

MOLOGEN AG

MOLOGEN AG:

**Interim Financial Statements
as of March 31, 2010**

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Foreword

Dear Shareholders,

Mologen AG reached further important milestones in the first quarter of the current financial year: In March, we received official approval to conduct a clinical study phase II with the cancer medicine MGN1703. After the cancer medicine in phase Ib has already proven its excellent tolerability, phase II of the study will test the efficacy of the drug in the treatment of metastasized colorectal cancer. The phase II study is planned to start promptly, and first results will be available after an interim assessment as soon as in nine to ten months.

We can also report positive news about our DNA-based vaccines: We received confirmation for subsidies for the pre-clinical development of a vaccine against Hepatitis B. The research project will be conducted in cooperation with the company Synvolux Therapeutics B.V from the Netherlands and is scheduled to be completed by the end of 2012. In addition, MOLOGEN will receive financial support for the development of a DNA-based vaccine against Leishmaniasis. We had already received a first advance payment in the amount of EUR 0.6 million last year. Currently, the first pre-clinical tests are being conducted with the new vaccine.

In the past months, we not only made progress in the development of our product candidates: With the successful execution of a further capital increase in January 2010, we also managed to financially secure all further scheduled activities and clinical studies until 2011.

The economic development of MOLOGEN also proceeded as planned in the first quarter. In total, we spent EUR 1.2 million in the first three months of the financial year for research projects. In the comparison period of the previous year the expenditures totaled EUR 0.7 million. The increase in expenses in R&D is specifically due to the preparations for the clinical phase II with the cancer medicine MGN1703. The deficit increased as expected to EUR -1.6 million in the reporting period as compared to the previous year's value. MOLOGEN's asset position continues to be characterized by a high proportion of liquid funds in the balance sheet total.

We are convinced that the therapeutic approach of immunomodulation will result in a breakthrough for cancer therapies with regard to efficacy and tolerability and have an optimistic outlook toward the financial year 2010. We would like to thank you, our shareholders, for the trust you have put in us.

Dr. Matthias Schroff
Chief Executive Officer

Jörg Petraß
Chief Financial Officer

**Mologen AG,
Berlin**

**Interim management report
for the period from January 1 to March 31, 2010**

Economic environment

General economic environment

There are more and more signs that point towards an improvement of the economic situation. The growth rates of real GDP in the U.S. and Japan in the fourth quarter of 2009 are indicative of a fast recovery in the world economy. In the Eurozone, however, the economic recovery slowed down. The market recovery is still being supported by extensive stabilization measures.

The outlook for the year 2010 remains positive. The Organization for Economic Cooperation and Development (OECD) expects an average growth of 1.9% for its member countries in 2010. The International Monetary Fund (IMF) expects a positive global growth of 3.9% for 2010. The prospects for global trade are equally positive. The IMF expects a significant recovery of +5.8%.

Overall growth remains fragile, however, and shows a highly varied development on a regional basis. In the industrial nations, the developments in the labor market as well as the banking and finance sectors continue to show uncertainties. The emerging markets, in particular, play a major role in the global economic recovery. They were less affected by the recession and are now recovering at a much faster pace.

The IMF expects a growth of 6.0% for emerging market nations. The generally very dynamic Asian economies, for which an economic growth of 8.4% has been forecast for 2010, are led by China with an expected growth of 10% in 2010. In the Western industrial nations, the IMF expects a GDP growth of 2.1% for 2010. The U.S. and Canada, in particular, lead with a forecast growth of 2.7% and 2.6%, respectively, in 2010.

The economy in the Eurozone also continues to recover, albeit at a slower pace. The intermediate projection of the EU Commission on February 25, 2010 forecast a GDP growth of 0.7% for the Eurozone for the entire year of 2010. This development, however, is currently overshadowed by the crisis in Greece.

The seasonally adjusted unemployment rate in the U.S. decreased slightly from 10.1% in October 2009 to 9.7% in February 2010. The seasonally adjusted unemployment rate in the Eurozone at 9.9% in January 2010 was significantly higher than the prior year value of 8.2%.

As per the Fed's decision of March 16, 2010, the key interest rate will be kept in a range of 0% to 0.25%. On March 4, 2010, the European Central Bank likewise decided to leave its prime rate at 1.0%.

Development of pharmaceutical industry and biotechnology sector

The global pharmaceutical market continues its unexpectedly strong development. As recently as October 2009, the market research institute, IMS Health, forecast a growth of 4-6% for the entire year of 2009. In reality, this past year the pharmaceutical market grew worldwide by 7% to USD 837 billion. In 2008, the growth rate was still at 4.8%.

The institute therefore adjusted its forecast upward by one percentage point for the period up to 2014. It now assumes an average annual growth rate of 5% to 8%. A growth of 4% to 6% is expected in 2010. With growth rates of more than 10%, the highest growth is thus expected in the areas of oncology, diabetes, multiple sclerosis and HIV.

Nevertheless, the industry faces enormous challenges. Core topics include the expansion of market shares for generic drugs, budgeting for health expenditure as well as regulatory and technological risks. The patent protection for blockbuster products that is set to expire over the next five years will further increase the innovational pressure on the large pharmaceutical companies. This primarily impacts the U.S., where six of the ten highest-selling products will lose their patent protection in 2011-2012 and will have to fight to protect their market share against generic products.

This will further influence the increasing integration of pharmaceutical and biotechnology companies. Thus, 2009 has been another year with a large number of extensive research and development cooperation projects.

The stock market environment for innovative biotechnology companies such as Mologen AG (hereinafter: MOLOGEN) continues to be seen as favorable over the long-term based on this development. Advancements in the area of clinical development programs are expected to have a positive impact on the perception of the company in the capital markets. To support this process, MOLOGEN upgraded into the Prime Standard at the Frankfurt Stock Exchange in June 2009 and since that time has been listed in this stock exchange segment, which is subject to the highest transparency and reporting requirements of the German Stock Exchange.

In the first three months of 2010, pharmaceutical and biotechnology stocks continued their recovery, as did the general market. On March 31, 2010, the German index of the pharmaceutical industry "DAXsector Pharma & Healthcare" was 7 percentage points above the value at the beginning of the year. The MOLOGEN shares advanced even more in the same period, and on March 31, 2010 they had increased in value by about 34% above levels at the beginning of the year.

Research and Development (R&D)

The objective of MOLOGEN is to develop highly innovative drugs based on proprietary platform technologies to treat cancer and severe infectious diseases. In this regard, important progress was

again made in the research and development strategy of the company in the first three months of the 2010 financial year.

The focus of the R&D activities in the reporting period was the preparation of the phase II clinical study with the DNA-based cancer medicine for colorectal cancer MGN1703. In the previous phase Ib study, MGN1703 had demonstrated a positive safety profile and exhibited superior tolerability. Moreover, the patient responses to the cancer medicine significantly exceeded expectations. While authorization for the phase II study was still in the application stage for Germany and Austria back in December 2009, at the beginning of March 2010 the official approvals were already available. The study will commence shortly; its start is anticipated for May 2010.

Further progress was also made in the preparations for a phase I/II clinical study of the cell-based gene therapy for renal cancer MGN1601. We expect to receive the official approvals for the execution of the study in the third quarter of 2010.

MOLOGEN also made progress in developing a MIDGE®-based DNA vaccine against Leishmaniasis in humans. As a member of an international project consortium, the corporation received a financial subsidy for the years 2009 to 2011 to develop a DNA vaccine against this infectious disease. The subsidy was granted as part of the 7th Framework Programme of the EU. Support for the project, which spans three years, totals EUR 3.0 million. The project was successfully started according to plan in January of 2009, and MOLOGEN had already received an initial advance payment of EUR 0.6 million in 2009. The project progress continues as planned; currently, initial pre-clinical tests are being conducted with the newly developed vaccine.

In addition, MOLOGEN also started a new research project. The project is executed in cooperation with the Dutch company Synvolux Therapeutics B.V. and its objective is to develop a new highly effective vaccine against an infection associated with Hepatitis B viruses. The vaccine is to be available for preventative (prophylactic) use as well as for treatment (therapeutic use). All the necessary pre-clinical studies will be carried out so that by the end of the project the vaccine will be available for testing in clinical trials. The project is being funded by the German Federal Ministry of Education and Research, as part of the EU's EuroTrans-Bio initiative, with around EUR 280,000, which equates to a 50% share of funding. The project is to be concluded by the end of 2012.

The achievement of the milestones in the R&D area provides an important foundation for the further positive development of the company. In the first three months of the financial year 2010, measures and investments in the amount of EUR 1.2 million were therefore carried out to benefit these milestones and were reported in the income statement (comparison period: EUR 0.7 million). The increase in expenses in R&D is specifically due to the preparations for the clinical phase II study with the cancer medicine MGN1703.

Financial performance and financial position

As expected, sales revenue in the first three months of 2010 was at a low level with EUR 18 thousand, similar to the first three months of the previous year (comparison period: EUR 14 thousand). At EUR 3 thousand, other operating income was significantly below the previous year's figure for the first three months (EUR 103 thousand) due to the fact that no subsidies were received in the reporting period.

In comparison to the value from the previous year's period, the loss thus increased in the reporting period to EUR -1,562 thousand (comparison period: EUR -914 thousand). Aside from lower other income, the negative development in operating income was due to increased expenses for materials (EUR -286 thousand; comparison period: EUR -174 thousand) as well as an increase in other operating expenses (EUR -583 thousand; comparison period: EUR -314 thousand). The increase in expenses for materials is primarily due to the purchase of raw materials in conjunction with the manufacturing of investigational medicinal products that are required for the clinical phase II study with MGN1703. In addition, increased expenditure was required for both clinical development programs with MGN1703 and MGN1601. The increase in other operating expenses was caused by an increased utilization of legal and consulting services. Personnel expenses of EUR -614 thousand were above the previous year's level of EUR -429 thousand. The increase is due to restructuring in relation to the new hires for the department of clinical development, one-time effects caused by bonus payments, and a moderate increase in salaries.

The financial status of MOLOGEN presented in the statement of financial position continues to be characterized by a high percentage of liquid assets as part of the balance sheet total. Cash and cash equivalents at the end of the first three months of 2010 were EUR 4,518 thousand (12/31/2009: EUR 6,174 thousand). Equity was at 86%, which corresponds to the value at the end of 2009.

In December 2009, the issuance of 512,000 new shares was affirmed, with the exclusion of subscription rights. The issuance took place by taking partial advantage of the available authorized capital. In January 2010, investments committed but not yet paid in the amount of EUR 10 thousand were deposited and entered in the Commercial Register.

Employees

In comparison to the reporting date in the previous year, the number of employees has not changed significantly. As of March 31, 2010, MOLOGEN had a total of 43 employees (March 31, 2009: 44 employees).

Opportunities and risk statement

The extraordinary revenue opportunities of the MOLOGEN business model are offset by technological, financial, regulatory, patent right-related and sales risks.

There has been no major change in the assessment of possible risks that may have an adverse effect on development compared with the situation outlined in the 2009 annual financial statements.

Forecast

For the current financial year 2010, MOLOGEN bases its forecast on the projections outlined in the Management Report 2009. These are in line with the following objectives:

- Start of a clinical phase II study to examine the efficacy of the cancer medicine MGN1703 for the treatment of metastasized colorectal cancer.

- Obtaining the official approval for the execution of a phase I/II clinical study to examine the safety and efficacy of the cancer medicine MGN1601 for the treatment of advanced renal cancer as well as the start of the study,
- Continuation of the activities as part of an international project consortium for the development of a prophylactic and therapeutic vaccination against Leishmaniasis in humans,
- Achievement of additional development milestones in the veterinary Leishmaniasis project through license partners.

Liquid assets will be sufficient to perform all scheduled activities and clinical studies into the year 2011 if the development proceeds as planned.

Additionally, the corporate strategy has been designed to focus on research and further development of the innovative product pipeline to achieve higher rates of return over the mid- to long-term. To achieve this objective, profit-and-loss-related spending measures and investments will therefore also be necessary in 2010, which will counter short-term positive profit-and-loss development.

Berlin, 11 May 2010

Mologen AG
Board of Directors

Dr. Matthias Schroff
Chief Executive Officer

Jörg Petraß
Chief Financial Officer

Molgen AG, Berlin, Germany
IFRS Statement of financial position as of March 31, 2010

	03/31/2010 in EUR '000	12/31/2009 in EUR '000
ASSETS		
Non-current assets	1,764	1,872
Property, plant and equipment	177	187
Intangible assets	1,583	1,681
Financial assets	0	0
Other non-current assets	4	4
Current assets	4,908	6,710
Cash and cash equivalents	4,518	6,174
Trade receivables	3	5
Inventories	18	20
Other current assets	349	491
Income tax receivables	20	20
Total assets	6,672	8,582
EQUITY AND LIABILITIES		
Non-current liabilities	83	86
Deferred revenues	83	86
Current liabilities	871	1,145
Trade payables	302	513
Other current liabilities and deferred revenue	562	627
Liabilities to banks	7	5
Shareholders' equity	5,718	7,351
Issued capital	10,655	10,143
Deposits to effect the agreed capital increase, entered in the commercial register on January 20, 2010	0	3,574
Capital reserve	31,789	28,798
Accumulated losses	-36,726	-35,164
Total equity and liabilities	6,672	8,582

Molgen AG, Berlin, Germany
IFRS Statement of comprehensive income
for the period of January 1 to March 31, 2010

	01/01/2010	01/01/2009
	-	-
	03/31/2010	03/31/2009
	in EUR '000	in EUR '000
Revenue	18	14
Other operating income	3	103
Cost of materials	-286	-174
Personnel expenses	-614	-429
Depreciation and amortization	-113	-119
Other operating expenses	-583	-314
Operating result	-1,575	-919
Finance income	13	5
Profit for the period before taxes	-1,562	-914
Tax income	0	0
Net loss for the period/ Comprehensive income	-1,562	-914
Net loss carried forward	-35,164	-28,880
Accumulated losses	-36,726	-29,794
Basic earnings per share (in EUR)	-0.15	-0.10
Diluted earnings per share (in EUR)	-	-

Mologen AG, Berlin, Germany
IFRS Statement of cash flows for the period
of January 1 to March 31, 2010

	01/01/2010 to 03/31/2010 in EUR '000	01/01/2009 to 03/31/2009 in EUR '000
Cash flows from operating activities		
Net loss for the period	-1,562	-914
Depreciation and amortization of fixed assets	113	119
Decrease in provisions	0	-58
Other non-cash expenses and income	47	43
Change in trade receivables, inventories and other assets	146	106
Change in trade payables and other liabilities	-274	137
Net cash used in operating activities	-1,530	-567
Cash flow from investing activities		
Payments for property, plant and equipment, classified as investing activities	-5	-14
Purchase of intangible assets, classified as investing activities	0	0
Net cash used in investing activities	-5	-14
Cash flow from financing activities		
Cash receipts from issue of capital	-121	2,611
Net cash used in financing activities	-121	2,611
Foreign currency effect on cash and cash equivalents	0	0
Total change in liquidity (Cash flow)	-1,656	2,030
Cash and cash equivalents at the beginning of the period	6,174	3,324
Cash and cash equivalents at the end of the period	4,518	5,354

Mologen AG, Berlin, Germany
IFRS Statement of changes in equity
as of March 31, 2010

in EUR '000 except for shares

	Issued capital	Deposits made to effect the agreed capital increase	Capital reserve	Accumulated losses	Shareholders' equity	
	Number of ordinary shares	Share capital				
As of December 31, 2008	9,378,348	9,378	0	24,745	-28,880	5,243
Capital increase in exchange for cash contributions			2,763	-152		2,611
Value of the services rendered by employees (acc. to IFRS 2)				48		48
Net loss for the period				-914		-914
As of March 31, 2009	9,378,348	9,378	2,763	24,641	-29,794	6,988
As of December 31, 2009	10,143,348	10,143	3,574	28,798	-35,164	7,351
Capital increase in exchange for cash contributions	512,000	512	-3,574	2,941		-121
Value of the services rendered by employees (acc. to IFRS 2)				50		50
Net loss for the period				-1,562		-1,562
As of March 31, 2010	10,655,348	10,655	0	31,789	-36,726	5,718

Mologen AG, Berlin, Germany

Condensed notes for the period of January 1 to March 31, 2010

A. General information on the company

Mologen AG (short: MOLOGEN) is a corporation headquartered in Berlin (Fabeckstraße 30, 14195 Berlin, Germany). It was founded on January 14, 1998 and is registered at the Berlin-Charlottenburg District Court under HRB65633. The shares of the company are listed on the regulated market (Prime Standard) of the Frankfurt Stock Exchange under ISIN DE0006637200.

The objective of the company is the research and development and the marketing of products in the field of molecular medicine. This particularly encompasses biomolecular vaccines, application-specific clinical research in the field of biomolecular tumor therapy, including somatic gene therapy. The main focus of research is on the MIDGE®- und dSLIM® technologies patented by MOLOGEN, which facilitate the use of DNA-based therapies to treat diseases that are currently untreatable or for which treatment is not sufficiently available.

B. General information on the financial statements

The present, unaudited condensed interim financial statements of MOLOGEN were prepared according to the IFRS as applied in the EU on the reporting date of March 31, 2010. They were prepared in accordance with IAS 34 (Interim Financial Reporting) and should be read in conjunction with MOLOGEN's audited annual financial statements of December 31, 2009 prepared according to the IFRS, as applied in the EU. The accounting and valuation methods have remained unchanged in comparison to the financial statements of December 31, 2009.

All new or adjusted accounting methods that were applied for the first time in the reporting period had no significant impact on the interim financial statements of MOLOGEN.

The reporting period of this condensed interim report is the period from January 1, 2010 to March 31, 2010. The comparison period for the comprehensive income statement as part of the condensed interim financial statements is the period from January 1, 2009 to March 31, 2009.

The functional currency and the presentation currency in the financial statements is the Euro (€). For better readability, the numbers have been rounded in accordance with standard business practice and are presented in thousands of EUR (EUR '000), unless otherwise indicated.

C. Selected notes to the balance sheet of March 31, 2010**Assets**Tangible assets / intangible assets

In the reporting period, tangible assets were purchased in the amount of EUR 5 thousand (12/31/2009: EUR 85 thousand); there were no significant disposals of tangible assets. There were no purchases or sales of intangible assets. There were no indications of any need to make unplanned impairment adjustments.

Cash and cash equivalents

Cash and cash equivalents comprise cash on hand and bank balances. Readily available bank balances are subject to variable interest rates. Short-term investments are undertaken for various periods for up to three months, based on the respective liquidity needs of the company. They are invested at fixed interest rates. The value of cash, cash equivalents and short-term investments as of the reporting date was EUR 4,518 thousand (12/31/2009: EUR 6,174 thousand). This is based on the nominal value of the Euro holdings and the valuation of a foreign currency account as of the reporting date on March 31, 2010.

Other current assets and income tax receivable

	03/31/2010 (EUR '000)	12/31/2009 (EUR '000)
Reimbursements from VAT	143	321
Accruals and deferrals	123	79
Income tax entitlements	20	20
Claims against tax authorities for investment subsidy	8	8
Other receivables	75	83
	369	511

No impairment charges were made with respect to other assets in the reporting period and the 2009 financial year.

Liabilities and shareholders' equityNon-current deferred liabilities

The amount stated as EUR 83 thousand (12/31/2009: EUR 86 thousand) comprises public grants for assets (EUR 12 thousand; 12/31/2009: EUR 13 thousand) and deferred income (EUR 71 thousand; 12/31/2009: EUR 73 thousand).

Current liabilities and deferred liabilities

	03/31/2010 (EUR '000)	12/31/2009 (EUR '000)
Deferred revenues	306	306
Trade payables	302	513
Liabilities from income and church tax	37	28
Payments received for orders	37	37
Liabilities to banks	7	5
Other liabilities	182	256
	871	1,145

The deferred liability position in the amount of EUR 306 thousand (12/31/2009: EUR 306 thousand) primarily stems from funds from the 7th Framework Programme of the EU in the amount of EUR 299 thousand (12/31/2009: EUR 299 thousand).

Shareholders' equity

A breakdown of the shareholders' equity and the development of the individual equity positions can be found in the statement of changes in shareholders' equity.

Issued capital

The issued capital of MOLOGEN is its share capital of EUR 10,655,348.00, divided into 10,655,348 no-par bearer shares, each with a notional share of EUR 1.00 per share unit.

In the reporting period, MOLOGEN implemented the following measures relevant to the subscribed capital:

On January 20, 2010, a capital increase performed in December 2009 was entered into the Commercial Register of the company's jurisdiction in exchange for cash contributions to the exclusion of subscription rights. Based on an authorization by the Annual General Meeting as well as the approval of the Supervisory Board of MOLOGEN, 512,000 ordinary shares in the name of the bearer (about 5% of the share capital) were issued to institutional investors. At an issue price of EUR 7.00 per share, the company as of December 31, 2009 had received total capital in the amount of EUR 3,574 thousand and additional capital in the amount of EUR 10 thousand between the effective date and the entry in the Commercial Register.

Authorized capital

Following the utilization of part of the authorized capital through issuance of 512,000 new shares in exchange for cash deposits for the capital increase entered into the Capital Register in January 2010, the authorized capital now amounts to EUR 3,706,424.00.

Conditional capital

Information on the conditional capital can be found in the annual financial statements for the financial year 2009. There were no changes in the reporting period, as no new employee participation programs were introduced and no stock options were exercised.

Capital reserve

The costs in the amount of EUR 131 thousand (01/01/2009 - 03/31/2009: EUR 152 thousand) incurred during the reporting period in connection with the capital measures entered into the Commercial Register were taken into consideration and reported in the capital reserve in accordance with IAS 32.37.

The application of IFRS 2 (Share-based payment) resulted in an addition to capital reserves in the amount of EUR 50 thousand (01/01/2009 - 03/31/2009: EUR 48 thousand) in the reporting period.

	03/31/2010 (EUR '000)	12/31/2009 (EUR '000)
Capital reserve	31,010	27,938
Employee compensation in equity instruments	2,716	2,666
Costs of equity procurement	-1,937	-1,806
	<u>31,789</u>	<u>28,798</u>

D. Selected notes to the income statementOther operating income

Other operating income (03/31/2010: EUR 3 thousand; 03/31/2009: EUR 103 thousand) decreased due to the fact that in contrast to the comparison period no subsidies were received (03/31/2009: EUR 100 thousand).

Personnel expenses

The increase in personnel expenses (03/31/2010: EUR 614 thousand; 03/31/2009: EUR 429 thousand) is due to restructuring in relation to the new hires for the department of clinical development, one-time effects caused by bonus payments, and a moderate increase in salaries.

Other operating expenses

Other operating expenses (03/31/2010: EUR 583 thousand; 03/31/2009: EUR 314 thousand) increased in contrast to the comparison period. The increase is due to an increase in research expenses.

Research and Development (R&D)

The resources available to the company are largely used directly for research projects. The expenditures for this area are in the amount of EUR 1.2 million (01/01/2009-03/31/2009: EUR 0.7 million). No development costs were incurred in the reporting or the comparison period.

E. Notes on the cash flow statement

The cash flow statement shows how the cash and cash equivalents of MOLOGEN have changed through cash inflows and outflows during the reporting period. According to IAS 7, a distinction is made between cash flow from operating activities and from investment and financing activities.

Cash flow from operations contains cash interest inflow in the amount of EUR 13 thousand (01/01/2009-03/31/2009: EUR 12 thousand). No interest was paid in the reporting or the comparison period.

F. Notes on the employee participation programs

During the reporting period, no employee participation programs were introduced and no stock options were exercised.

G. Notes on the types and management of financial risks

Information with regard to risks in connection with the use of financial instruments and with regard to our financial risk management can be found in the Notes to the Annual Financial Statements 2009 section H. There is nothing to be added to the risks that are described there.

H. Information on affiliated personsDirectors' Dealings

During the reporting period, the following securities transactions of executive employees were reported to the company as required according to Section 15a WpHG (Wertpapierhandelsgesetz - Securities Trading Act):

Name Function	Date	Transaction	Number of shares	Price	Trading volume	Trading location
Hella Schlichting, Natural person in close relationship to a member of the Supervisory Board	03/05/2010	Sale	1,900	EUR 8.90	EUR 16,910.00	XETRA / FFM

I. Other information

Approval of the interim financial statements

The interim financial statements were approved on May 11, 2010 by the Board of Directors and released for publication.

Berlin, 11 May 2010

Mologen AG
Board of Directors

Dr. Matthias Schroff
Chief Executive Officer

Jörg Petraß
Chief Financial Officer

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