UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549

FORM 6-K

Report of Foreign Private Issuer
Pursuant to Rule 13a-16 or 15d-16 of
the Securities Exchange Act of 1934

August 3, 2023

Commission File Number: 001-36622

PROQR THERAPEUTICS N.V.

Zernikedreef 9
2333 CK Leiden
The Netherlands
Tel: +31 88 166 7000
(Address, Including ZIP Code, and Telephone Number,
Including Area Code, of Registrant's Principal Executive Offices)

Indicate by check mark whether the registrant files or will file annual reports under cover of Form 20-F or Form 40-F.

Form 20-F ☒ Form 40-F ☐

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(1): ☐

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(7): ☐

Furnished as Exhibit 99.1 to this Report on Form 6-K are the unaudited financial statements of ProQR Therapeutics N.V. (the “Company”) for the three and six month periods ended June 30, 2023, and furnished as Exhibit 99.2 to this Report on Form 6-K is a press release of ProQR Therapeutics N.V. dated August 3, 2023, announcing the Company’s results for the three and six month periods ended June 30, 2023.

On August 3, 2023, the Company issued a press release titled, “ProQR Announces Second Quarter 2023 Operating and Financial Results,” announcing the Company’s results for the three and six month periods ended June 30, 2023 and providing a business update. A copy of this press release is attached hereto as Exhibit 99.2 and is incorporated herein by reference.

INDEX TO EXHIBITS

Number	Description
99.1	Unaudited financial statements of ProQR Therapeutics N.V. for the three and six month periods ended June 30, 2023.
99.2	Press Release of ProQR Therapeutics N.V. dated August 3, 2023, announcing the Company's results for the three and six month periods ended June 30, 2023.

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned, thereunto duly authorized.

PROQR THERAPEUTICS N.V.

Date: August 3, 2023

By: /s/ Jurriaan Dekkers
Jurriaan Dekkers
Chief Financial Officer

Table of Contents

Exhibit 99.1

PROQR THERAPEUTICS N.V.
Index to Unaudited Condensed Consolidated Financial Statements

PROQR THERAPEUTICS N.V.
Unaudited Condensed Consolidated Statement of Financial Position

		June 30, 2023	December 31, 2022
		€1,000	*€1,000*
Assets			
Current assets			
Cash and cash equivalents	5	128,562	94,775
Prepayments and other receivables		2,187	59,078
Other taxes		400	607
Total current assets		**131,149**	**154,460**
Property, plant and equipment	6	16,220	16,240
Investments in financial assets	16	621	621
Total assets		**147,990**	**171,321**
Equity and liabilities			
Equity			
Equity attributable to owners of the Company		51,790	67,064
Non-controlling interests		(290)	(384)
Total equity	**11**	**51,500**	**66,680**
Current liabilities			
Borrowings	8	2,920	2,500
Lease liabilities	9	1,334	1,387
Derivative financial instruments	8	372	1,263
Trade payables		119	392
Social securities and other taxes		1,064	1,118
Deferred income	10	13,137	5,641
Other current liabilities	7	4,030	8,687
Total current liabilities		**22,976**	**20,988**
Borrowings	8	3,575	4,271
Lease liabilities	9	13,700	13,813
Deferred income	10	56,239	65,569
Total liabilities		**96,490**	**104,641**
Total equity and liabilities		**147,990**	**171,321**

The accompanying notes are an integral part of these unaudited interim condensed consolidated financial statements.

PROQR THERAPEUTICS N.V.
Unaudited Condensed Consolidated Statement of Profit or Loss and OCI
(€ in thousands, except share and per share data)

		Three month period ended June 30,		Six month period ended June 30,	
		2023	2022	2023	2022
		€1,000	*€1,000*	*€1,000*	*€1,000*
Revenue	12	**1,205**	**930**	**1,860**	**2,060**
Other income	13	**38**	**99**	**80**	**200**

Table of Contents

Research and development costs	14	(5,909)	(11,449)	(11,969)	(24,816)
General and administrative costs	15	(4,145)	(5,412)	(8,171)	(10,320)
Total operating costs		**(10,054)**	**(16,861)**	**(20,140)**	**(35,136)**
Operating result		**(8,811)**	**(15,832)**	**(18,200)**	**(32,876)**
Finance income and expense		470	1,163	(74)	342
Results related to associates		—	—	—	(8)
Results related to financial liabilities measured at fair value through profit or loss	8	221	62	891	3,826
Results on derecognition of financial liabilities	17	101	1,144	509	1,144
Result before corporate income taxes		**(8,019)**	**(13,463)**	**(16,874)**	**(27,572)**
Income taxes	18	42	(20)	42	(27)
Result for the period		**(7,977)**	**(13,483)**	**(16,832)**	**(27,599)**
Other comprehensive income (foreign exchange differences on foreign operation)		7	689	(212)	911
Total comprehensive income		**(7,970)**	**(12,794)**	**(17,044)**	**(26,688)**
Result attributable to					
Owners of the Company		(7,993)	(13,700)	(16,926)	(27,808)
Non-controlling interests		16	217	94	209
		(7,977)	**(13,483)**	**(16,832)**	**(27,599)**
Total comprehensive income attributable to					
Owners of the Company		(7,986)	(13,011)	(17,138)	(26,897)
Non-controlling interests		16	217	94	209
		(7,970)	**(12,794)**	**(17,044)**	**(26,688)**
Share information					
Weighted average number of shares outstanding[1]		80,939,392	71,362,088	80,913,751	71,359,642
Earnings per share attributable to owners of the Company (Euro per share)					
Basic loss per share[1]		(0.10)	(0.19)	(0.21)	(0.39)
Diluted loss per share[1]		(0.10)	(0.19)	(0.21)	(0.39)

The accompanying notes are an integral part of these unaudited interim condensed consolidated financial statements.

1. For these periods the potential exercise of share options is not included in the diluted earnings per share as the Company was loss-making. Due to the anti-dilutive nature of the outstanding options, basic and diluted earnings per share are equal.

PROQR THERAPEUTICS N.V.
Unaudited Condensed Consolidated Statement of Changes in Equity

	Attributable to owners of the Company									
	Number of shares	**Share Capital**	**Share Premium**	**Equity settled Employee Benefit Reserve**	**Option premium on convertible loan**	**Translation Reserve**	**Accumulated Deficit**	**Total**	**Non-controlling interests**	**Total Equity**
		€1,000	*€1,000*	*€1,000*	*€1,000*	*€1,000*	*€1,000*	*€1,000*	*€1,000*	*€1,000*
Balance at January 1, 2022	**74,865,381**	**2,995**	**398,309**	**28,443**	**1,426**	**430**	**(316,890)**	**114,713**	**(604)**	**114,109**
Result for the period	—	—	—	—	—	—	(27,808)	(27,808)	209	(27,599)
Other comprehensive income	—	—	—	—	—	911	—	911	—	911
Recognition of share-based payments	—	—	—	1,921	—	—	—	1,921	—	1,921
Treasury shares transferred	(71,283)	—	—	—	—	—	—	—	—	—
Share options lapsed	—	—	—	(380)	—	—	380	—	—	—
Share options exercised	71,283	—	33	(256)	—	—	256	33	—	33
Balance at June 30, 2022	**74,865,381**	**2,995**	**398,342**	**29,728**	**1,426**	**1,341**	**(344,062)**	**89,770**	**(395)**	**89,375**
Balance at January 1, 2023	**84,246,967**	**3,370**	**412,540**	**29,052**	**—**	**1,212**	**(379,110)**	**67,064**	**(384)**	**66,680**
Result for the period	—	—	—	—	—	—	(16,926)	(16,926)	94	(16,832)
Other comprehensive income	—	—	—	—	—	(212)	—	(212)	—	(212)
Recognition of share-based payments	—	—	—	1,860	—	—	—	1,860	—	1,860
Treasury shares transferred	(122,584)	—	—	—	—	—	—	—	—	—
Share options lapsed	—	—	—	(3,873)	—	—	3,873	—	—	—
Share options exercised / RSUs vested	122,584	—	4.00	(231)	—	—	231	4	—	4
Balance at June 30, 2023	**84,246,967**	**3,370**	**412,544**	**26,808**	**—**	**1,000**	**(391,932)**	**51,790**	**(290)**	**51,500**

The accompanying notes are an integral part of these unaudited interim condensed consolidated financial statements

PROQR THERAPEUTICS N.V.
Unaudited Condensed Consolidated Statement of Cash Flows

		Three month period ended June 30,		Six month period ended June 30,	
		2023	2022	2023	2022
		€1,000	*€1,000*	*€1,000*	*€1,000*
Cash flows from operating activities					
Net result		(7,977)	(13,483)	(16,832)	(27,599)
Adjustments for:					
— Depreciation		594	591	1,143	1,161
— Share-based compensation		765	738	1,860	1,921
— Financial income and expenses		(470)	(1,163)	74	(342)
— Results related to associates		—	—	—	8
— Results related to financial liabilities measured at fair value through profit or loss		(221)	(62)	(891)	(3,826)
— Results on derecognition of financial liabilities	17	(101)	(1,144)	(509)	(1,144)
— Income tax expenses	18	—	20	—	27
Changes in working capital		(3,622)	916	48,668	(3,035)
Cash (used in)/generated from operations		*(11,032)*	*(13,587)*	*33,513*	*(32,829)*
Corporate income tax paid		—	(20)	—	(27)
Interest received		685	—	865	—
Interest paid		—	(1,237)	—	(2,455)
Net cash (used in)/generated from operating activities		***(10,347)***	***(14,844)***	***34,378***	***(35,311)***
Cash flow from investing activities					
Purchases of property, plant and equipment		(294)	(231)	(430)	(475)
Sales of property, plant and equipment		—	—	47	—
Net cash used in investing activities		***(294)***	***(231)***	***(383)***	***(475)***
Cash flow from financing activities					
Proceeds from exercise of share options	11	4	—	4	33
Repayment of lease liability	9	(647)	(357)	(906)	(933)
Net cash used in financing activities		***(643)***	***(357)***	***(902)***	***(900)***
Net (decrease)/increase in cash and cash equivalents		**(11,284)**	**(15,432)**	**33,093**	**(36,686)**
Currency effect cash and cash equivalents		860	4,222	694	5,564
Cash and cash equivalents, at beginning of the period		138,986	167,612	94,775	187,524
Cash and cash equivalents at the end of the period		**128,562**	**156,402**	**128,562**	**156,402**

The accompanying notes are an integral part of these unaudited interim condensed consolidated financial statements.

PROQR THERAPEUTICS N.V.
Notes to Unaudited Condensed Consolidated Financial Statements

1. General information

ProQR Therapeutics N.V., or "ProQR" or the "Company", is a biotechnology company domiciled in the Netherlands that primarily focuses on the discovery and development of novel therapeutic medicines.

Since September 18, 2014, the Company's ordinary shares have been listed on Nasdaq. They are currently trading at Nasdaq Capital Market under ticker symbol PRQR.

The Company was incorporated in the Netherlands, on February 21, 2012 (Chamber of Commerce no. 54600790) and was reorganized from a private company with limited liability to a public company with limited liability on September 23, 2014. The Company has its statutory seat in Leiden, the Netherlands. The address of its headquarters and registered office is Zernikedreef 9, 2333 CK Leiden, the Netherlands.

ProQR Therapeutics N.V. is the ultimate parent company of the following entities:

- ProQR Therapeutics Holding B.V. (100%);
- ProQR Therapeutics I B.V. (100%);
- ProQR Therapeutics II B.V. (100%);
- ProQR Therapeutics III B.V. (100%);
- ProQR Therapeutics IV B.V. (100%);
- ProQR Therapeutics V B.V. (100%);
- ProQR Therapeutics VI B.V. (100%);
- ProQR Therapeutics VII B.V. (100%);
- ProQR Therapeutics VIII B.V. (100%);
- ProQR Therapeutics IX B.V. (100%);
- ProQR Therapeutics I Inc. (100%);
- Amylon Therapeutics B.V. (80%);

ProQR Therapeutics N.V. is also statutory director of Stichting Bewaarneming Aandelen ProQR ("ESOP Foundation") and has full control over this entity. The Company holds a 5.1% minority shareholding in Yarrow Biotechnology, Inc.

As used in these condensed consolidated financial statements, unless the context indicates otherwise, all references to "ProQR" or the "Company" refer to ProQR Therapeutics N.V. including its subsidiaries and the ESOP Foundation.

Revision of comparative figures

In the Company's application of IAS 21 *The Effects of Changes in Foreign Exchange Rates*, certain deferred income positions were incorrectly treated as monetary items in 2021 and 2022. To correct for the effects of this error, which is immaterial for all affected prior periods, the comparative figures for the year ended December 31, 2022 and the three and six month periods ended June 30, 2022 have been revised as follows:

- in the Statement of financial position as at December 31, 2022, equity attributable to owners of the Company increased by € 1,567,000 and total deferred income decreased by € 1,567,000.
- In the Statement of profit or loss and OCI for the three and six month periods ended June 30, 2022, revenue decreased by € 95,000 and € 199,000, respectively, and net finance expenses decreased by €1,282,000 and €1,720,000 respectively. Net loss for the three and six month periods ended June 30, 2022 decreased by €1,187,000 and €1,521,000, respectively.
- In the Statement of changes in equity, accumulated deficit at January 1, 2022 decreased by €880,000.
- In the Statement of cash flows for the three and six month periods ended June 30, 2022, changes in working capital decreased by €95,000 and €199,000, respectively. Net cash used in operating activities for the three and six month periods ended June 30, 2022 was not affected by the revision.

2. Significant Accounting Policies

These interim condensed consolidated financial statements for the three and six month periods ended June 30, 2023 have been prepared in accordance with IAS 34 Interim Financial Statements. They should be read in conjunction with the

Company's annual financial statements for the year ended December 31, 2022. These interim condensed consolidated financial statements do not include all information required for a complete set of financial statements prepared in accordance with IFRS Standards. However, selected explanatory notes are included to explain events and transactions that are significant to an understanding of the changes in the Company's financial position and performance since the last annual financial statements. In the opinion of management, all events and transactions that are significant to an understanding of the changes in financial position and performance of the Company since the end of the last annual reporting period are disclosed in these interim condensed consolidated financial statements. The accounting policies adopted in the preparation of the interim condensed consolidated financial statements are consistent with those applied in the preparation of the Company's annual financial statements for the year ended December 31, 2022.

The Company's financial results have varied substantially, and are expected to continue to vary, from period to period. The Company believes that its ordinary activities are not linked to any particular seasonal factors.

The management of ProQR has, upon preparing and finalizing these interim condensed consolidated financial statements, assessed the Company's ability to fund its operations for a period of at least one year after the date of signing these interim condensed consolidated financial statements. Management expects the Company to continue as a going concern based on its existing funding, taking into account the Company's current cash position and the projected cash flows based on the activities under execution on the basis of ProQR's business plan and budget. Based on our current operating plan, we believe that the existing cash and cash equivalents will be sufficient to fund our anticipated level of operations at least into mid-2026. Thus, we continue to adopt the going concern basis of accounting in preparing the interim condensed consolidated financial statements.

The carrying amount of all financial assets and financial liabilities is a reasonable approximation of the fair value and therefore information about the fair values of each class has not been disclosed.

The Company operates in one reportable segment, which comprises the discovery and development of innovative, RNA based therapeutics.

3. Adoption of new and revised International Financial Reporting Standards

New Standards and Interpretations, which became effective as of January 1, 2023, did not have a material impact on our condensed consolidated financial statements.

4. Critical Accounting Estimates and Judgments

In the application of the Company's accounting policies, management is required to make judgments, estimates and assumptions about the carrying amounts of assets and liabilities that are not readily apparent from other sources. The estimates and associated assumptions are based on historical experience and other factors that are considered to be relevant. Actual results may differ from these estimates.

The estimates and underlying assumptions are reviewed on an ongoing basis. Revisions to accounting estimates are recognized in the period in which the estimate is revised if the revision affects only that period or in the period of the revision and future periods if the revision affects both current and future periods.

The significant judgements made by management in applying the Company's accounting policies and the key sources of estimation uncertainty were the same as those described in the Company's annual financial statements for the year ended December 31, 2022.

Revenue recognition for the Eli Lilly collaboration and license agreement

a. Identification of the performance obligation

Note 12 describes the Company's original research and collaboration agreement with Eli Lilly and Company, and the amended and restated research and collaboration agreement (collectively, the "Collaboration agreement"). Under the Collaboration agreement, ProQR provides Eli Lilly with a license (with a right to sub-license) to exploit compounds resulting from the collaboration. A significant amount of judgement is required to determine whether the license is distinct from the other promises in the contract. The license was concluded not to be distinct from the other promises in the contract based on the following considerations:

- the license has no stand-alone value to Eli Lilly without the Company being involved in the research and development collaboration, and;
- there are significant interdependencies between the license and the research and development services to be provided by the Company.

b. Determining the timing of satisfaction of performance obligations

Under the Collaboration agreement, the Company recognizes revenue over time, using an input method that estimates the satisfaction of the performance obligation as the percentage of labor hours incurred compared to the total estimated labor hours required to complete the promised services. As our estimate of the total labor hours required is dependent on the evolution of the research and development activities, it may be subject to change. If the progression and/or outcome of certain research and development activities would be different from the assumptions that were made during the preparation of these financial statements, this could lead to material adjustments to the total estimated labor hours, which might result in a reallocation of revenue between current and future periods. Our total deferred revenue balance related to this Eli Lilly performance obligation amounts to € 69,376,000 at June 30, 2023 (December 31, 2022: € 71,210,000).

c. Determining the transaction price

The Company applied judgement to determine whether the equity investments made by Eli Lilly in ProQR are part of the transaction price for the Collaboration agreement. The Company concluded that the differences between the prices that Eli Lilly paid for the shares and the ProQR stock closing prices on the days of entering into the equity investment agreements arose because of the Company's existing obligations to deliver research and development services to Eli Lilly under the terms of the Collaboration agreement. Therefore, the above differences between the closing share prices on the agreement effective dates and the equity investment prices paid by Eli Lilly are considered to be part of the transaction price of the contract and are initially allocated to deferred revenue.

The contract also includes variable consideration, but no variable consideration was included in the transaction price, as it is not highly probable that a significant reversal in the amount of cumulative revenue recognized will not occur when the uncertainty associated with the variable consideration is subsequently resolved.

Research and development expenditures
Research expenditures are reflected in the income statement. Development expenses are currently also reflected in the income statement because the criteria for capitalization are not met. At each balance sheet date, the Company estimates the level of service performed by the vendors and the associated costs incurred for the services performed.

Although we do not expect the estimates to be materially different from amounts actually incurred, the understanding of the status and timing of services performed relative to the actual status and timing of services performed may vary and could result in reporting amounts that are too high or too low in any particular period.

5. Cash and cash equivalents

At June 30, 2023, the Company's cash and cash equivalents were € 128,562,000 as compared to € 94,775,000 at December 31, 2022. The cash balances are held at banks with investment grade credit ratings, which meet our defined minimum credit ratings. The cash at banks is at full disposal of the Company.

6. Property, plant and equipment

At June 30, 2023 and December 31, 2022, property plant and equipment consisted of buildings and leasehold improvements, laboratory equipment and other assets. Buildings and leasehold improvements include a right-of-use asset relating to the lease of our Leiden office and laboratory space, with a carrying amount of € 14,363,000 at June 30, 2023 (December 31, 2022: € 14,484,000).

7. Other current liabilities

At June 30, 2023, other current liabilities amount to € 4,030,000 (December 31, 2022: € 8,687,000). At June 30, 2023 and December 31, 2022, other current liabilities consisted principally of accruals for services provided by vendors not yet billed, payroll related accruals and other miscellaneous liabilities.

8. Borrowings

	June 30, 2023	**December 31, 2022**
	€1,000	€1,000
Innovation credit	3,907	3,907
Accrued interest on innovation credit	1,231	1,035
Convertible notes	992	1,369
Accrued interest on convertible notes	365	460
Total borrowings	**6,495**	**6,771**
Current portion	(2,920)	(2,500)
	3,575	**4,271**

On December 10, 2018 ProQR was awarded an Innovation credit for the sepofarsen program. Amounts were drawn under this facility from 2018 through 2022. The credit of € 3,907,000 was used to conduct the Phase 2/3 clinical study and efforts to obtain regulatory and ethical market approval (NDA/MAA) of sepofarsen for LCA10. The received amount of € 3,907,000 is recognized under borrowings at June 30, 2023 and December 31, 2022. The credit and accrued interest of 10% per annum is repayable depending on the future development of the sepofarsen program.

Convertible loans

Convertible loans were issued to Amylon Therapeutics B.V. (‘Amylon’) and are interest-bearing at an average rate of 8% per annum. They are convertible into a variable number of ordinary shares within 36 months at the option of the holder or the Company in case financing criteria are met. Any unconverted loans become payable on demand after 24 – 36 months in equal quarterly terms.

In 2023 and 2022, Amylon entered into waiver agreements with certain lenders. Such lenders’ loan agreements with Amylon are severed and any claims to repayment of any outstanding debt and accumulated interest are renounced. The amount of convertible loans and accumulated interest waived under these agreements in the six month period ended June 30, 2023 is € 509,000 (six month period ended June 30, 2022: € 1,144,000). The resulting gain was recognized as a gain on derecognition of financial liabilities (refer to note 17).

In September 2022, ProQR extinguished its debt with Pontifax and Kreos by repaying all outstanding principal amounts. Pontifax’ and Kreos’ warrants remain in place until their five-year economic life expires. These warrants are accounted for as embedded derivatives and were recognized separately from the host contract as derivative financial liabilities at fair value through profit or loss.

9. Lease liabilities

At June 30, 2023 and December 31, 2022, lease liabilities primarily consisted of the Company’s lease of office and laboratory facilities at Zernikedreef in Leiden, the Netherlands.

The Company leases office and laboratory facilities of 4,818 square meters at Zernikedreef in Leiden, the Netherlands, where our headquarters and our laboratories are located. The current lease agreement for these facilities terminates on June 30, 2031. The lease agreement contains no significant dismantling requirements.

The initial 10-year lease agreement for the Leiden office and laboratory facilities was accounted for as of commencement date July 1, 2020. This 10-year period was extended by 1 year to an 11-year period in December 2020. The lease contract may be extended for subsequent 5-year periods. As the Company is not reasonably certain to exercise these extension options, these are not included in the lease term.

The carrying amount of the right-of-use asset is disclosed in note 6.

10. Deferred income

The following table summarizes details of deferred income at June 30, 2023 and December 31, 2022. The nature of the deferred income is described in Note 12.

	June 30, 2023	**December 31, 2022**
	€1,000	*€1,000*
Eli Lilly up-front payment and equity consideration: current portion	13,137	5,641
Eli Lilly up-front payment and equity consideration: non-current portion	56,239	65,569
Total deferred income	**69,376**	**71,210**

11. Shareholders’ equity

The authorized share capital of the Company amounting to € 13,600,000 consists of 170,000,000 ordinary shares and 170,000,000 preference shares with a par value of € 0.04 per share. At June 30, 2023, 84,246,967 ordinary shares were issued. 80,939,812 ordinary shares were fully paid and 3,307,155 ordinary shares were held by the Company as treasury shares (December 31, 2022: 3,429,888).

In December 2022, the Company issued 9,381,586 shares to Lilly pursuant to the amended and restated licensing and research collaboration between the Company and Lilly, resulting in gross proceeds of € 14,122,000, with no significant transaction costs.

Translation reserve

The translation reserve comprises all foreign currency differences arising from the translation of the financial statements of foreign operations.

Share options
The Company operates an equity-settled share-based compensation plan, which was introduced in 2013. Options and RSUs may be granted to employees, members of the Supervisory Board, members of the Management Board and consultants. The compensation expenses included in operating costs for this plan in the six month period ended June 30, 2023 were € 1,860,000 (six month period ended June 30, 2022: €1,921,000), of which €1,644,000 was recorded in general and administrative costs (six month period ended June 30, 2022: €1,537,000) and €216,000 was recorded in research and development costs (six month period ended June 30, 2022: €384,000).

12. Revenue

Eli Lilly

In September 2021, the Company entered into a global licensing and research collaboration with Eli Lilly and Company ('Lilly') focused on the discovery, development, and commercialization of potential new medicines for genetic disorders in the liver and nervous system. ProQR and Lilly will use ProQR's proprietary Axiomer® RNA editing platform to progress new drug targets toward clinical development and commercialization.

Under the terms of the agreement, ProQR received an upfront payment and equity consideration, and is eligible to receive milestone payments and royalties on the net sales of any resulting products. In September 2021, the Company issued 3,989,976 shares to Lilly, resulting in net proceeds of € 23,223,000. This amount included a price premium of € 2,144,000, which was determined to be part of the transaction price and as such was initially recognized as deferred revenue. An up-front payment of € 17,651,000 was received in October 2021.

With regard to its original collaboration with Lilly, the Company concluded as follows:

- There is one single performance obligation under IFRS 15, which is the transfer of a license combined with the performance of research and development activities. The Company concluded that the license is not capable of being distinct and is not distinct in the context of the contract.

- The transaction price of this agreement currently only includes fixed parts, consisting of an up-front fee and an equity component. The agreement also contains variable parts, but those are not yet included in the transaction price. Milestone payments will only be included to the extent that it is highly probable that a significant reversal in the amount of cumulative revenue recognized will not occur when the uncertainty associated with the milestones is subsequently resolved. Sales-based milestones and sales-based royalties will be included as the underlying sales occur.

- The Company recognizes revenue over time, using an input method that estimates the satisfaction of the performance obligation as the percentage of labor hours incurred compared to the total estimated labor hours required to complete the promised services.

In December 2022, the Company and Lilly amended their research and collaboration agreement described above, which expanded the collaboration. Under the amended and restated research and collaboration agreement, Lilly will gain access to additional targets in the central nervous system and peripheral nervous system with ProQR's Axiomer platform.

As described under Note 11, pursuant to the amended and restated agreement, the Company issued 9,381,586 shares to Lilly in December 2022, resulting in gross proceeds of $ 15,000,000 (€ 14,122,000). These shares were issued at a discount of $ 480,000 (€ 451,000), which is accounted for as a reduction of the transaction price. In February 2023, ProQR also received an upfront payment of $ 60,000,000 (€ 56,254,000), which was recognized under Other Receivables at December 31, 2022. Lilly has the ability to exercise an option to further expand the partnership for a consideration of $ 50,000,000.

With regard to the amended and restated research and collaboration agreement with Lilly, the Company concluded as follows:

- There is one single performance obligation under IFRS 15, which is the transfer of a license combined with the performance of research and development activities. The Company concluded that the license is not capable of being distinct and is not distinct in the context of the contract.

- The transaction price of this agreement currently only includes fixed components, consisting of an up-front fee and an equity component (discount). The agreement also contains variable components, but those are not yet included in the transaction price. Milestone payments will only be included to the extent that it is highly probable that a significant reversal in the amount of cumulative revenue recognized will not occur when the uncertainty associated with the milestones is subsequently resolved. Sales-based milestones and sales-based royalties will be included as the underlying sales occur.

- The Company recognizes revenue over time, using an input method that estimates the satisfaction of the performance obligation as the percentage of labor hours incurred compared to the total estimated labor hours required to complete the promised services.

Yarrow Biotechnology

In May 2021, the Company entered into an exclusive worldwide license and discovery collaboration for an undisclosed target with Yarrow Biotechnology, Inc. (“Yarrow”). Under the terms of the agreement, ProQR received an upfront payment, equity consideration and reimbursement for ongoing R&D services. In May 2021, ProQR received an up-front payment of € 419,000 and 8% of the shares of Yarrow’s common stock, which was subsequently diluted to 5.1%. In 2022, ProQR was also entitled to receive reimbursements for R&D services performed amounting to € 272,000.

Although ProQR only owns 5.1% of Yarrow’s shares, the Company has significant influence over Yarrow by virtue of its right to appoint one of Yarrow’s three board members, as well as its participation in Yarrow’s policy-making process, amongst other factors. As such, our interest in Yarrow amounting to € nil at June 30, 2023 and December 31, 2022 is recognized as an investment in associate.

With regard to its collaboration with Yarrow, the Company concluded as follows:

- There is one single performance obligation under IFRS 15, which is the transfer of a license combined with the performance of research and development activities. The Company concluded that the license is not capable of being distinct and is not distinct in the context of the contract.

- The transaction price of this agreement currently includes both fixed and variable parts. The fixed part consists of an up-front fee and an equity component. The variable part consists of a cost reimbursement for research and development activities. The agreement also contains other variable parts, but those are not yet included in the transaction price. Milestone payments will only be included to the extent that it is highly probable that a significant reversal in the amount of cumulative revenue recognized will not occur when the uncertainty associated with the milestones is subsequently resolved. Sales-based milestones and sales-based royalties will be included as the underlying sales occur.

- The Company recognizes revenue over time, using an input method that estimates the satisfaction of the performance obligation as the percentage of labor hours incurred compared to the total estimated labor hours required to complete the promised services.

The Yarrow collaboration was terminated in Q3 2022.

	Six month period ended June 30,	
	2023	**2022**
	€1,000	*€1,000*
Eli Lilly collaboration revenue	1,860	1,703
Yarrow collaboration revenue	-	357
	1,860	**2,060**

The revenues relating to providing IP licenses and research and development services under the Company's collaboration agreements have no directly associable cost of sales. Costs incurred to fulfill the associated performance obligations are recognized in research and development expenses, due to their being part of the Company's primary activities of biopharmaceutical research and development.

13. Other income

	Six month period ended June 30,	
	2023	**2022**
	€1,000	*€1,000*
Grant income	76	193
Other income	4	7
	80	**200**

On February 9, 2018, the Company entered into a partnership agreement with Foundation Fighting Blindness (FFB), under which FFB has agreed to provide funding of $ 7.5 million for the pre-clinical and clinical development of *ultervursen* for Usher syndrome type 2A targeting mutations in exon 13, of which $ 6.8 million was granted. In the third quarter of 2022, the Company started winding down the clinical studies for *ultevursen*. As of that moment, the Company has ceased recognizing grant income for the FFB grant.

Grants are recognized in other income in the same period in which the related R&D costs are recognized.

14. Research and development costs

Research and development costs amount to € 11,969,000 for the six month period ended June 30, 2023 (six month period ended June 30, 2022: € 24,816,000) and are comprised of allocated employee costs including share-based payments, the costs of materials and laboratory consumables, outsourced activities, license and intellectual property costs and other allocated costs. Research and development costs decreased by € 12,847,000 compared to the same period in the prior year, mainly because the Company's ophthalmology clinical trials were wound down and are no longer active in the first half of 2023, whereas they were ongoing in the first half of 2022.

15. General and administrative costs

General and administrative costs amount to € 8,171,000 for the six month period ended June 30, 2023 (six month period ended June 30, 2022: € 10,320,000).

16. Investment in financial asset

At June 30, 2023, the investment in financial asset amounting to € 621,000 (December 2022: € 621,000) consists of the Company's investment in Phoenicis Therapeutics Inc. In January 2021, Wings Therapeutics Inc. merged into Phoenicis Therapeutics Inc. by means of a non-cash transaction. ProQR holds a 3.9% interest in Phoenicis Therapeutics Inc.

17. Results related to derecognition of financial liabilities

	Six month period ended June 30,	
	2023	**2022**
	€1,000	*€1,000*
Gain on waiver of Amylon convertible loans	509	1,144
	509	**1,144**

Refer to note 8 for a description of convertible loans issued to Amylon.

18. Income taxes

The current income tax liability amounts to € nil at June 30, 2023 (December 31, 2022: € nil). No significant temporary differences exist between accounting and tax results. Realization of deferred tax assets is dependent on future earnings, if any, the timing and amount of which are uncertain. Accordingly, the Company has not yet recognized any deferred tax asset related to operating losses.

Tax losses may be carried forward indefinitely. However, the offset of losses will be limited in a given year against the first € 1 million of taxable profit. For taxable profit in excess of this amount, losses may only be offset up to 50% of this excess.

19. Events after balance sheet date

In August, ProQR announced it will divest its late-stage ophthalmic programs, sepofarsen and ultevursen, to Théa. Under the terms of the agreement, ProQR will receive an initial payment of €12.5M. The transaction is expected to close in the third quarter of 2023, subject to the satisfaction of customary closing conditions. The carrying amounts and classifications of assets and liabilities recognized on the statement of financial position at June 30, 2023 are not affected by the planned divestiture of ProQR's late-stage ophthalmic programs.

Exhibit 99.2

ProQR Announces Second Quarter 2023 Operating and Financial Results

- Divestment of sepofarsen and ultevursen ophthalmic programs supports ProQR's strategic focus on Axiomer® RNA editing platform technology and continued advancement of pipeline
- €129 M cash and cash equivalents as of June 30, 2023 providing runway into mid-2026

LEIDEN, Netherlands & CAMBRIDGE, Mass., August 3, 2023 – ProQR Therapeutics NV. (Nasdaq: PRQR) (ProQR), a company dedicated to changing lives through transformative RNA therapies based on its proprietary Axiomer® RNA editing technology platform, today reported its financial and operating results for the second quarter ended June 30, 2023, and provided a business update.

"As we continue to advance our Axiomer RNA editing platform technology and pipeline with an initial focus on liver-originated diseases, we were very pleased to announce the divestment of our sepofarsen and ultevursen programs to Théa, where they will continue the development of those programs for patients with rare genetic eye disease," said Daniel A. de Boer, Chief Executive Officer of ProQR. "Over the next several quarters, we anticipate presenting and publishing additional preclinical data including for our first two pipeline programs, AX-0810 for patients suffering from cholestatic diseases and AX-1412 for patients with cardiovascular risk. Along with our preclinical proof of concept data for the platform, a partnership with Eli Lilly, a leading IP position, and cash runway into mid-2026, ProQR is well positioned to execute on our strategy."

Recent Progress

- In August, ProQR **announced it will divest its late-stage ophthalmic assets, sepofarsen and ultevursen, to Théa**. Under the terms of the agreement, ProQR will receive an initial payment of €12.5M and will also be eligible for up to €135M in further development, regulatory and commercial payments, as well as additional earn outs up to high teen percentage based on commercial sales in the US and EU. The transaction is expected to close in the third quarter of 2023, subject to the satisfaction of customary closing conditions.
- In July, ProQR presented **an overview of its Axiomer RNA editing platform at the RNA Editing Summit** in Boston. The presentation described ongoing efforts led by ProQR to put in place effective strategies to increase RNA editing yields of ADARs using chemical modifications and oligonucleotide design. Additionally highlighted was the potential of the Axiomer platform in multiple organs, as well as its applicability for the treatment of liver-originated disorders, including data from preclinical models.

Anticipated Upcoming Events

- Present various platform updates over the next several quarters, including liver NHP data, at scientific conferences, as well as research related to ongoing discovery efforts and pipeline programs.
- Continue to execute on existing partnership with Eli Lilly.

- ProQR may selectively form new partnerships, which could include multi-target discovery alliances, similar to the Company's partnership with Lilly, or product alliances on specific programs.
- ProQR is on track to advance AX-0810 targeting NTCP and AX-1412 targeting B4GALT1 into clinical development in late 2024/early 2025.

Financial Highlights

On June 30, 2023, ProQR held cash and cash equivalents of €128.6 million, compared to €94.8 million on December 31, 2022. Net cash used in operating activities during the three-month period ended June 30, 2023 was €10.3 million, compared to €14.8 million for the same period last year.
Research and development costs were €5.9 million for the quarter ended June 30, 2023 compared to €11.4 million for the same period last year.
General and administrative costs were €4.1 million for the quarter ended June 30, 2023 compared to €5.4 million for the quarter ended June 30, 2022.
Net loss for the three-month period ended June 30, 2023 was €8.0 million, or €0.10 per diluted share, compared to €13.5 million, or €0.19 per diluted share, for the same period last year. For further financial information for the period ending June 30, 2023, please refer to the financial statements appearing at the end of this release.

About Axiomer®

ProQR is pioneering a next-generation RNA base editing technology called Axiomer®, which could potentially yield a new class of medicines for diverse types of diseases. Axiomer® "Editing Oligonucleotides", or EONs, mediate single nucleotide changes to RNA in a highly specific and targeted way using molecular machinery that is present in human cells called ADAR (Adenosine Deaminase Acting on RNA). Axiomer® EONs are designed to recruit and direct endogenously expressed ADARs to change an Adenosine (A) to an Inosine (I) in the RNA – an Inosine is translated as a Guanosine (G) – correcting an RNA with a disease-causing mutation back to a normal (wild type) RNA, modulating protein expression, or altering a protein so that it will have a new function that helps prevent or treat disease.

About Biliary Atresia (BA) and Primary Sclerosing Cholangitis (PSC)

Cholestatic disorders refer to a group of diseases presenting excessive and toxic buildup of bile acids in the liver due to bile ducts dysfunction. This leads to liver damage and a range of debilitating symptoms. Without treatment, liver damage can progress through various stages, ultimately leading to liver failure and elevated risk of liver malignancy, affecting life expectancy. Cholestatic diseases remain leading causes of liver transplantation. There are no approved therapies for primary sclerosing cholangitis (PSC) for adults and biliary atresia (BA) for pediatrics It is estimated that 80,000 and 20,000 individuals have PSC and BA, respectively, in North America and in Europe.

About AX-0810 targeting NTCP

The majority of the bile acids present in the liver cells originate from the enterohepatic reuptake cycle. The key transporter responsible for hepatic uptake of bile acids from portal circulation is the sodium (Na+)-taurocholate cotransporting polypeptide (NTCP, SLC10A1 gene) expressed in the liver. AX-0810 is designed to introduce a loss of function variant in SLC10A1 RNA that has been found in human genetics to prevent re-uptake of bile acids in liver via NTCP. Based on its mechanism of action, AX-0810 has the potential to become a disease modifying treatment for PSC and BA primarily among other cholestatic diseases.

About Cardiovascular Diseases

Cardiovascular diseases (CVDs) are a group of health conditions that affect the heart and blood vessels, such as atherosclerosis which can lead to severe problems like heart attacks, heart failure, and stroke. CVDs represent the leading cause of disability and death in the world. Approximately 18 million people die every year from CVDs representing one third of all the global deaths. Despite available lipid lowering therapies and hypertension medications, the risk of CVDs is still projected to increase rapidly over the coming years.

About AX-1412 targeting B4GALT1

Gene–based analysis of rare beta-1,4-galactosyltransferase 1 (*B4GALT1*) missense variant (p.Asn352Ser) is known to lead to B4GALT1 protein loss of function and showed an association with decreased coronary artery disease. These beneficial effects are mediated by hypo-galactosylation of the apolipoprotein B100 and fibrinogen, known – independent – drivers of increased risk of CVDs. AX-1412 introduces a protective variant into *B4GALT1* RNA to address the remaining residual risk of developing cardiovascular diseases. ProQR intends to advance AX-1412 targeting B4GALT1 to early clinical proof of concept stage, then would seek to partner this program.

About ProQR

ProQR Therapeutics is dedicated to changing lives through the creation of transformative RNA therapies. ProQR is pioneering a next-generation RNA technology called Axiomer®, which uses a cell's own editing machinery called ADAR to make specific single nucleotide edits in RNA to reverse a mutation or modulate protein expression and could potentially yield a new class of medicines for both rare and prevalent diseases with unmet need. Based on our unique proprietary RNA repair platform technologies we are growing our pipeline with patients and loved ones in mind.

Learn more about ProQR at **www.proqr.com**.

Forward Looking Statements

This press release contains forward-looking statements. All statements other than statements of historical fact are forward-looking statements, which are often indicated by terms such as "anticipate," "believe," "could," "estimate," "expect," "goal," "intend," "look forward to", "may," "plan," "potential," "predict," "project," "should," "will," "would" and similar expressions. Such forward-looking statements include, but are not limited to, statements regarding our business, preclinical model data, our initial pipeline targets, our Axiomer platform , the divestment and further development of sepofarsen and ultevursen and the potential payments and earnouts arising out of the divestment, the expected timing for the closing of the divestment, as well as the potential of our technologies and product candidates, the collaboration with Eli Lilly and Company ("Lilly") and the intended benefits thereof, and our financial position and cash-runway. Forward-looking statements are based on management's beliefs and assumptions and on information available to management only as of the date of this press release. Our actual results could differ materially from those anticipated in these forward-looking statements for many reasons, including, without limitation, the risks, uncertainties and other factors in our filings made with the Securities and Exchange Commission, including certain sections of our annual report filed on Form 20-F. These risks and uncertainties include, among others, the clinical development activities to be performed by Théa and the condition of successful market access for sepofarsen and ultevursen, the cost, timing and results of preclinical studies and clinical trials and other development activities by us and our collaborative partners whose operations and activities may be slowed or halted shortage and pressure on supply and logistics on the global market; the likelihood of our preclinical and clinical programs being initiated and executed on timelines provided and reliance on our contract research organizations and predictability of timely enrollment of subjects and patients to advance our clinical trials and maintain their own operations; our reliance on contract manufacturers to supply materials for research and development and the risk of supply interruption from a contract manufacturer; the potential for future data to alter initial and preliminary results of early-stage clinical trials; the unpredictability of the duration and results of the regulatory review of applications or clearances that are necessary to initiate and continue to advance and progress our clinical programs; the ability to secure, maintain and realize the intended benefits of collaborations with partners, including the collaboration with Lilly; the possible impairment of, inability to obtain, and costs to obtain intellectual property rights; possible safety or efficacy concerns that could emerge as new data are generated in research and development; and general business, operational, financial and accounting risks, and risks related to litigation and disputes with third parties. Given these risks, uncertainties and other factors, you should not place undue reliance on these forward-looking statements, and we assume no obligation to update these forward-looking statements, even if new information becomes available in the future, except as required by law.

ProQR Therapeutics N.V.

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Financial Tables

PROQR THERAPEUTICS N.V.
Unaudited Condensed Consolidated Statement of Financial Position

	June 30, 2023	December 31, 2022
	€1,*000*	€1,*000*
Assets		
Current assets		
Cash and cash equivalents	128,562	94,775
Prepayments and other receivables	2,187	59,078
Other taxes	400	607
Total current assets	**131,149**	**154,460**
Property, plant and equipment	16,220	16,240
Investments in financial assets	621	621
Total assets	**147,990**	**171,321**
Equity and liabilities		
Equity		
Equity attributable to owners of the Company	51,790	67,064
Non-controlling interests	(290)	(384)
Total equity	**51,500**	**66,680**
Current liabilities		
Borrowings	2,920	2,500
Lease liabilities	1,334	1,387
Derivative financial instruments	372	1,263
Trade payables	119	392
Social securities and other taxes	1,064	1,118
Deferred income	13,137	5,641
Other current liabilities	4,030	8,687
Total current liabilities	**22,976**	**20,988**
Borrowings	3,575	4,271
Lease liabilities	13,700	13,813
Deferred income	56,239	65,569
Total liabilities	**96,490**	**104,641**
Total equity and liabilities	**147,990**	**171,321**

PROQR THERAPEUTICS N.V.

Unaudited Condensed Consolidated Statement of Profit or Loss and OCI

(€ in thousands, except share and per share data)

	Three month period ended June 30,		Six month period ended June 30,	
	2023	2022	2023	2022
	€*1,000*	€*1,000*	€*1,000*	€*1,000*
Revenue	**1,205**	**930**	**1,860**	**2,060**
Other income	**38**	**99**	**80**	**200**
Research and development costs	(5,909)	(11,449)	(11,969)	(24,816)
General and administrative costs	(4,145)	(5,412)	(8,171)	(10,320)
Total operating costs	**(10,054)**	**(16,861)**	**(20,140)**	**(35,136)**
Operating result	**(8,811)**	**(15,832)**	**(18,200)**	**(32,876)**
Finance income and expense	470	1,163	(74)	342
Results related to associates	—	—	—	(8)
Results related to financial liabilities measured at fair value through profit or loss	221	62	891	3,826
Results on derecognition of financial liabilities	101	1,144	509	1,144
Result before corporate income taxes	**(8,019)**	**(13,463)**	**(16,874)**	**(27,572)**
Income taxes	42	(20)	42	(27)
Result for the period	**(7,977)**	**(13,483)**	**(16,832)**	**(27,599)**
Other comprehensive income (foreign exchange differences on foreign operation)	7	689	(212)	911
Total comprehensive income	**(7,970)**	**(12,794)**	**(17,044)**	**(26,688)**
Result attributable to				
Owners of the Company	(7,993)	(13,700)	(16,926)	(27,808)
Non-controlling interests	16	217	94	209
	(7,977)	**(13,483)**	**(16,832)**	**(27,599)**
Total comprehensive income attributable to				
Owners of the Company	(7,986)	(13,011)	(17,138)	(26,897)
Non-controlling interests	16	217	94	209
	(7,970)	**(12,794)**	**(17,044)**	**(26,688)**
Share information				
Weighted average number of shares outstanding[1]	80,939,392	71,362,088	80,913,751	71,359,642
Earnings per share attributable to owners of the Company (Euro per share)				
Basic loss per share[1]	(0.10)	(0.19)	(0.21)	(0.39)
Diluted loss per share[1]	(0.10)	(0.19)	(0.21)	(0.39)

PROQR THERAPEUTICS N.V.

Unaudited Condensed Consolidated Statement of Changes in Equity

	Attributable to owners of the Company									
	Number of shares	**Share Capital**	**Share Premium**	**Equity settled Employee Benefit Reserve**	**Option premium on convertible loan**	**Translation Reserve**	**Accumulated Deficit**	**Total**	**Non-controlling interests**	**Total Equity**
		€*1,000*	€*1,000*	€*1,000*	€*1,000*	€*1,000*	€*1,000*	€*1,000*	€*1,000*	€*1,000*
Balance at January 1, 2022	**74,865,381**	**2,995**	**398,309**	**28,443**	**1,426**	**430**	**(316,890)**	**114,713**	**(604)**	**114,109**
Result for the period	—	—	—	—	—	—	(27,808)	(27,808)	209	(27,599)
Other comprehensive income	—	—	—	—	—	911	—	911	—	911
Recognition of share-based payments	—	—	—	1,921	—	—	—	1,921	—	1,921
Treasury shares transferred	(71,283)	—	—	—	—	—	—	—	—	—
Share options lapsed	—	—	—	(380)	—	—	380	—	—	—
Share options exercised	71,283	—	33	(256)	—	—	256	33	—	33
Balance at June 30, 2022	**74,865,381**	**2,995**	**398,342**	**29,728**	**1,426**	**1,341**	**(344,062)**	**89,770**	**(395)**	**89,375**
Balance at January 1, 2023	**84,246,967**	**3,370**	**412,540**	**29,052**	**—**	**1,212**	**(379,110)**	**67,064**	**(384)**	**66,680**
Result for the period	—	—	—	—	—	—	(16,926)	(16,926)	94	(16,832)
Other comprehensive income	—	—	—	—	—	(212)	—	(212)	—	(212)
Recognition of share-based payments	—	—	—	1,860	—	—	—	1,860	—	1,860
Treasury shares transferred	(122,584)	—	—	—	—	—	—	—	—	—
Share options lapsed	—	—	—	(3,873)	—	—	3,873	—	—	—
Share options exercised / RSUs vested	122,584	—	4.00	(231)	—	—	231	4	—	4
Balance at June 30, 2023	**84,246,967**	**3,370**	**412,544**	**26,808**	**—**	**1,000**	**(391,932)**	**51,790**	**(290)**	**51,500**

PROQR THERAPEUTICS N.V.

Unaudited Condensed Consolidated Statement of Cash Flows

	Three month period ended June 30,		Six month period ended June 30,	
	2023	**2022**	**2023**	**2022**
	€1,*000*	€1,*000*	€1,*000*	€1,*000*
Cash flows from operating activities				
Net result	(7,977)	(13,483)	(16,832)	(27,599)
Adjustments for:				
— Depreciation	594	591	1,143	1,161
— Share-based compensation	765	738	1,860	1,921
— Financial income and expenses	(470)	(1,163)	74	(342)
— Results related to associates	—	—	—	8
— Results related to financial liabilities measured at fair value through profit or loss	(221)	(62)	(891)	(3,826)
— Results on derecognition of financial liabilities	(101)	(1,144)	(509)	(1,144)
— Income tax expenses	—	20	—	27
Changes in working capital	(3,622)	916	48,668	(3,035)
Cash (used in)/generated from operations	*(11,032)*	*(13,587)*	*33,513*	*(32,829)*
Corporate income tax paid	—	(20)	—	(27)
Interest received	685	—	865	—
Interest paid	—	(1,237)	—	(2,455)
Net cash (used in)/generated from operating activities	***(10,347)***	***(14,844)***	***34,378***	***(35,311)***
Cash flow from investing activities				
Purchases of property, plant and equipment	(294)	(231)	(430)	(475)
Sales of property, plant and equipment	—	—	47	—
Net cash used in investing activities	***(294)***	***(231)***	***(383)***	***(475)***
Cash flow from financing activities				
Proceeds from exercise of share options	4	—	4	33
Repayment of lease liability	(647)	(357)	(906)	(933)
Net cash used in financing activities	***(643)***	***(357)***	***(902)***	***(900)***
Net (decrease)/increase in cash and cash equivalents	**(11,284)**	**(15,432)**	**33,093**	**(36,686)**
Currency effect cash and cash equivalents	860	4,222	694	5,564
Cash and cash equivalents, at beginning of the period	138,986	167,612	94,775	187,524
Cash and cash equivalents at the end of the period	**128,562**	**156,402**	**128,562**	**156,402**