. The CTA was filed in the fourth quarter 2015 and in May 2016 we announced that the dose-escalation part of the Phase I study was completed and that initial results confirm the compound to be safe and well-tolerated across all doses in healthy volunteers. A regulatory filing to commence a 100 patient multicenter Phase II study in Germany was submitted to the German authorities in August, 2017.

ABY-039 - Autoimmune Diseases

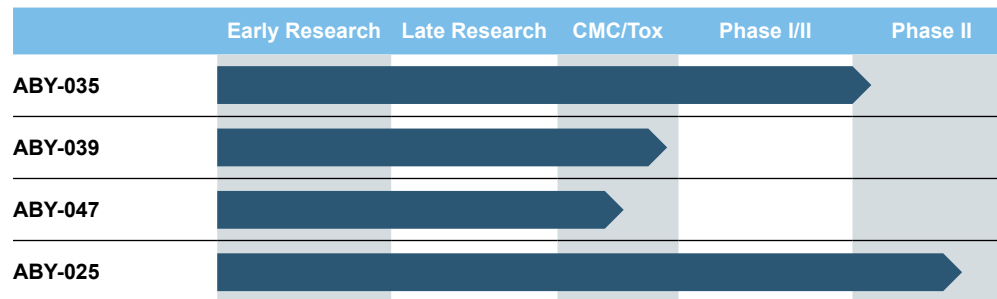
The goal of the ABY-039 project is to offer a treatment for people suffering from antibody mediated autoimmune diseases. Preclinical PoC has been demonstrated in animals with a lead Affibody® molecule and process development has been initiated. A successful scientific advice meeting with UK's regulatory agency MHRA was held in August 2017.

ABY-047 - Liver disease

The goal with ABY-047 is to develop a treatment within a broad spectrum of inflammatory diseases, initially within liver diseases. Preclinical PoC has been demonstrated in animals.

ABY-025 - Breast Cancer Imaging

ABY-025 provides a new non-invasive cost-effective approach to diagnose global HER2-expression in metastatic breast cancer patients using PET imaging. Affibody is currently working together with academic institutions to explore the clinical utility of ABY-025 further.



Collaborations

Products on the Market

GE Healthcare

The product MabSelect Sure™ was launched by GE Healthcare Bio-Sciences AB in 2004, as a result of a collaboration with Affibody for the development of affinity ligands for large scale affinity purification. The product generates royalties and constitutes Affibody's largest revenue source. The product generates royalties until 2019.

Projects in Clinical Development

Daiichi Sankyo

In 2013, Affibody signed an initial license agreement with Daiichi Sankyo regarding the use of Albumod™. The technology will be applied to increase the efficacy of Daiichi Sankyo's proprietary compounds by prolonging the half-life in the circulation. In the lead program the first patient was dosed in a first-in-human clinical trial during the fourth quarter 2015.

Projects in Preclinical Research and Development

AbClon

In 2013, Affibody and AbClon signed a license agreement regarding the use of Affibody® molecules in combination with AbClon's proprietary and/or generic antibodies to create multispecific drugs (AffiMabs).

Biotest

In 2015, Biotest AG and Affibody AB signed a Research License and Option Agreement regarding the use of Albumod™. The technology will be applied to compounds from Biotest's portfolio of proprietary molecules to increase the efficacy by prolonging the half-life in the circulation.

Financial Summary - Second Quarter 2017

Significant Events during the Reporting Period and After Close of the Reporting Period

During the second quarter 2017, the ongoing work with our proprietary programs developed well and according to plan, resulting in substantial and increasing costs for research and development. The most advanced program, ABY-035, is currently in a first-in-human study to establish clinical safety and first signs of efficacy, and a regulatory filing to commence a 100 patient multicenter Phase II study in Germany was submitted to the German authorities in August, 2017. Regarding ABY-039, a successful scientific advice meeting with UK's regulatory agency MHRA was held in August 2017. During the first quarter, the first patient was dosed and evaluated in the clinical trial with ABY-029, to guide cancer surgery.

Revenue

Revenue for the quarter amounted to SEK 26.2 (33.6) m, and for the first six months to SEK 57.4 (54.6) m, where the majority of the revenue comes from royalties and research payments from commercial partners. Revenue for the second quarter in 2016 included a substantial license fee related to Sobi.

Operating Costs

Total operating costs for the quarter amounted to SEK 45.9 (24.5) m, and for the first six months to SEK 74.8 (56.8) m. The costs consisted of research and development costs of SEK 41.7 (20.9) m for the quarter, and for the first six months to SEK 67.2 (45.4) m, mainly related to the accelerated work with our proprietary programs. Administrative costs amounted to SEK 3.9 (2.8) m for the quarter, and for the first six months to SEK 6.9 (9.2) m, and were in 2016 affected by changes in provisions for ESOP related pay-roll taxes. Marketing

and sales costs amounted to SEK 0.4 (0.8) m for the quarter, and for the first six months to SEK 0.6 (2.3) m. Depreciation of fixed assets, included in the operating cost mentioned above, amounted to SEK 0.3 (0.1) m for the quarter, and for the first six months to SEK 0.6 (0.3) m, and were related to laboratory equipment.

Operating Result

The operating result for the quarter amounted to SEK -19.7 (9.2) m, and for the first six months to SEK -17.4 (-2.3) m.

Financial Items

Financial income for the quarter amounted to SEK 0.1 (0.1) m, and for the first six months to SEK 0.1 (0.1) m, and consisted of interest income. Financial costs for the quarter amounted to SEK 0.0 (-0.1) m, and for the first six months to SEK 0.0 (0.1) m, and consisted mainly of fees related to a credit facility.

Taxes

No corporate income tax was reported during the period (-). The Group's unutilized tax losses have not been assigned any value in the balance sheet as they are not expected to be utilized within the conventional period.

Net Result

Net result for the quarter amounted to SEK -19.6 (9.4) m, and for the first six months to SEK -17.3 (-2.3) m.

Cash Flow

Cash flow from current operations, before changes in working capital, amounted to SEK -19.3 (9.8) for the quarter, and for the first six months to SEK -16.7 (3.6) m. The numbers include non-cash items of SEK 0.3 (0.4) m for the quarter, and for the first six months SEK 0.6

(5.9) m, mainly related to the depreciation of tangible assets and, in 2016, employee stock ownership plans. The cash flow from working capital changes for the period amounted to SEK -34.3 (-33.7) m, and for the first six months to SEK -18.9 (-7.4) m, a consequence of the changed payment terms related to royalties from a product. Capital expenditure for the quarter amounted to SEK 0.5 (0.3) m, and for the first six months to SEK 0.5 (0.3) m. The cash flow from financing activities for the quarter amounted to SEK - (129.3) m, and for the first six months to SEK - (129.3) m. Cash flow for the quarter amounted to SEK -54.2 (105.0) m, and for the first six months to SEK -36.2 (125.2) m.

Financial Position

As of June 30, 2017, cash amounted to SEK 90.9 (132.3) m. The equity ratio at the end of the quarter was 87 (91) %.

Shareholders' Equity

Total equity in the Group as of June 30, 2017 was SEK 148.9 (170.1) m.

Significant Risks and Uncertainties

No changes in the company's risk assessment have taken place during the period. A detailed presentation of significant risks and uncertainties is available in the Annual Report.

General Information

Affibody Medical AB (previously Affibody Holding AB) (publ) (registration number 556714-5601) is a public limited company with registered office in Stockholm in Sweden. "Affibody" and "the Company" refers to Affibody Medical AB, and where appropriate, including subsidiaries.

Parent Company

Affibody Medical AB's revenue for the six month period amounted to SEK 2.5 (2.4) m. The costs, mainly consisting of administrative costs in relation to management and financing activities amounted to 4.1 (4.1) m. Net result amounted to SEK -1.5 (-1.6) m. Cash and cash equivalents as of June 30, 2017 amounted to SEK 87.1 (28.1) m and the equity amounted to 369.4 (372.3) m.

Employees

Per June 30, 2017 the number of employees amounted to 32 (30).

Financial Instruments

The extent and nature of financial assets and liabilities are essentially the same as at 31 December 2016. Similar to what was the case at the end of 2016; the recorded values are the same as fair values.

Forward-looking Statement

This interim report includes statements that are forward looking. Actual results may differ from those stated. Internal factors such as the successful management of research and intellectual property rights may affect future results. There are also external conditions such as the economic climate, political changes and competing research that may affect Affibody's results.

Accounting Principles

This report has been prepared in accordance with IAS 34, Interim Financial Reporting. The accounting principles are in accordance with International Financial Reporting Standards (IFRS), as approved by EU and Chapter 9 of the Annual Accounts Act. This report

has been prepared using the same accounting policies and methods of computation as the Annual Report for 2016. No new IFRS standards effective from 2017 have had any effects on Affibody's financial statements. The Parent Company follows the Swedish Financial Reporting Board and the recommendation RFR 2, meaning that the parent company, in the reporting of the legal entity, shall apply all EU-approved IFRS and statements as far as possible within the framework of the Annual Accounts Act, the Pension Obligations Vesting Act and considering the relationship between accounting and taxation.

AGM

The Annual General Meeting (AGM) in 2017 was held on June 29, and reelected Håkan Åström, Jakob Lindberg, Mathias Uhlén and Jonathan Knowles as Directors of the Board.

The Share

As of June 30, 2017 the registered share capital amounted to 67 684 955 SEK divided into 13 536 991 shares. Affibody Medical AB has only one share class and the shares carry one vote each and are entitled to equal shares of distributable earnings.

Other

Amounts are expressed in SEKk (thousands Swedish kronor) unless otherwise stated. Figures in parentheses refer to the corresponding period last year. The Board of Directors and the CEO of Affibody Medical provide their assurance that the interim report provides a fair and true overview of the parent company's and the group's operations, financial position and results, and describes material risks and uncertainties faced by the parent

company and the companies in the group. See under the heading "Significant Risks and Uncertainties" and in other information provided for a description of the operational risks.

Stockholm on August 23, 2017

Håkan Åström
Chairman

Jonathan Knowles
Board Member

Mathias Uhlén
Board Member

Jakob Lindberg
Board Member

David Bejker
President and CEO

This report has not been subject to review by the company's auditor.

For further information please contact:

David Bejker, CEO, Phone: +46 706 454 948
Johan Stuart, CFO, Phone: +46 706 644 096

Financial Calendar

» The interim report for January-September 2017 will be published on Nov 17, 2017

Affibody Medical AB (publ)
Gunnar Asplunds Allé 24
171 69 Solna, Sweden
Phone: +46 8 59 88 38 00
www.affibody.com
Reg 556714-5601

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Financial Statements for the Group

Income Statement

(SEKk)	Apr - Jun 2017	Apr - Jun 2016	Jan - Jun 2017	Jan - Jun 2016	12m 2016
Sales	21 911	31 004	50 952	50 449	97 250
Other revenue	4 334	2 641	6 410	4 133	7 357
Total	26 245	33 645	57 362	54 582	104 607
Operating costs					
Marketing and sales costs	-356	-787	-611	-2 305	-3 387
Administrative costs	-3 892	-2 765	-6 916	-9 184	-10 624
Research and development costs	-41 681	-20 939	-67 227	-45 351	-98 110
Total operating costs	-45 928	-24 491	-74 754	-56 841	-112 121
Operating profit / loss	-19 683	9 154	-17 392	-2 259	-7 515
Net financial items					
Other interest income and similar profit/loss items	51	85	108	86	182
Other interest expense and similar profit/loss items	-8	118	-16	-127	-161
Total net financial items	43	203	92	-42	21
Profit / loss after financial items	-19 640	9 357	-17 300	-2 300	-7 494
Income tax	-	-	-	-	-
Net result	-19 640	9 357	-17 300	-2 300	-7 494
Other comprehensive income					
Comprehensive income	-19 640	9 357	-17 300	-2 300	-7 494

The result is in total attributable to the parent company's shareholders.

Consolidated Balance Sheet

(SEKk)	2017-06-30	2016-06-30	2016-12-31
ASSETS			
Non-current assets			
Property, plant and equipment	4 123	1 430	4 180
Total non-current assets	4 123	1 430	4 180
Current assets			
Other receivables			
Accounts receivable	50 094	42 941	33 182
Other receivables	7 565	5	826
Prepaid expenses and accrued income	19 252	11 242	27 330
Total receivables	76 912	54 188	61 338
Cash and cash equivalents	90 859	132 290	127 020
Total current assets	167 770	186 478	188 358
Total assets	171 893	187 908	192 538

(SEKk)	2017-06-30	2016-06-30	2016-12-31
EQUITY AND LIABILITIES			
Equity			
Share capital	67 685	66 654	67 685
Other capital contribution	694 179	693 905	694 179
Accumulated result including result for the period	-612 994	-590 500	-595 694
Total equity	148 871	170 059	166 170
Non-current liabilities			
Provisions	-	8 353	-
Total non-current liabilities	-	8 353	-
Current liabilities			
Accounts payable	4 877	2 831	13 056
Other payables	2 736	1 971	4 018
Accrued expenses and deferred income	15 410	4 695	9 294
Total current liabilities	23 023	9 497	26 368
Total equity and liabilities	171 893	187 908	192 538

Consolidated Changes in Equity

(SEKk)	Share capital	Other capital contribution	Accumulated losses	Total
Closing balance Dec 31 2015	52 900	578 048	-588 200	42 748
Net result Jan-Jun 2016	-	-	-2 300	-2 300
Employee StockOwnership Plan	-	324	-	324
Rights issue of shares	13 754	115 533	-	129 287
Closing balance Jun 30 2016	66 654	693 905	-590 500	170 059
Closing balance Dec 31 2016	67 685	694 179	-595 694	166 170
Net result Jan-Jun 2017	-	-	-17 300	-17 300
Closing balance Jun 30 2017	67 685	694 179	-612 994	148 871

The equity is in total attributable to the parent company's shareholders.

Cash Flow Analysis

(SEKk)	Apr - Jun 2017	Apr - Jun 2016	Jan - Jun 2017	Jan - Jun 2016	12m 2016
Current operations					
Profit / loss after financial items	-19 640	9 357	-17 300	-2 300	-7 494
Adjustments for non-cash flow items					
Depreciation	298	137	589	271	725
Other non-cash flow items	-	284	-	5 653	-2 425
Cash flow from current operations before income tax	-19 342	9 777	-16 711	3 624	-9 195
Income tax paid	-	-	-	-	-
Cash flow from current operations before changes in working capital	-19 342	9 777	-16 711	3 624	-9 195
Cash flow from working capital changes					
Change in trade, other receivables and current assets	-35 342	-29 789	-15 574	-5 612	-12 761
Change in trade, other payables and other current liabilities	1 017	-3 956	-3 345	-1 789	15 082
Cash flow from current operations	-53 667	-23 967	-35 629	-3 777	-6 874
Investment activities					
Investments in property, plant and equipment	-536	-310	-532	-310	-3 513
Sale of property, plant and equipment	-	-	-	-	-
Cash flow from investment activities	-536	-310	-532	-310	-3 513
Financing activities					
New issue	-	129 287	-	129 287	129 287
Exercise of ESOPs	-	-	-	-	1 031
Cash flow from financing activities	-	129 287	-	129 287	130 318
Cash flow for the period	-54 203	105 010	-36 161	125 200	119 930
Cash and cash equivalents at beginning of period	145 061	27 280	127 020	7 090	7 090
Cash and cash equivalents at end of period	90 859	132 290	90 859	132 290	127 020

Financial Statements for the Parent Company

Income for the Parent Company

(SEKk)	Apr - Jun 2017	Apr - Jun 2016	Jan - Jun 2017	Jan - Jun 2016	12m 2016
Revenue	1 260	1 200	2 520	2 400	4 850
Total	1 260	1 200	2 520	2 400	4 850
Operating expenses					
Marketing and sales costs	-	-	-	-	-
Administrative costs	-2 057	-1 856	-4 091	-4 054	-8 994
Total operating expenses	-2 057	-1 856	-4 091	-4 054	-8 994
Operating profit / loss	-797	-656	-1 571	-1 654	-4 144
Net financial items					
Other interest income and similar profit/loss items	-30	39	-0	39	89
Other interest expense and similar profit/loss items	68	-15	68	-15	-15
Total net financial items	38	24	67	24	73
Profit / loss after financial items	-760	-633	-1 503	-1 630	-4 071
Income tax	-	-	-	-	-
Net loss	-760	-633	-1 503	-1 630	-4 071

Report of Comprehensive Income for the Parent Company

Other comprehensive income	-	-	-	-	-
Comprehensive income	-760	-633	-1 503	-1 630	-4 071

Parent Company Balance Sheet

(SEKk)	2017-06-30	2016-06-30	2016-12-31
ASSETS			
Non-current assets			
Shares in group companies	220 000	220 200	220 000
Total non-current assets	220 000	220 200	220 000
Current assets			
Other receivables			
Accounts receivable	25	-	25
Other receivables	1 120	2	63
Prepaid expenses and accrued income	76	65	32
Receivables from group companies	62 740	125 641	127 240
Total receivables	63 961	125 708	127 360
Cash and cash equivalents	87 140	28 053	27 261
Total current assets	151 101	153 761	154 621
TOTAL ASSETS	371 101	373 961	374 621

(SEKk)	2017-06-30	2016-06-30	2016-12-31
EQUITY AND LIABILITIES			
Equity			
Restricted equity			
Share capital	67 685	66 654	67 685
Total restricted equity	67 685	66 654	67 685
Non restricted equity			
Share premium reserve	330 646	330 646	330 646
Profit/loss brought forward	-27 475	-23 404	-23 404
Accumulated loss for the period	-1 503	-1 630	-4 071
Total non restricted equity	301 668	305 612	303 172
Total equity	369 353	372 266	370 856
Current liabilities			
Accounts payable	223	38	338
Other payables	354	271	1 766
Liabilities to group companies	-	100	-
Accrued expenses and deferred income	1 170	1 286	1 660
Total liabilities	1 747	1 695	3 764
Total equity and liabilities	371 100	373 961	374 621

The Parent Company's Changes in Equity

(SEKk)	RESTRICTED EQUITY		NONE RESTRICTED EQUITY		Total equity
	Share capital	Share premium reserve	Profit/loss brought forward	Accumulated loss for the period	
Closing balance Dec 31 2015	52 900	215 113	-22 435	-969	244 609
Result for the period Jan - Jun 2016	-	-	-	-1 630	-1 630
Accounting of loss 2015	-	-	-969	969	-
Rights issue of shares	13 754	115 533	-	-	115 533
Closing balance Jun 30 2016	66 654	330 646	-23 404	-1 630	372 266
Closing balance Dec 31 2016	67 685	330 646	-23 404	-4 071	370 856
Result for the period Jan - Jun 2017	-	-	-	-1 503	-1 503
Accounting of loss 2016	-	-	-4 071	4 071	-
Closing balance Jun 30 2017	67 685	330 646	-27 475	-1 503	369 353

Cash Flow Statement for the Parent Company

(SEKk)	Apr - Jun 2017	Apr - Jun 2016	Jan - Jun 2017	Jan - Jun 2016	12m 2016
Current operations					
Profit / loss after financial items	-760	-633	-1 503	-1 630	-4 071
Adjustments for non-cash flow items					
Other non-cash flow items	-	-	-	-	-
Cash flow from current operations before income tax	-760	-633	-1 503	-1 630	-4 071
Income tax paid	-	-	-	-	-
Cash flow from working capital changes					
Change in trade, other receivables and current assets	-6 632	-99 730	63 400	-99 497	157
Change in trade, other payables and other current liabilities	-770	-1 155	-2 017	-660	1 410
Cash flow from current operations	-8 161	-101 518	59 879	-101 787	-2 504
Investment activities					
Investments	-	-	-	-	-101 306
Divestments	-	-	-	-	200
Cash flow from investment activities	-	-	-	-	-101 106
Financing activities					
New issue	-	129 287	-	129 287	129 287
Exercise of ESOPs	-	-	-	-	1 031
Cash flow from financing activities	-	129 287	-	129 287	130 318
Cash flow for the period	-8 161	27 769	59 879	27 500	26 708
Cash and cash equivalents at beginning of period	95 301	284	27 261	553	553
Cash and cash equivalents at end of period	87 140	28 053	87 140	28 053	27 261



Affibody Medical AB (publ) (556714-5601)
Gunnar Asplunds Allé 24
SE-171 69 Solna, Sweden

E-mail: reception@affibody.com
Phone: +46 (0) 8 59 88 38 00

www.affibody.com

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