



Unlocking the potential of innovative medicines

FIRST QUARTER REPORT

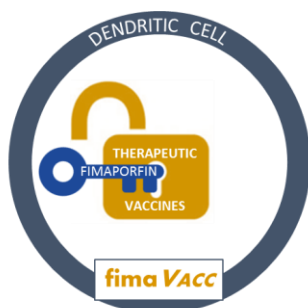
2018

LEVERAGING THE PCI-TECHNOLOGY IN THREE DISTINCT AREAS

TRIGGERED ENDOSOMAL RELEASE



Enabling approved drugs to fulfil unmet local treatment need



Enhancing cellular immune responses important for therapeutic vaccines



Providing a delivery solution for nucleic acid therapeutics

ABOUT PCI BIOTECH

PCI Biotech is a cancer focused biopharmaceutical company headquartered in Norway and listed on the Oslo Stock Exchange. The company develops therapeutic products based on its proprietary photochemical internalisation (PCI) technology. Originating from world leading research at the Oslo University Hospital - the Radium Hospital, the PCI technology works by inducing light-triggered endosomal release and may be used to unlock the true potential of a wide array of therapeutic modalities, such as small molecules, vaccines and nucleic acids.

PCI Biotech's lead candidate is the photosensitiser fimaporfin (Amphinex®). A Phase I study of fimaporfin in cancer patients has been completed at University College Hospital in London and published in Lancet Oncology. Promising early signs of tumour response were seen in all 22 patients and the treatment seemed to be well tolerated, providing the first clinical proof-of-concept of the fimaporfin technology.

HIGHLIGHTS

- **fimaCHEM**
 - Encouraging interim overall survival data from Phase I in bile duct cancer
 - Preparations for pivotal phase progressing towards initiation 2H'18
- **fimaVACC**
 - Promising interim clinical results from Phase I suggesting enhancement of several parameters of importance for vaccination
 - Completed dose-finding part of the Phase I study
- **fimaNAC**
 - Extension of the top-10 pharma collaboration
- **CORPORATE**
 - Oslo Børs listing, as a transfer from Oslo Axess (subsequent event)

KEY FIGURES

<i>(In NOK 1,000)</i>	2018 Q1	2017 Q1	2017 FY
Other income	2 238	2 428	10 250
Operating costs	16 900	12 281	53 681
Operating results	-14 663	-9 854	-43 431
Financial items	123	221	590
Comprehensive income	-14 540	-9 632	-42 841
Cash & cash equivalents	38 586	69 929	50 789
Net cash flow from operating activities	-12 203	-9 105	-29 943

The dose-escalation part of the fimaCHEM Phase I study in bile duct cancer continues to deliver encouraging survival data, with the latest update from March 2018 providing an interim average overall survival of 17.4 months, with 25% of the patients still being alive. The company is currently focusing on completion of discussions with advisors and contract research organisations on the full study design, as well as completion of the extension study that may allow for repeated fimaCHEM treatment in the pivotal study.

The initial results from the fimaVACC Phase I study in healthy volunteers reported last year indicate enhanced overall T-cell responses at tolerable dose levels, as well as early responses and high response rates. The dose-finding part of the study is completed with more than 90 subjects included, offering a large library of clinical fimaVACC test samples. The near term focus is now on in-depth analysis and characterisation of the immune responses.

Our alliances in the fimaNAC programme are progressing well and the research collaboration with a top-10 pharma company, which entered into a new stage (in vivo) in Q3 2017, has been further extended to the end of Q2 2018.

The listing of PCI Biotech was on 27th April transferred from Oslo Axess to Oslo Børs. This is seen as a natural, important development step, as the Company is approaching a pivotal clinical phase, and the market cap and shareholder base have developed positively over the last years. The listing on Oslo Børs is expected to enhance the visibility, strengthen the Company profile and make the shares more attractive to investors.

OPERATIONAL REVIEW

fimaCHEM

The fimaCHEM programme aims to fulfil unmet medical needs by providing local enhancement of approved chemotherapies. The lead project – local enhancement of gemcitabine in bile duct cancer – is in clinical development with Amphinex, the intravenous formulation of fimaporfin.

ENCOURAGING INTERIM OVERALL SURVIVAL DATA

Per March 2018 the interim average overall survival from Phase I in the study of fimaCHEM for treatment of inoperable extrahepatic bile duct cancer (cholangiocarcinoma) patients was 17.4 months, with 25% of the patients still being alive. The median overall survival ended at 14.4 months. The survival data includes all dose cohorts, 16 patients in total and are encouraging when seen in relation to the most appropriate published comparator data.

PREPARATIONS FOR PIVOTAL PHASE PROGRESSING TOWARDS INITIATION 2H'18

The promising early signs of efficacy in the Phase I study were based on a single fimaCHEM treatment in addition to standard of care treatment. A Phase I extension study was initiated with the objective to determine safety and tolerability of repeated treatments with fimaCHEM, as this may well increase the promising signs of efficacy even further. In this study, the second fimaCHEM treatment is in this study done approximately 3-4 months after the initial treatment. The extension study will include a minimum of 6 evaluable patients. The study is on schedule for readout of safety and tolerability of a minimum of 6 evaluable patients in second half of 2018. The fimaCHEM treatment in the pivotal study may include one or two treatments, depending on the outcome of the extension study.

Last year, the Company initiated processes to assess the fastest way to market for fimaCHEM in this life-threatening orphan disease without approved treatments. In December 2017 PCI Biotech announced the preliminary outcome of meetings with European Medicines Agency (EMA) and the U.S. Food and Drug Administration (FDA). The interactions with the regulatory authorities were initiated with the intent to determine the fastest route to market for fimaporfin, including the design of a pivotal study to support future marketing authorisation applications in EU and the USA. A common understanding was reached on several important factors for a pivotal study with fimaporfin in inoperable cholangiocarcinoma, including the sufficiency of a single randomised two-arm study and the potential for accelerated/conditional approval based on interim results. The randomised study will provide the opportunity to generate robust comparative data of importance for market acceptance of fimaporfin as a first line treatment. Further information on the study and the development strategy will be announced following completion of discussions with clinical advisors.

About bile duct cancer and the fimaCHEM technology

Bile duct cancer originates in the ducts that drain bile from the liver into the small intestine. It is a rare cancer without approved chemotherapies and the development pipeline is weak. The annual incidence rate is 1-2 cases per 100,000 in the Western world, but rates are higher in most Asian countries. The majority of cases present as inoperable and there is a high-unmet need for improved treatment technologies.

Surgery is currently the only curative option for these patients, yet the majority of the tumours are inoperable. Standard treatment for inoperable patients is stenting to keep the bile duct open, followed by chemotherapy. Combination of the chemotherapeutics gemcitabine and cisplatin has become standard treatment, but there is a need to increase overall survival and quality of life.

Bile duct cancer is characterised by a remarkable resistance to common chemotherapy, and there is a high need for new drug classes or alternative methods. The most studied and used drug is gemcitabine, which also is one of the drugs significantly enhanced by the fimaCHEM technology in preclinical studies. Light access for fimaCHEM treatment is easy through routinely used endoscopic methods.

About comparator data for inoperable bile duct cancer

The median overall survival (OS) in the studies that established gemcitabine and cisplatin as standard treatment in cholangiocarcinoma (CCA) was 11.7 and 11.2 months respectively (Valle et al. NEJM (2010) 362:1273-81 and Okusaka et al. BJC (2010) 103:469-74). Gallbladder cancer patients had a poorer outcome in the latter study and the median OS was 13 months when these patients were excluded. These results represents the best available published comparator data, but are not directly comparable to the data in the fimaCHEM Phase I study. The published studies include a wide range of different inoperable CCA patients, while the fimaCHEM Phase I study focuses on inoperable perihilar CCA patients.

fimaVACC

The **fimaVACC** programme aims to enhance the cellular immune responses important for therapeutic effect of vaccines. This proprietary vaccination technology has entered clinical development, and is also subject to one active research collaboration.

PROMISING INITIAL CLINICAL RESULTS

Improving immunogenicity of vaccine candidates is a main priority in the immunotherapy industry and a successful translation of the promising fimaVACC technology into man is therefore a very important step to establish PCI Biotech in the immunotherapy field.

The company initiated clinical validation of the promising fimaVACC technology in September 2016 through a Phase I, healthy volunteer study, in up to 170 healthy volunteers in the UK. The main objective of the study is to determine safety, tolerability and immune responses for fimaVACC.

The interim clinical results on overall T-cell responses indicate that vaccination with well-tolerated doses of fimaVACC enhance cellular immune responses important for therapeutic effect of vaccines. The data also suggest that fimaVACC triggers early T-cell responses and provides high response rates, which are two highly sought-after features of vaccination platforms. The best responses were seen at the lowest dose level tested and the study was therefore expanded with further dose cohorts investigating lower dose levels. The dose-finding part of the study is now completed, with more than 90 subjects included.

The study has provided a vast number of clinical test samples and the near-term focus is to further characterise the fimaVACC immune response by performing targeted in-depth immune analysis of relevant samples.

The fimaVACC programme is supported by a grant from the Research Council of Norway (BIA-programme) of up to NOK 13.8 million and the grant is distributed over the course of three and a half years, 2017-2020.

About immunotherapy with the fimaVACC technology

The pharmaceutical industry has long recognised the potential of therapeutic cancer vaccination, i.e. vaccines that treat cancer by inducing or strengthening an immune response. Several companies have reported failed clinical studies in the past years, but the potential of combining vaccination with checkpoint inhibitors has triggered a renewed interest in therapeutic cancer vaccines. However, there are still important unsolved issues and improving the immunogenicity of vaccine candidates is a main priority in immunotherapy. PCI Biotech believes the fimaVacc technology may play an important role in solving this challenge.

Effective induction of cytotoxic T-cells is key to realise the huge potential of therapeutic cancer vaccination, but vaccines often fail to generate such responses. One of the most important reasons is probably insufficient delivery of vaccine antigens to the appropriate presentation pathway in immune cells. The fimaVacc technology may solve this challenge by effectively enhancing the vaccine presentation through this pathway.

fimaNAC

The **fimaNAC** programme provides a targeted intracellular delivery technology for nucleic acid therapeutics. It is a preclinical stage opportunistic programme subject to four active research collaborations.

PRECLINICAL RESEARCH COLLABORATION

PCI Biotech has four active preclinical research collaborations in the area of nucleic acid therapeutics, established to evaluate technologically compatibilities and synergies and thereafter explore the potential for further partnerships.

A collaboration with an undisclosed top-10 pharma company, aiming to evaluate synergistic effects of fimaNAC with their nucleic acid therapeutics technology was signed in September 2015. This agreement has been extended several times, most recently until the end of June 2018 with the possibility for further extension. In 2017 the collaboration was expanded to include evaluation of technological compatibility and synergy based on *in vivo* studies, aimed at determining whether PCI Biotech's fimaNAC technology has the potential to enhance the therapeutic effect of the partner's nucleic acid therapeutics.

About the fimaNAC and nucleic acid therapy

The fimaNAC technology may enhance the delivery of most types of nucleic acids. Several forms of nucleic acids are widely acknowledged to have a large therapeutic potential, and numerous clinical trials are underway. The therapeutic potential of such compounds is challenged by the obstacles to achieve adequate intracellular access, which the fimaNAC technology may resolve.

The fimaNAC programme has four active research collaborations with key players in the field of nucleic acid therapeutics. These aim to explore synergies between partners proprietary nucleic acid technologies and the fimaNAC technology. The collaboration partners span from an undisclosed big pharma company to three mid-/small-size biotechs: BioNTech, eTheRNA immunotherapies and RXi Pharmaceuticals.

FINANCIAL REVIEW

Income Statement

(Figures in brackets = same period 2017 unless stated otherwise).

The Group did not record revenues for Q1 2018. Grants received from various public sources such as the Norwegian Research Council and "SkatteFUNN" were recorded as other income. Other income for Q1 amounted to NOK 2.2 million (NOK 2.4 million).

Expenditure on research activities is recognised as an expense in the period in which it was incurred. The Group has no development expenditure that qualifies for recognition as an asset under IAS 38 and all research expenses are recorded in the profit and loss statement, in line with previous years. Research and development (R&D) costs for Q1 totalled NOK 13.3 million (NOK 9.4 million). Operating costs ended at NOK 16.9 million (NOK 12.3 million). Operating expenses are mainly driven by the R&D activity level and for 2018 there are two clinical phase I trials running, increasing the cost level compared to 2017.

Net loss for the quarter was NOK 14.5 million (NOK 9.6 million).

Cash flow and balance sheet

The Group held cash and cash equivalents of NOK 38.6 million at the end of the quarter, compared to NOK 50.8 million at year-end 2017, reflecting net negative cash flow from operating activities. All cash and cash equivalents were placed as bank deposits at the end of the quarter.

Cash flow from operations is mainly dependent on R&D activities. Net cash flow from operating activities was NOK -12.2 million in the quarter (NOK -9.1 million). The increase in total short-term liabilities from NOK 14.6 million at year-end 2017 to NOK 16.6 million at the end of the quarter is mainly due to the higher R&D activity level

OTHER

Risks and uncertainty factors for 2018

PCI Biotech is exposed to uncertainties and risk factors, which may influence some or all of the company's activities. As described in the Annual Report 2017, the most important risks the company is exposed to in 2018 are associated with progress and performance of R&D programmes, and the associated regulatory affairs and market risk. No circumstances have been identified that significantly change the uncertainties and risk factors described in the Annual Report 2017.

Related party transactions

PCI Biotech is relying on services provided by third parties, including related parties, as a result of its organisational set-up. PCI Biotech considers its business relationship with The Norwegian Radium Hospital Research Foundation as the only material ordinary related party transactions in Q1 2018.

Post-closing events

One participant in the Company's share option program has on 12 April 2018 exercised a total number of 5,000 share options at a strike price of NOK 9.11 and a total number of 3,000 share options at a strike price of NOK 3.79, corresponding to a total number of 8,000 shares. At the same time another 4,000 share options lapsed.

Following the exercise of share options, the Company's Board of Directors, pursuant to an authorisation granted by the Company's Annual General Meeting on 29 May 2017, has decided to increase the Company's share capital with NOK 24,000 by issuing 8,000 new shares, each share of par value NOK 3.00. The capital increase was registered in the Norwegian Register of Business Enterprises 17 April 2018 and has thus been completed. Subsequent to the transaction, the Company's share capital is NOK 74,984,670 divided into 24,994,890 shares. The capital increase resulted in gross proceeds of NOK 56,920. Please see note 12 for further details.

On 27 April 2018 PCI Biotech was listed at Oslo Børs, as a transfer from Oslo Axess.

PCI Biotech is not aware of any other post-closing events, which could materially influence this interim financial statement.

OUTLOOK

PCI Biotech's lead project is clinical development of *fimaCHEM* (fimaporfin) in combination with gemcitabine for treatment of inoperable bile duct cancer; an orphan disease with high unmet medical need. Based on the promising early signs of efficacy in Phase I, the company has reached out to key regulators and received important guidance for a pivotal phase study. The final pivotal study design requires further discussions with clinical advisors and the development strategy will be announced following completion of these discussions.

PCI Biotech believes the PCI technology has potential to also play a role in the realisation of several new therapeutic modalities, including cancer immunotherapy (*fimaVACC*) and nucleic acid therapeutics (*fimaNAc*). The active research collaborations show that external companies share this view.

Clinical validation of the promising *fimaVACC* technology is essential for PCI Biotech's role within the immunotherapy space and the Phase I study in healthy volunteers will provide results on clinical translation of the technology. Initial results are promising and the study is expected to provide important results for determination of the development strategy.

The *fimaNAc* programme will continue to follow an opportunistic approach, pursuing out-licensing opportunities.

The main priorities of PCI Biotech are to:

- Effectively drive the *fimaCHEM* development programme in inoperable bile duct cancer towards the market;
- Progress and finalise the *fimaVACC* Phase I study in healthy volunteers;
- Alliance management and partnering activities across all commercially interesting areas for the PCI platform.

The Board of Directors and CEO
PCI Biotech Holding ASA
Oslo, 7 May 2018

Hans Peter Bøhn
Chairman (sign)

Christina Herder
Director (sign)

Hilde H. Steineger
Director (sign)

Kjetil Taskén
Director (sign)

Lars Viksmoen
Director (sign)

Per Walday
CEO (sign)

CONDENSED INTERIM CONSOLIDATED FINANCIAL INFORMATION

PROFIT AND LOSS

<i>(In NOK 1,000)</i>	Note	2018 Q1	2017 Q1	2017 FY
Other income	5	2 238	2 428	10 250
Research and development	8	13 335	9 378	40 988
General and administrative		3 565	2 903	12 693
Operating costs		16 900	12 281	53 681
Operating results		-14 663	-9 854	-43 431
Financial income and costs				
Financial income		123	221	677
Financial expenses		0	0	87
Net financial result		123	221	590
Profit/loss before income tax		-14 540	-9 632	-42 841
Income tax	9	0		0
Net profit/loss	4	-14 540	-9 632	-42 841
Other comprehensive income		0		0
Comprehensive income		-14 540	-9 632	-42 841

BALANCE SHEET

<i>(In NOK 1,000)</i>	Note	2018 31.03	2017 31.03	2017 31.12
Fixed and intangible assets				
Operating assets		21	4	22
Total fixed and intangible assets		21	4	22
Current assets				
Short term receivables	7	8 838	9 117	7 625
Cash & cash equivalents	7	38 586	69 929	50 789
Total current assets		47 424	79 045	58 414
Total assets		47 445	79 049	58 436
Shareholders' equity and liabilities				
Shareholders' equity				
Paid in capital		232 109	230 411	232 109
Other reserves		-203 551	-161 458	-190 266
Total equity	10	28 558	68 953	41 842
Other long term liabilities		2 260	0	2 009
Total long term liabilities		2 260	0	2 009
Trade debtors		2 109	842	1 497
Other short term liabilities		14 519	9 254	13 088
Total short term liabilities		16 627	10 096	14 585
Total liabilities		18 887	10 096	16 594
Total shareholders' equity and liabilities		47 445	79 049	58 436

CHANGE IN SHAREHOLDERS EQUITY

<i>(In NOK '000)</i>	2018 Q1	2017 Q1	2017 FY
Equity at beginning of period	41 842	13 086	13 086
Capital increase	-	65 032	66 729
Share option scheme	1 256	467	4 867
Comprehensive income in the period	-14 540	-9 632	-42 841
Equity at end of period	28 558	68 953	41 842

CASH FLOW

<i>(In NOK '000)</i>	2018 Q1	2017 Q1	2017 FY
Ordinary profit before taxes	-14 540	-9 632	-42 841
Depreciation, amortisation and write off	1	1	6
Share-based payments	1 256	467	4 867
Net financials	-123	-221	-677
Changes in working capital	1 080	59	8 025
Cash flow from operating activities	-12 326	-9 327	-30 620
Net financials	123	221	677
Taxes paid	-	-	-
Net cash flow from operating activities	-12 203	-9 105	-29 943
Cash flow from financial activities			
Net proceeds from share issues	-	65 032	66 730
Net cash flow from financial activities	-	65 032	66 730
Net change in cash during the period	-12 203	55 927	36 787
Cash and cash equivalents at the beginning of the period	50 789	14 002	14 002
Cash and cash equivalents at the end of the period	38 586	69 929	50 789

SELECTED EXPLANATORY NOTES:

1. Nature of operation

PCI Biotech Holding ASA (PCI Biotech) was established in 2008, and comprises PCI Biotech Holding ASA, the fully owned subsidiary PCI Biotech AS and the dormant Icelandic Branch PCI Biotech Utibu. PCI Biotech AS was a subsidiary of Photocure ASA until June 2008. The PCI Biotech shares have been listed on the Oslo Axess since 18 June 2008 under the ticker PCIB. On 27 April 2018 the listing was transferred from Oslo Axess to Oslo Børs. The company is headquartered in Oslo, Norway.

PCI Biotech has developed a unique and patented photochemical intracellular drug delivery technology for use in cancer therapy and other diseases. The technology may also be used to enhance the immunological response of vaccines. The company collaborates closely with The Norwegian Radium Hospital in Oslo, Norway and receives substantial funding on several projects from the Research Council of Norway. The company has an extensive international collaboration network with recognised expert groups in both drug delivery and vaccination. Photochemical Internalisation (PCI) is a proprietary technology for light-directed intracellular drug delivery by triggered endosomal release.

The PCI technology has potential to improve the efficacy of both existing drugs and new classes of drugs, such as therapeutic vaccines, gene therapy and other therapies based on nanotechnology or on biotechnological principles. The company's objective is to prove the clinical usefulness of the technology with various drugs and subsequently license out the technology to partners for further development and marketing. Revenues will be generated at the time of partnering and onwards from up-front payments, milestone payments and royalties from sales. PCI Biotech works on the development of PCI products for enhanced delivery of existing cancer drugs (fimaCHEM), and as a platform that may both potentiate the effect of vaccines (fimaVACC) and delivery of nucleic acids (fimaNAC). PCI Biotech has one active clinical development project in the fimaCHEM programme, with the lead candidate fimaporfin (Amphinex) in combination with the chemotherapeutic agent gemcitabine for treatment of bile duct cancer. The company also has one active study in the fimaVACC programme, a phase I study in healthy volunteers, for clinical proof of concept of fimaVACC's ability to enhance and direct the response of vaccines towards a stronger cellular type immunity. The fimaNAC programme is in preclinical stage.

2. Basis of presentation

These condensed interim financial statements have been prepared in accordance with IAS 34 Interim Financial Reporting. These condensed interim financial statements should be read in conjunction with the consolidated financial statements for the year ended 31 December 2017 (hereafter 'the Annual Financial Statements'), as they provide an update of previously reported information. The accounting policies used are consistent with those used in the Annual Financial Statements. The presentation of the condensed interim financial statements is consistent with the Annual Financial Statements. This interim report has not been subject to an audit. The going concern assumption has been applied when preparing this interim financial report. The board of directors approved the condensed interim financial information on 7 May 2018.

PCI Biotech has Norwegian kroner (NOK) as its functional currency and presentation currency. In the absence of any statement to the contrary, all financial information is reported in whole thousands. As a result of rounding adjustments, the figures in the condensed interim financial statements may not add up to the totals.

3. Summary of significant accounting policies

The accounting policies applied and the presentation of the interim condensed consolidated financial information is consistent with the consolidated financial statements for the year ended 31 December 2017.

The new standards and interpretations or amendments to published standards that were effective for the annual period beginning on January 1, 2018 or later and that could affect PCI Biotech are discussed in accounting principles, part 4, to the consolidated financial statements for 2017. In the 2017 financial statements, PCI Biotech made evaluations that at current stage *IFRS 9 Financial Instruments*, *IFRS 15 Revenue from contract with customers* and *IFRS 16 Leases* are not expected to have a significant impact on PCI Biotech's financial position, performance and/or disclosure.

4. Important accounting valuations, estimates and assumptions

Estimates and judgments are evaluated on an on-going basis and are based on historical experience and other factors, including expectations of future events that are considered to be relevant.

In preparing these condensed interim financial statements, the significant judgements made by management in applying the group's accounting policies and the key sources of estimation uncertainty were the same as those applied to the consolidated financial statements for the year ended December 31st, 2017.

5. Earnings per share

Earnings per share

	2018 Q1	2017 Q1	2017 FY
Result allocated to shareholders (NOK'000)	-14 540	-9 632	-42 841
Weighted average of outstanding shares ('000)	24 988	22 593	24 348
Earnings per share (NOK per share)	-0.58	-0.43	-1.76

Diluted earnings per share:

	2018 Q1	2017 Q1	2017 FY
Result allocated to shareholders (NOK'000)	-14 540	-9 632	-42 841
Weighted average of outstanding shares ('000)	25 727	22 968	24 826
Earnings per share (NOK per share)	-0.58	-0.43	-1.76

Weighted average of outstanding diluted shares is weighted number of average number of shares adjusted with share options that are in the money. Earnings per share is not affected by the dilution if negative results in the period.

6. Segment information

The Company reports only one segment and had no revenues for the reporting period. Government grants are recognised at the value of the contribution at the transaction date. Grants are not recognised until it is probable that the conditions attached to the contribution will be achieved. The grants are recognised in the statement of profit and loss in the same period as the related costs, and are disclosed as other income. The Company has recognised Norwegian grants and tax incentive scheme (SkatteFUNN) in the period.

7. Related party transactions

PCI Biotech is relying on services provided by third parties, included related parties, as a result of its organisational set-up. PCI Biotech considers that its business relationship with The Norwegian Radium Hospital Research Foundation regarding research and overall PCI technology development represent

related party transactions. The following table shows the extent of such transactions in the reported periods (all figures in NOK '000):

Purchase of services	2018 Q1	2017 Q1	2017 FY
The Norwegian Radium Hospital Research Foundation	459	870	2 600

At the end of the quarter PCI Biotech had NOK 0.3 million in short-term liability to The Norwegian Radium Hospital Research Foundation.

8. Credit risk, foreign currency risk and interest risk

Credit risk

PCI Biotech has no sales for 2017 and 2018 and faces therefore no credit risk.

Maturity profile on short-term receivables at the end of the quarter (all figures in '000 NOK):

	Not due (prepaid expenses)	Less than 3 months	3 to 12 months	More than 12 months	Total
Trade receivables	-	-	-	-	-
Other receivables	348	1 523	5 717	1 250	8 838
Total receivables	348	1 523	5 717	1 250	8 838

A majority of the short-term receivables relates to accrued, not received grants (BIA) and tax incentive scheme (SkatteFUNN).

Foreign currency risk

PCI Biotech has transactional currency exposure arising from purchases in currencies other than the functional currency (NOK). PCI Biotech has not implemented any hedging strategy to reduce foreign currency risk.

Interest risk

PCI Biotech has no interest bearing debt.

9. Research and Development costs

All figures in '000 NOK

	Q1 2018	Q1 2017	FY 2017
Clinical studies	9 993	5 465	23 886
Pre-clinical studies	1 531	2 869	12 539
CMC and equipment	1 239	405	1 770
Patents	572	639	2 793
Other costs	0	0	0
Total	13 335	9 378	40 988

PCI Biotech has no development expenditure that qualifies for recognition of an asset under IAS 38 Intangible assets and all research expenditures are charged through the income statement, in line with previous years.

PCI Biotech reviewed in 2017 the internal allocation of operating expenses for disclosure of the sub categories in the statement of comprehensive income; research and development expenses versus general and administrative expenses. The review was made based on the current operational set-up of the organisation which has changed and developed over the years, from an early stage clinical company towards a pivotal stage ready company. The outcome of the review has led to reallocation of expenses, for the interim figures previously reported for 2017, between these two relevant P&L sub categories with no net change in the disclosed total operating expenses. In the statement of comprehensive income 2017 for the Company the new allocation routines are applied prospectively, as this reflects the underlying operations.

10. Deferred tax and deferred tax assets

At the end of the quarter, the group held NOK 81.3 million in non-capitalised deferred tax assets, which mainly relates to carry forward losses.

11. Share options

Share options outstanding at the end of the period have the following expiry date and exercise prices:

Expiry date	Exercise price in NOK per share	Number of shares	
		31.12.2017	31.03.2018
2018 - Q3	10.55	85 000	85 000
2018 - Q3	10.02	40 000	40 000
2020 - Q3	9.11	73 500	73 500
2020 - Q3	3.79	110 000	110 000
2022 - Q3	24.95	340 000	340 000
2022 - Q3	22.35	90 000	90 000
Total		738 500	738 500

As a subsequent event, one participant in the Company's share option program has on 12 April 2018 exercised a total number of 5,000 share options at a strike price of NOK 9.11 and a total number of 3,000 share options at a strike price of NOK 3.79, corresponding to a total number of 8,000 shares. At the same time another 4,000 share options lapsed.

Overview options 2018, Senior executives	Total holdings 31.12.2017	Allocated	Lapsed	Exercised	Expired	Total holdings 31.03.2018
Per Walday, CEO	104 000	0	0	0	0	104 000
Ronny Skuggedal, CFO	116 000	0	0	0	0	116 000
Anders Høgset, CSO	66 000	0	0	0	0	66 000
Gaël L'Hévéder, CBDO	106 000	0	0	0	0	106 000
Kristin Eivindvik, PD	33 500	0	0	0	0	33 500
Hans Olivecrona, CMO	90 000	0	0	0	0	90 000
Sum	515 500	0	0	0	0	515 500

12. Share capital

The Company's share capital is NOK 74,960,670 divided into 24,986,890 shares, each giving one vote at the Company's general meeting.

	No. of shares	Nominal value per share in NOK	Share capital in NOK
31.12.2017	24 986 890	3.00	74 960 670
Rights Issue			
Exercise of share options			
31.03.2018	24 986 890	3.00	74 960 670

Following the exercise of share options on 12 April 2018, the Company's Board of Directors, pursuant to an authorisation granted by the Company's Annual General Meeting on 29 May 2017, decided to increase the Company's share capital with NOK 24,000 by issuing 8,000 new shares, each share of par value NOK 3.00. Subsequent to the transaction, the Company's share capital will be NOK 74,984,670 divided into 24,994,890 shares after the subsequent event. The capital increase results in gross proceeds of NOK 56,920.

The Annual General Meeting held 29 May 2017 authorised the Board of Directors to execute share capital increases by issuing up to 1,865,000 shares with a nominal value of NOK 3 in connection with the company's employee incentive program. The authorisation is valid for 2 years. The Board of Directors have utilised the authorisation to issue 86,500 shares by end of September 2017 and 8,000 shares in April 2018 as a subsequent event.

The Annual General Meeting held 29 May 2017 authorised the Board of Directors to execute share capital increases with up to NOK 8,029,600 in connection with private placements. The authorisation shall not be used to increase the share capital by an amount in excess of 10% of the share capital, based on the share capital per 29 May 2017 and potential share capital increases in relation to the employee incentive programme. The authorisation may be used for general corporate purposes. The authorisation is valid for 2 years.

The Company has more than 3,500 shareholders at the end of the quarter.

10 largest shareholders per 31 March 2018:

Name	No. of shares	Ownership
FONDSAVANSE AS	2 540 840	10,17 %
MP PENSJON PK	1 394 522	5,58 %
RADIUMHOSPITALET'S FORSKNINGSSSTIFTELSE	1 261 262	5,05 %
Myrlid AS	1 240 000	4,96 %
NORDNET LIVSFORSIKRING	663 870	2,66 %
GRESSLIEN ODD ROAR	530 000	2,12 %
BERG-LARSEN ALEXANDER	480 181	1,92 %
Nordnet Bank AB	467 233	1,87 %
SYVERTSEN SVEIN ERIK	393 907	1,58 %
Jandersen Kapital AS	360 000	1,44 %
Total 10 largest shareholders	9 331 815	37,35 %
<i>Others</i>	15 655 075	62,65 %
<i>Total</i>	24 986 890	100 %

Shares owned, directly or indirectly, by members of the board, senior executives and their personally related parties per end of the quarter:

Name	Position	No. of shares	
		31.03.2018	31.12.2017
Hans Peter Bøhn	Chairman	83 556	83 556
Christina Herder	Board member	8 355	8 355
Kjetil Taskén (Kjetil Taskén AS)	Board member	4 000	4 000
Lars Viksmoen (Stocken Invest AS)	Board member	4 000	4 000
Hilde H. Steineger	Board member	0	0
Per Walday	CEO	65 133	65 133
Anders Høgset	CSO	62 456	62 456
Ronny Skuggedal	CFO	25 066	25 066
Gaël L'Hévéder	CBDO	10 000	10 000
Kristin Eivindvik	PD	17 948	17 948
Total		280 514	280 514

13. Other short term liabilities

Other short term liabilities mainly consist of accrued R&D and salary related costs and public duties.

14. Subsequent events

One participant in the Company's share option program has on 12 April 2018 exercised a total number of 5,000 share options at a strike price of NOK 9.11 and a total number of 3,000 share options at a strike price of NOK 3.79, corresponding to a total number of 8,000 shares. At the same time another 4,000 share options lapsed.

Following the exercise of share options on 12 April 2018, the Company's Board of Directors, pursuant to an authorisation granted by the Company's Annual General Meeting on 29 May 2017, has decided to increase the Company's share capital with NOK 24,000 by issuing 8,000 new shares, each share of par value NOK 3.00. Subsequent to the transaction, the Company's share capital will be NOK

74,984,670 divided into 24,994,890 shares. The capital increase results in gross proceeds of NOK 56,920.

On 27 April 2018 PCI Biotech was listed at Oslo Børs, as a transfer from Oslo Axess.

PCI Biotech is not aware of any other post-closing events, which could materially influence this interim financial statement.

DEFINITIONS AND GLOSSARY

Amphinex:	Trade name of the clinical intravenous formulation of fimaporfin
FDA:	US Food and Drug Administration
Fimaporfin:	Generic name of the photosensitiser active ingredient TPCS2a
IND	Investigational New Drug
<i>In vitro</i> :	Studies performed with cells or biological molecules studied outside their normal biological context; for example proteins are examined in solution, or cells in artificial culture medium.
<i>In vivo</i> :	Studies in which the effects of various biological entities are tested on whole, living organisms usually animals.
ODD:	Orphan Drug Designation
PCI:	Photochemical internalisation
PFS:	Progression Free Survival
R&D:	Research and Development
FY:	Financial year (1 st January – 31 st December)
NOK:	Norwegian kroner
Q1:	First quarter (1 st January – 31 st March)

FINANCIAL CALENDAR

Q2 Report 2018	30 August 2018
Q3 Report 2018	13 November 2018

INVESTOR CONTACT

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FORWARD LOOKING STATEMENTS

This Report contains certain forward-looking statements relating to the business, financial performance and results of the Company and/or the industry in which it operates. Forward-looking statements concern future circumstances and results and other statements that are not historical facts, and are sometimes identified by the words “believes”, “expects”, “predicts”, “intends”, “projects”, “plans”, “estimates”, “aims”, “foresees”, “anticipates”, “targets”, and similar expressions. The forward-looking statements contained in this Report, including assumptions, opinions and views of the Company or cited from third party sources, are solely opinions and forecasts which are subject to risks, uncertainties and other factors that may cause the actual results, performance or achievements of the Company to be materially different from any future results, performance or achievements that are expressed or implied by statements and information in the Report, including, among others, risks or uncertainties associated with the Company’s business, segments, development, growth management, financing, market acceptance and relations with customers, and, more generally, general economic and business conditions, changes in domestic and foreign laws and regulations, taxes, changes in competition and pricing environments, and fluctuations in currency exchange rates and interest rates. None of the Company or any of its subsidiaries or any such person’s directors, employees or advisors provide any assurance that the assumptions underlying forward-looking statements expressed in this Report are free from errors nor does any of them accept any responsibility for the future accuracy of such forward-looking statements.

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