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

For the month of November 2019

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 x Form 40-F o

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): o

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): o

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 nine month periods ended September 30, 2019, and furnished as Exhibit 99.2 to this Report on Form 6-K is a press release of the Company, dated November 6, 2019, announcing the Company's results for the three and nine month periods ended September 30, 2019.

The Company hereby incorporates by reference the information contained herein into the Company's registration statement on Form F-3 (File No. 333-228251).

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: November 12, 2019

By: /s/ Smital Shah

Smital Shah

Chief Financial Officer

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Unaudited Condensed Consolidated Statement of Financial Position

	September 30, 2019 € 1,000	December 31, 2018 € 1,000
Assets		,,,,,,
Current assets		
Cash and cash equivalents	74,792	105,580
Prepayments and other receivables	2,450	1,544
Social securities and other taxes	830	1,243
Total current assets	78,072	108,367
Property, plant and equipment	2.413	1,864
Investments in associates	579	
Total assets	81,064	110,231
Equity and liabilities		
Equity		
Equity attributable to owners of the Company	59,871	92,915
Non-controlling interests	(452)	(230)
Total equity	59,419	92,685
Current liabilities		
Borrowings	260	_
Lease liabilities	890	_
Trade payables	668	135
Social securities and other taxes	15	_
Pension premiums	13	7
Deferred income	984	545
Other current liabilities	8,305	7,473
Total current liabilities	11,135	8,160
Borrowings	10,510	9,386
Lease liabilities		
Total liabilities	21,645	17,546
Total equity and liabilities	<u>81,064</u>	110,231

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

PROQR THERAPEUTICS | ZERNIKEDREEF 9 | 2333 CK LEIDEN | THE NETHERLANDS | +31 88 166 7000 | WWW.PROQR.COM

Unaudited Condensed Consolidated Statement of Profit or Loss and OCI

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

	Three month p ended Septemb		Nine month p ended Septemb	
	2019	2018	2019	2018
Otherstone	€ 1,000	€ 1,000	€ 1,000	€ 1,000
Other income	530	2,958	1,509	4,428
Research and development costs	(11,074)	(6,297)	(32,560)	(19,972)
General and administrative costs	(2,903)	(2,579)	(8,970)	(7,900)
Total operating costs	(13,977)	(8,876)	(41,530)	(27,872)
Operating result	(13,447)	(5,918)	(40,021)	(23,444)
Finance income and expense	1,375	(74)	1,339	(664)
Results related to associates	(119)	_	579	_
De liberario de la constanta d	(40.404)	(F.002)	(20.402)	(24.100)
Result before corporate income taxes Income taxes	(12,191)	(5,992)	(38,103) (64)	(24,108) (1)
income taxes	-	_	(04)	(1)
Result for the period	(12,191)	(5,992)	(38,167)	(24,109)
Other comprehensive income	147	(4)	121	(15)
Total comprehensive income (attributable to owners of the				
Company)	(12,044)	(5,996)	(38,046)	(24,124)
Result attributable to				
Owners of the Company	(12,139)	(5,959)	(37,945)	(23,974)
Non-controlling interests	(52)	(33)	(222)	(135)
	(12,191)	(5,992)	(38,167)	(24,109)
Share information				
Weighted average number of shares outstanding(1)	38,912,701	33,355,327	38,902,203	32,440,220
Earnings per share attributable to the equity holders of the Company (expressed in Euro per share)				
Basic loss per share(1)	(0.31)	(0.18)	(0.98)	(0.74)
Diluted loss per share(1)	(0.31)	(0.18)	(0.98)	(0.74)

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

⁽¹⁾ For this period presented in these financial statements, the potential exercise of share options is not included in the diluted earnings per share calculation as the Company was loss-making in all periods. Due to the anti-dilutive nature of the outstanding options, basic and diluted earnings per share are equal in this period.

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

	Attributable to owners of the Company								
				Equity Settled Employee				Non-	
	Number of shares	Share Capital € 1,000	Share Premium € 1,000	Benefit Reserve € 1,000	Translation Reserve € 1,000	Accumulated Deficit € 1.000	Total € 1,000	controlling interests € 1,000	Total Equity € 1,000
Balance at January 1, 2018	36,425,014	1,457	148,763	8,377	136	(119,370)	39,363	(38)	39,325
Result for the period						(23,974)	(23,974)	(135)	(24,109)
Other comprehensive income	_	_	_	_	(15)		(15)	<u> </u>	(15)
Recognition of share-based									
payments	_			2,245		_	2,245		2,245
Issuance of ordinary shares	6,612,500	265	84,032	_	_	_	84,297	_	84,297
Share options exercised			659				659		659
Balance at									
September 30, 2018	43,037,514	1,722	233,454	10,622	121	(143,344)	102,575	(173)	102,402
Balance at January 1, 2019	43,149,987	1,726	235,744	10,780	108	(155,443)	92,915	(230)	92,685
Result for the period	_	_	_	_	_	(37,945)	(37,945)	(222)	(38,167)
Other comprehensive income	_	_	_	_	121	_	121	_	121
Recognition of share-based									
payments	_	_	_	4,614	_	_	4,614	_	4,614
Share options lapsed	_	_	_	(33)		33	_	_	
Share options exercised			166	(115)		115	166		166
Balance at									
September 30, 2019	43,149,987	1,726	235,910	15,246	229	(193,240)	59,871	(452)	59,419

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

Unaudited Condensed Consolidated Statement of Cash Flows

	Three month p ended Septeml		Nine month p ended Septeml	
	2019	2018	2019	2018
	€ 1,000	€ 1,000	€ 1,000	€ 1,000
Cash flows from operating activities	(12.101)	(F.002)	(20.167)	(24.100)
Net result	(12,191)	(5,992)	(38,167)	(24,109)
Adjustments for:	F06	2.42	4.540	E0.5
— Depreciation	506	242	1,543	725
— Share-based compensation	1,226	734	4,614	2,245
— Financial income and expenses	(1,375)	74	(1,339)	664
— Results related to associates	119		(579)	
— Net foreign exchange gain / (loss)	148	(4)	122	(15)
Changes in working capital	2,718	656	1,744	1,074
Cash used in operations	(8,849)	(4,290)	(32,062)	(19,416)
Corporate income tax paid	_	1	(64)	_
Interest received	90	32	176	25
Interest paid	(13)		(64)	_
Net cash used in operating activities	(8,772)	(4,257)	(32,014)	(19,391)
Cash flow from investing activities				
Purchases of intangible assets				
Purchases of property, plant and equipment	(32)	(99)	(341)	(285)
ruchases of property, plant and equipment	(32)	(99)	(341)	(203)
Net cash used in investing activities	(32)	(99)	(341)	(285)
Cash flow from financing activities				
Proceeds from issuance of shares, net of transaction costs	_	84,295	_	84,295
Proceeds from exercise of share options	2	637	166	660
Proceeds from borrowings	_	_	_	101
Proceeds from convertible loans	_	115	690	430
Repayment of lease liability	(290)		(861)	
Net cash (used in)/generated by financing activities	(288)	85,047	(5)	85,486
Net increase/(decrease) in cash and cash equivalents	(9,092)	80,691	(32,360)	65,810
·		<u> </u>		<u> </u>
Currency effect cash and cash equivalents	1,420	57	1,572	(193)
Cash and cash equivalents, at beginning of the period	82,464	32,968	105,580	48,099
Cash and cash equivalents at the end of the period	74,792	113,716	74,792	113,716

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

Notes to Unaudited Condensed Consolidated Financial Statements

1. General information

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

Since September 18, 2014, the Company's ordinary shares are listed on the NASDAQ Global Market under ticker symbol PRQR.

The Company was incorporated in the Netherlands, on February 21, 2012 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%);
- ProOR Therapeutics IV 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%);
- ProOR Therapeutics I Inc. (100%):
- · Amylon Therapeutics B.V. (80%);
- Amylon Therapeutics Inc. (80%).

ProQR Therapeutics N.V. is also statutory director of Stichting Bewaarneming Aandelen ProQR ("ESOP Foundation") and has full control over this entity. ProQR Therapeutics N.V. holds a minority shareholding in Wings Therapeutics 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.

2. Significant Accounting Policies

These condensed consolidated financial statements have been prepared in accordance with IAS 34 - Interim Financial Reporting. Certain information and disclosures normally included in financial statements prepared in accordance with IFRS have been condensed or omitted. Accordingly, these condensed consolidated financial statements should be read in conjunction with the Company's annual financial statements for the year ended December 31, 2018. In the opinion of management, all adjustments, consisting of normal recurring nature, considered necessary for a fair presentation have been included in the condensed consolidated financial statements.

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 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

The accounting policies adopted in the preparation of the 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, 2018, except for the change in accounting policies resulting from the implementation of IFRS 16 *Leases*.

IFRS 16 specifies how an entity recognizes, measures, presents and discloses leases. The standard provides a single lessee accounting model, requiring lessees to recognize assets and liabilities for all leases unless the lease term is 12 months or less or the underlying asset has a low value. Upon implementation of the standard on January 1, 2019, the Company recognized a lease liability and a corresponding right-of-use asset. As at September 30, 2019, the carrying amount of the lease liability is \in 0.9 million and the carrying amount of the right-of-use asset is \in 0.9 million.

The impact on the income statement is that operating expenses are replaced by depreciation expenses on the right-of-use asset and interest expenses on the lease liability. The main impact on the statement of cash flows is higher cash flows from operating activities, since cash payments for the principal part of the lease liability are classified as cash flows used in financing activities, whereas such payments were previously classified as cash flows used in operating activities. (effect on nine month period ended September 2019: € 0.9 million).

Other new Standards and Interpretations, which became effective as of January 1, 2019, 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.

(a) Share-based payments

Share options granted to employees and consultants are measured at the fair value of the equity instruments granted. Fair value is determined through the use of the Black-Scholes option-pricing model, which is considered the most appropriate model for this purpose by management.

Initially, the Company's ordinary shares were not publicly traded and consequently the Company needed to estimate the fair value of its share and the expected volatility of that value. Please refer to the Company's annual financial statements for the year ended December 31, 2018 for the assumptions used in those estimates. The value of the underlying shares

was determined on the basis of the prior sale of company stock method. As such, the Company has benchmarked the value per share to external transactions of Company shares and external financing rounds.

For options granted from the moment of listing, the Company uses the closing price of the ordinary shares on the previous business day as exercise price of the options granted.

The result of the share option valuations and the related compensation expense is dependent on the model and input parameters used. Even though Management considers the fair values reasonable and defensible based on the methodologies applied and the information available, others might derive a different fair value for the Company's share options.

(b) Corporate income taxes

The Company recognizes deferred tax assets arising from unused tax losses or tax credits only to the extent that the Company has sufficient taxable temporary differences or there is convincing evidence that sufficient taxable profit will be available against which the unused tax losses or unused tax credits can be utilized. Management's judgment is that such convincing evidence is currently not sufficiently available and a deferred tax asset is therefore only recognized to the extent that the Company has sufficient taxable temporary differences.

(c) Grant income

Grant income is not recognized until there is reasonable assurance that the Company will comply with the conditions attached to them. Grants are recognized in profit or loss on a systematic basis over the period the Company recognizes as expenses the related costs for which the grants are expected to compensate.

(d) Research and development expenditures

Research expenditures are currently not capitalized but are 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 September 30, 2019, the Company's cash and equivalents were \in 74,792,000 as compared to \in 105,580,000 at December 31, 2018. An amount of \in 35,166,000 of the cash balance is denominated in US dollars. The cash balances are held at banks with investment grade credit ratings. The cash at banks is at full disposal of the Company.

6. Current liabilities

At September 30, 2019 and December 31, 2018, the other current liabilities consisted principally of accruals for services provided by vendors not yet billed, payroll related accruals and other miscellaneous liabilities.

7. Borrowings

	September 30, 2019 € 1,000	December 31, 2018 € 1,000
Innovation credit	5,164	5,164
Accrued interest on innovation credit	2,915	1,871
Convertible notes	2,473	1,783
Accrued interest on convertible notes	218	568
Total borrowings	10,770	9,386
Current portion	(260)	
	10,510	9,386

On June 1, 2012, ProQR was awarded an Innovation credit by the Dutch government, through its agency RVO of the Ministry of Economic Affairs, for the Company's cystic fibrosis program. Amounts were drawn under this facility in the course of the years 2013 through 2017. The credit covers 35% of the costs incurred in respect of the program up to € 5.0 million.

The credit is interest-bearing at a rate of 10% per annum. Early October 2018 ProQR received a conditional waiver of the €5 million Innovation credit. Consequently, the repayment of the total loan of €7.9 million, including interest, has been waived if conditions are met, which will be reviewed annually for 3 years. The assets which are co-financed with the granted innovation credit are subject to a right of pledge for the benefit of RVO.

On December 10, 2018 ProQR was awarded an Innovation credit for the QR-110 program. Amounts will be drawn under this facility from 2018 through 2021. The credit of € 4.7 million through December 31, 2021 will be used to conduct the Phase 2/3 clinical study and efforts to obtain regulatory and ethical market approval (NDA/MAA) of QR-110 for LCA10, of which €0.2 million has been received at September 30, 2019. The credit, including accrued interest of 10% per annum, is repayable depending on obtaining market approval.

Convertible loans

Convertible loans were issued to Amylon Therapeutics B.V. 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.

8. Shareholders' equity

The authorized share capital of the Company amounting to \in 7,200,000 consists of 90,000,000 ordinary shares and 90,000,000 preference shares with a par value of \in 0.04 per share. At September 30, 2019, 43,149,987 ordinary shares were issued and fully paid in cash, of which 4,236,792 were held by the Company as treasury shares (December 31, 2018: 4,277,051).

In November 2018, the Company issued 112,473 shares in the aggregate amount of \$2.5 million, at \$22.23 (€19.46) per share to Ionis Pharmaceuticals, Inc. Under the terms of the agreement, an upfront payment in ordinary shares to its common stock, was made to Ionis upon signing the worldwide license agreement. The Company was granted an exclusive worldwide license to QR-1123 and relevant patents. The Company will also make future milestone payments, certain of which will be made in equity and others in cash or equity at the company's discretion, and royalties on net sales of 20% through the royalty term.

On November 7, 2018, the Company filed a shelf registration statement, which permitted: (a) the offering, issuance and sale by the Company of up to a maximum aggregate offering price of \$300,000,000 of its ordinary shares, warrants and/or units; and (b) as part of the \$300,000,000, the offering, issuance and sale by us of up to a maximum aggregate

offering price of \$75,000,000 of its ordinary shares that may be issued and sold under a sales agreement with H.C. Wainwright & Co in one or more at-the-market offerings. In 2018, no shares were issued pursuant to our ATM facility.

In September 2018, the Company consummated an underwritten public offering and concurrent registered direct offering of 6,612,500 ordinary shares at an issue price of \$ 15.75 per share. The gross proceeds from this offering amounted to & 89,983,000 while the transaction costs amounted to & 5,792,000, resulting in net proceeds of & 84,191,000.

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 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 three month period ended September 30, 2019 were € 1,226,000 (three month period ended September 30, 2018: € 735,000), of which € 835,000 (2018: € 550,000) was recorded in general and administrative costs and € 391,000 (2018: € 185,000) was recorded in research and development costs.

9. Other income

	Three month p Septemb	
	2019 € 1,000	2018 € 1,000
Grant income	518	2,835
Other income	12	123
	530	2,958

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 QR-421a for Usher syndrome type 2A targeting mutations in exon 13.

In addition, funding was received for our Huntington's disease program.

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

10. Research and development costs

Research and development costs amount to \in 11,074,000 for the three month period ended September 30, 2019 compared to \in 6,297,000 for same period in 2018 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.

11. General and administrative costs

General and administrative costs amount to € 2,903,000 for the three month period ended September 30, 2019 compared to € 2,579,000 for the same period in 2018.

12. Results related to associates

Results related to associates for the three month period ended September 30, 2019 consist of our share of the net loss of Wings Therapeutics Inc. amounting to € 119,000.

13. Income taxes

Due to the operating losses incurred since inception the Company has no tax provisions as of the balance sheet date. Furthermore, 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.

14. Events after balance sheet date

In October 2019, the Company consummated an underwritten public offering of 10,454,545 ordinary shares at an issue price of \$5.50 per share. The gross proceeds from this offering amounted to \$57,500,000 while the transaction costs amounted to \$3,575,000, resulting in net proceeds of \$53,925,000.

FINAL — FOR RELEASE

ProQR Announces Recent Progress and Financial Results for the Third Quarter of 2019

- · Encouraging clinical data reported from Phase 1/2 trial of sepofarsen for LCA10
- · Initial clinical data from Phase 1/2 trial of QR-421a for Usher syndrome type 2 on track for Q1 2020
- · QR-1123 Investigational New Drug application active for autosomal dominant retinitis pigmentosa
- €49.3 million net proceeds from public offering extends cash runway into 2022

LEIDEN, Netherlands & CAMBRIDGE, Mass., Nov 06, 2019 — ProQR Therapeutics N.V. (Nasdaq: PRQR) (the "Company"), a company dedicated to changing lives through the creation of transformative RNA medicines for severe genetic rare diseases, today announced results for the third quarter of 2019.

"During the third quarter we made great progress towards our mission to bring novel RNA therapies to patients and we recently strengthened our capital position," said Daniel A. de Boer, chief executive officer of ProQR. "In October we announced that LCA10 patients experienced a significant improvement in vision after treatment with sepofarsen in a Phase 1/2 trial. These results strengthen our confidence in the design of the ongoing pivotal *Illuminate* Phase 2/3 trial and our broader inherited retinal diseases pipeline. The Phase 1/2 trial of QR-421a in Usher syndrome is on track to deliver first clinical data in Q1 2020 and we are about to dose a first patient in a Phase 1/2 trial of QR-1123 for adRP. We strengthened our financial position with a public offering in October that will allow us to fund operations into 2022, well beyond the expected top-line readout of Illuminate."

Corporate Highlights and Business Update

Sepofarsen (QR-110) for Leber's congenital amaurosis 10 (LCA10)

- Presented positive top-line results from the Phase 1/2 clinical trial. In the trial, sepofarsen was observed to be well tolerated with rapid, significant and durable improvements in vision observed at month twelve. The target registration dose for the ongoing Phase 2/3 *Illuminate* trial showed a favorable benefit/risk profile in the Phase 1/2 trial.
- The top-line results from the Phase 1/2 trial support confidence in the design of the ongoing Phase 2/3 *Illuminate* trial that could be the sole registration trial for the program. Top-line data from *Illuminate* are expected during the first half of 2021.
- · Received Rare Pediatric Disease designation from the Food and Drug Administration (FDA) for the treatment of LCA10.

QR-421a for Usher syndrome type 2

 The Phase 1/2 Stellar trial of QR-421a in patients with Usher syndrome type 2 is ongoing and on track to deliver interim data during the first quarter of 2020

QR-1123 for autosomal dominant retinitis pigmentosa (adRP)

- The FDA cleared the Investigational New Drug (IND) application to start a first-in-human clinical trial in patients with adRP. ProQR plans to start enrolling patients in the Phase 1/2 *Aurora* trial in 2019.
- · Received Fast Track designation from the FDA.

Business Updates

· Closed an underwritten public offering of 9,090,909 ordinary shares on October 18, 2019 at a price of \$5.50 per share with full exercise of underwriters' option to purchase 1,363,636 additional ordinary shares. Net proceeds totaled approximately €49.3 million. With the addition of this capital ProQR's operations are funded into 2022.

Financial Highlights

At September 30, 2019, prior to the offering on October 18^{th} , ProQR held cash and cash equivalents of €74.8 million, compared to €105.6 million at December 31, 2018. The cash balance at September 30, 2019 excludes the net proceeds of €49.3 million from the underwritten offering of ordinary shares in October 2019. Net cash used in operating activities during the three-month period ended September 30, 2019 was €8.8 million, compared to €4.3 million for the same period last year.

Research and development costs increased to €11.1 million for the quarter ended September 30, 2019 compared to €6.3 million for the same period last year due to increased clinical trial activity.

General and administrative costs increased to €2.9 million for the quarter ended September 30, 2019 compared to €2.6 million for the same period last year.

Net loss for the three-month period ended September 30, 2019 was €12.2 million or €0.31 per share, compared to a €6.0 million loss or €0.18 per share for the same period last year.

In October 2019 the Company sold an aggregate of 10.5 million ordinary shares, with net proceeds to the Company of €49.3 million.

For further financial information for the period ended September 30, 2019, please refer to the financial statements appearing at the end of this release.

About Sepofarsen

Sepofarsen (QR-110) is a first-in-class investigational RNA-based oligonucleotide designed to address the underlying cause of Leber's congenital amaurosis 10 due to the p.Cys998X mutation (also known as the c.2991+1655A>G mutation) in the *CEP290* gene. The p.Cys998X mutation leads to aberrant splicing of the mRNA and non-functional CEP290 protein. Sepofarsen is designed to enable normal splicing, resulting in restoration of normal (wild type) *CEP290* mRNA and subsequent production of functional CEP290 protein. Sepofarsen is intended to be administered through intravitreal injections in the eye and has been granted orphan drug designation in the United States and the European Union and received fast-track designation and rare pediatric disease designation from the FDA as well as access to the PRIME scheme by the EMA.

About QR-421a

QR-421a is a first-in-class investigational RNA-based oligonucleotide designed to address the underlying cause of vision loss in Usher syndrome type 2 and non-syndromic retinitis pigmentosa (RP) due to mutations in exon 13 of the *USH2A* gene. QR-421a is designed to restore functional Usherin protein by using an exon skipping approach with the aim to stop or reverse vision loss in patients. QR-421a is intended to be administered through intravitreal injections in the eye and has been granted orphan drug designation in the United States and the European Union and received fast-track designation from the FDA.

About QR-1123

QR-1123 is a first-in-class investigational RNA-based oligonucleotide that was discovered and developed by Ionis Pharmaceuticals using Ionis' proprietary antisense technology for the treatment of adRP due to the P23H mutation in the *RHO* gene. The therapy aims to inhibit the formation of the mutated toxic version of the rhodopsin protein by specifically binding the mutated *RHO* mRNA. Binding of QR-1123 causes allele specific knockdown of the mutant mRNA by a mechanism called RNase H mediated cleavage without affecting the normal *RHO* mRNA. QR-1123 is intended to be administered through intravitreal injections in the eye and received IND clearance in August 2019.

About ProQR

ProQR Therapeutics is dedicated to changing lives through the creation of transformative RNA medicines for the treatment of severe genetic rare diseases such as Leber's congenital amaurosis 10, Usher syndrome and autosomal dominant retinitis pigmentosa. Based on our unique proprietary RNA repair platform technologies we are growing our pipeline with patients and loved ones in mind.

Since 2012

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 sepofarsen (QR-110) and the clinical development and the therapeutic potential thereof, statements regarding our pipeline of programs targeting inherited retinal dystrophies, statements regarding QR-421a, and the clinical development and therapeutic potential thereof, statements regarding QR-1123 and the clinical development and therapeutic potential 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. 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.

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

onducted condensed consolidated statement of 1 manetar 1 ostaon		
	September 30, 2019	December 31, 2018
•	€ 1,000	€ 1,000
Assets		
Current assets	74.700	105 500
Cash and cash equivalents	74,792	105,580
Prepayments and other receivables	2,450	1,544
Social securities and other taxes	830	1,243
Total current assets	78,072	108,367
Property, plant and equipment	2,413	1,864
Investments in associates	579	
Total assets	81, 064	110,231
Equity and liabilities		
Equity		
Equity attributable to owners of the Company	59,871	92,915
Non-controlling interests	(452)	(230)
Total equity	59,419	92,685
Current liabilities		
Borrowings	260	
Lease liabilities	890	-
Trade payables	668	135
Social securities and other taxes	15	
Pension premiums	13	7
Deferred income	984	545
Other current liabilities	8,305	7,473
Total current liabilities	11,135	8,160
Borrowings	10,510	9,386
Lease liabilities	<u> </u>	
Total liabilities	21,645	17,546
		2.,310
Total equity and liabilities	81,064	110,231

Unaudited Condensed Consolidated Statement of Profit or Loss and OCI

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

	Three month pended Septemb		Nine month p ended Septeml	
	2019	2018	ended Septemt 2019	2018
	€ 1,000	€ 1,000	€ 1,000	€ 1,000
Other income	530	2,958	1,509	4,428
Research and development costs	(11,074)	(6,297)	(32,560)	(19,972)
General and administrative costs	(2,903)	(2,579)	(8,970)	(7,900)
Total operating costs	(13,977)	(8,876)	(41,530)	(27,872)
Operating result	(13,447)	(5,918)	(40,021)	(23,444)
Finance income and expense	1,375	(74)	1,339	(664)
Results related to associates	(119)		579	_
Result before corporate income taxes	(12,191)	(5,992)	(38,103)	(24,108)
Income taxes	_	— —	(64)	(1)
Result for the period	(12,191)	(5,992)	(38,167)	(24,109)
Other comprehensive income	147	(4)	121	(15)
Total comprehensive income (attributable to owners of the				
Company)	(12,044)	(5,996)	(38,046)	(24,124)
Result attributable to				
Owners of the Company	(12,139)	(5,959)	(37,945)	(23,974)
Non-controlling interests	(52)	(33)	(222)	(135)
Troit controlling interests	(12,191)	(5,992)	(38,167)	(24,109)
Share information				
Weighted average number of shares outstanding(1)	38,912,701	33,355,327	38,902,203	32,440,220
Earnings per share attributable to the equity holders of the				
Company (expressed in Euro per share)				
Basic loss per share(1)	(0.31)	(0.18)	(0.98)	(0.74)
Diluted loss per share(1)	(0.31)	(0.18)	(0.98)	(0.74)

⁽¹⁾ For this period presented in these financial statements, the potential exercise of share options is not included in the diluted earnings per share calculation as the Company was loss-making in all periods. Due to the anti-dilutive nature of the outstanding options, basic and diluted earnings per share are equal in this period.

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

	Attributable to owners of the Company								
	Number of shares	Share Capital € 1,000	Share Premium	Equity Settled Employee Benefit Reserve	Translation Reserve	Accumulated Deficit	Total € 1,000	Non- controlling interests	Total Equity € 1,000
Balance at January 1, 2018	36,425,014	1,457	€ 1,000 148,763	€ 1,000 8,377	€ 1,000 136	€ 1,000 (119,370)	39,363	€ 1,000 (38)	39,325
Result for the period						(23,974)	(23,974)	(135)	(24,109)
Other comprehensive income	_	_	_	_	(15)		(15)	_	(15)
Recognition of share-based					` /		, ,		` /
payments	_	_		2,245	_	_	2,245	_	2,245
Issuance of ordinary shares	6,612,500	265	84,032	_	_	_	84,297	_	84,297
Share options exercised	_	_	659	_	_	_	659	_	659
Balance at September 30, 2018	43,037,514	1,722	233,454	10,622	121	(143,344)	102,575	(173)	102,402
Balance at January 1, 2019	43,149,987	1,726	235,744	10,780	108	(155,443)	92,915	(230)	92,685
Result for the period						(37,945)	(37,945)	(222)	(38,167)
Other comprehensive income	_	_			121		121	_	121
Recognition of share-based									
payments	_	_	_	4,614	_	_	4,614	_	4,614
Share options lapsed	_	_	_	(33)		33	_	_	_
Share options exercised			166	(115)		115	166		166
Balance at September 30, 2019	43,149,987	1,726	235,910	15,246	229	(193,240)	59,871	(452)	59,419

Unaudited Condensed Consolidated Statement of Cash Flows

	Three month p ended Septeml		Nine month p ended Septeml	
	2019	2018	2019	2018
Cook floors from an arresting a stimition	€ 1,000	€ 1,000	€ 1,000	€ 1,000
Cash flows from operating activities	(12.101)	(F 003)	(20.167)	(24.100)
Net result	(12,191)	(5,992)	(38,167)	(24,109)
Adjustments for:	FOC	242	1 5 40	725
— Depreciation	506		1,543	725
— Share-based compensation	1,226	734	4,614	2,245
— Financial income and expenses	(1,375)	74	(1,339)	664
— Results related to associates	119	<u> </u>	(579)	(15)
— Net foreign exchange gain / (loss)	148	(4)	122	(15)
Changes in working capital	2,718	656	1,744	1,074
Cash used in operations	(8,849)	(4,290)	(32,062)	(19,416)
Corporate income tax paid	_	1	(64)	_
Interest received	90	32	176	25
Interest paid	(13)		(64)	_
Net cash used in operating activities	(8,772)	(4,257)	(32,014)	(19,391)
Cash flow from investing activities				
Purchases of intangible assets	<u> </u>	_	_	_
Purchases of property, plant and equipment	(32)	(99)	(341)	(285)
Net cash used in investing activities	(32)	(99)	(341)	(285)
ivet cash used in investing activities	(32)	(99)	(341)	(203)
Cash flow from financing activities				
Proceeds from issuance of shares, net of transaction costs	_	84,295	_	84,295
Proceeds from exercise of share options	2	637	166	660
Proceeds from borrowings	_	_	_	101
Proceeds from convertible loans	_	115	690	430
Repayment of lease liability	(290)		(861)	
Net cash (used in)/generated by financing activities	(288)	85,047	(5)	85,486
Net increase/(decrease) in cash and cash equivalents	(9,092)	80,691	(32,360)	65,810
	1 400		1.550	(100)
Currency effect cash and cash equivalents	1,420	57	1,572	(193)
Cash and cash equivalents, at beginning of the period	82,464	32,968	105,580	48,099
Cash and cash equivalents at the end of the period	74,792	113,716	74,792	113,716

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