

Quantum Genomics Corp.

(ALQGC.PA – Paris)

REFRESH - First Patient Enrolled

Based on our DCF model and a 15% discount rate, Quantum Genomics is valued at approximately €16.00 per share. Our model applies a 50% probability of ultimate approval and commercialization for firibastat in hypertension and a 15% probability for heart failure. The model includes contributions from the United States, European Union, Latin America, Asia, Canada, South Korea and Oceania.

Current Price (7/9/2021)

€3.76

Valuation

€16.00

OUTLOOK

Quantum Genomics is developing lead candidate firibastat for difficult-to-treat and resistant hypertension (HTN) and heart failure (HF). The BAPAI class drug blocks the conversion of A2 to A3 thereby preventing A3 binding with AT1 receptors which controls blood pressure via a triple mechanism of action.

Quantum is conducting two Ph3 trials for HTN in difficult-to-treat and resistant populations (FRESH & REFRESH), a Ph2b trial in heart failure (QUORUM) and a Ph1 in 1x/day HTN. The first patient in the REFRESH Ph3 safety study for HTN was enrolled in July 2021.. The timeline also anticipates a HTN NDA filing in 2023.

HTN is a highly prevalent disease and a material portion of this population does not have the disease under control. Firibastat seeks to address this unmet need via a differentiated pathway complementary to currently approved therapies.

Our valuation assumes a 2023 regulatory submission in multiple geographies and subsequent commercialization. Partner Biolab is expected to pursue approval in select Latin American countries and commercialize throughout that region in 2023 and 2024 respectively.

SUMMARY DATA

52-Week High	5.68
52-Week Low	2.18
One-Year Return (%)	40.8
Beta	0.89
Average Daily Volume (sh)	173,411

Shares Outstanding (mil)	26.8
Market Capitalization (€mil)	101
Short Interest Ratio (days)	N/A
Institutional Ownership (%)	6.42
Insider Ownership (%)	10.3

Annual Cash Dividend	\$0.00
Dividend Yield (%)	0.00

5-Yr. Historical Growth Rates	
Sales (%)	N/A
Earnings Per Share (%)	N/A
Dividend (%)	N/A

P/E using TTM EPS	N/A
P/E using 2020 Estimate	N/A
P/E using 2021 Estimate	N/A

Zacks Rank	N/A
------------	-----

Risk Level	Above Average
Type of Stock	Small-Growth
Industry	Med-Biomed/Gene

ZACKS ESTIMATES

Revenue

(In millions of EUR)

	Q1	Q2	Q3	Q4	Year
	(Mar)	(Jun)	(Sep)	(Dec)	(Dec)
2019	€0.0 A	€0.3 A	€0.0 A	€0.1 A	€0.4 A
2020	€0.0 A	€0.4 A	€0.0 A	€1.9 A	€2.3 A
2021	€0.0 E	€6.0 E	€0.0 E	€6.0 E	€12.0 E
2022					€15.0 E

Earnings per Share

	Q1	Q2	Q3	Q4	Year
2019	€0.00 A	-€0.24 A	€0.00 A	-€0.27 A	-€0.50 A
2020	€0.00 A	-€0.25 A	€0.00 A	-€0.25 A	-€0.43 A
2021	€0.00 E	-€0.19 E	€0.00 E	-€0.26 E	-€0.44 E
2022					-€0.20 E

WHAT'S NEW

REFRESH First Patient Enrolled

Quantum [announced](#) on July 8, 2021 that the first patient had been enrolled in its pivotal Phase III REFRESH trial of once-daily firobostat in difficult-to-treat and resistant hypertension. REFRESH is the last step in an FDA-endorsed development plan needed before submission for market authorization in 4Q:23. The study is being conducted with partners DongWha for the South Korean market and Orient Europharma for Southeast Asia, Australia and New Zealand. The multicenter, multinational study is targeting total enrollment of 750 patients with difficult to treat or resistant hypertension, defined by persistent hypertension despite administration of two antihypertensive drugs at maximum tolerated doses and three antihypertensive drugs including a diuretic at maximum tolerated doses, respectively. Quantum anticipates participation of 96 study sites across Europe, Canada, US, Taiwan and South Korea. Efficacy results and six month safety results are expected in mid-2023.

Phase III REFRESH Launched

On January 18, 2021, Quantum [announced](#) the launch of REFRESH, a Phase III pivotal trial of once daily firobostat in difficult-to-treat¹ hypertension and resistant² hypertension, a key milestone in the pursuit of global commercialization for Quantum's lead candidate. The study is part of an overall Phase III evaluation of firobostat. If successful, the once-daily administration will provide additional convenience and compliance compared to the twice-daily regimen being investigated in the FRESH trials. The goal of REFRESH is to assess both long-term safety and three-month efficacy in the once-daily dose. The launch of the trial will not impact the anticipated timeline for the filing of firobostat which is expected in 2023.

Dosing for REFRESH will be 1000 mg firobostat, administered once per day to individuals with treatment-resistant hypertension. The total number of target sites has not yet been determined as Quantum is still finalizing territories and associated partnerships. Some clinical sites are already being used in other Quantum trials easing site selection, while others, in new geographies, will need to be established.

For the first three months of treatment, patients in REFRESH will receive 1000 mg firobostat once-daily in addition to their current regimen. The primary endpoint is reduction in systolic automated office blood pressure from baseline. Following the three-month efficacy evaluation, treatment will continue with follow-up for six months. 100 patients will receive follow up exams at 12 months to assess long-term safety. As hypertension often requires chronic use of medication, the long-term safety data generated in REFRESH is necessary for submission of a New Drug Application (NDA).

Key Milestone for Quantum: QUORUM Study Results Announcement

[Topline](#) results for the [QUORUM](#) study will be presented at the [European Society of Cardiology](#) (ESC) Congress being held from August 27 – 30, in a virtual format. The communication of the data will take place on August 27 at 10:30 am Central European Summer Time (CEST). QUORUM (Firobostat Or Ramipril after Acute Myocardial infarction to prevent left ventricular dysfunction) is a multi-center, multinational, randomized, double-blind, active-controlled trial designed to assess the efficacy and the safety of firobostat compared to ramipril. Our valuation model attributes approximately 14% of Quantum's valuation to success of firobostat in post-myocardial infarction heart failure (MI HF). Strong results from this trial could potentially support even greater value for post MI HF indication compared with the hypertension target given our estimates of a higher penetration rate into this smaller, but less competitive market.

Quantum representatives will also be [presenting](#) other information at ESC. On August 28th, [Dr. Catherine Llorens-Cortes](#), one of the inventors of BAPAI, will be presenting the details of a preclinical study of QGC606 in heart failure after myocardial infarction. The title of the presentation is *Comparison of QGC606, a novel orally active brain-penetrating aminopeptidase A inhibitor prodrug with firobostat and ramipril for treating heart failure following myocardial infarction*, which will be offered at the Late Breaking Basic & Translational Science session.

¹ Hypertension that is not controlled despite two antihypertensive classes, including a diuretic, at maximum tolerated doses.

² Hypertension that is not controlled despite treatment with at least three antihypertensive classes, including a diuretic, at maximum tolerated doses.

Other Events

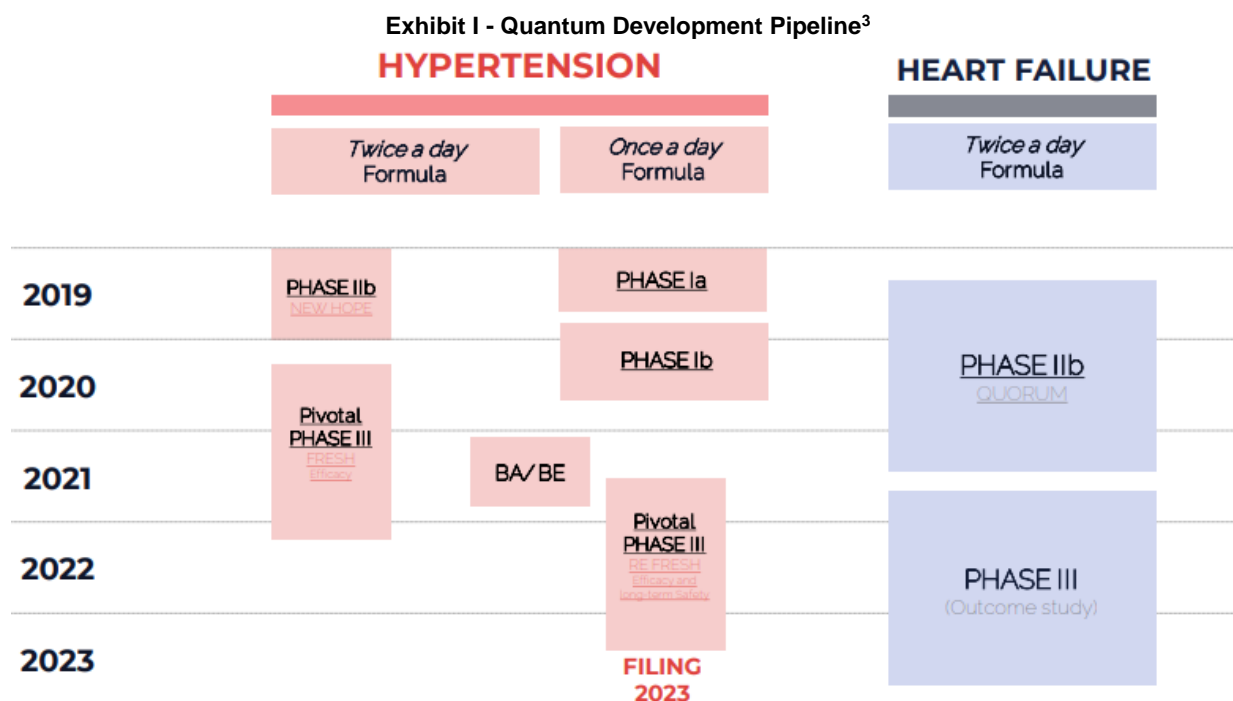
Since our previous update in April, Quantum has provided several updates related to partners, financing, publications and anticipated conference presentation in addition to advancing the REFRESH trial. On April 20, the company [announced](#) the end of its collaboration with Qilu in China. The two companies were not able to reconcile the objectives for firibastat development in the Chinese market and aborted their agreement to commercialize the region. Quantum has recovered development rights and is now opening discussions with other potential collaborators. No risk of litigation or penalties were identified by Quantum Genomics.

In late April, Quantum secured €3 million in non-dilutive financing from government-backed, low interest loans. Half comes from BNP at a 0.25% interest rate and half from a research and development loan from BPIfrance at a 0.72% rate. The first loan comes due in one year and the second has a seven year maturity. As of the end of 2020, Quantum held €28.5 million in cash on the balance sheet and is expected to burn about €1 million per month. The loans should extend the funding of operations by about a quarter at a very favorable cost of capital.

In May, Quantum [reported](#) the publication of a scientific article in Biomedicine & Pharmacotherapy entitled "[Effects of firibastat in combination with enalapril and hydrochlorothiazide on blood pressure and vasopressin release in hypertensive DOCA-salt rats.](#)" The paper examined oral administration of firibastat in hypertensive rats. Firibastat efficacy was evaluated in combination with enalapril and hydrochlorothiazide. Administration of oral firibastat in triple combination produced a significant decrease in blood pressure whereas enalapril and hydrochlorothiazide alone had minimal impact. The triple combination also performed better than using firibastat alone. The animal study was supportive of firibastat being used in combination therapy for blood pressure control in difficult to treat or resistant patient populations.

Pipeline

Quantum's pipeline consists of one drug, firibastat, with various formulations pursuing multiple indications. It is being investigated in difficult-to-treat and resistant hypertension, heart failure, renal failure also in once per day formulations. The candidate is currently in one Phase III trial for hypertension, one Phase II trial for heart failure and Phase I trials for QD formulation hypertension and renal failure.



³ Source: Quantum Genomics March 2021 Corporate Presentation

Milestones

- FRESH first patient enrolled – July 2020
- €20 million private placement – December 2020
- Launch REFRESH study – January 2021
- Orient EuroPharma Equity Stake – February 2021
- First patient enrolled in REFRESH study – July 2021
- QUORUM Phase II results, presented at ESC – August 2021
- FRESH study results – 4Q:21
- With supportive data launch Ph3 outcome study in HF – 2021/2022
- Topline results for REFRESH study – Mid-year 2023

Summary

Quantum Genomics continues to make headway in the clinic, reaching its latest milestone of enrolling its first patient in the REFRESH study. Quantum is building a global market for firibastat with development and commercialization partnerships around the world. Additional opportunity remains for other deals in countries such as Japan, India, North America, China and unpenetrated areas in Europe. Despite the Qilu setback, there are plenty of other catalysts that are expected in the near future. The most prominent is the presentation of topline data from the Phase IIb QUORUM trial, which will provide proof of concept for the use of firibastat in heart failure. New sources of capital and cash reserves of almost €29 million suggest a runway beyond 2022. We expect additional funds in coming quarters which will further support development programs and perhaps obviate the need for further capital raises. We maintain our target price of €16 per share.

PROJECTED FINANCIALS

Quantum Genomics Corp. - Income Statement⁴

Quantum Genomics Corp.	1H A	2H A	2019 A	1H A	2H A	2020 A	2021 E	2022 E
Total Revenues (€,000)	€ 285	€ 76	€ 361	€ 377	€ 1,884	€ 2,262	€ 12,000	€ 15,000
YOY Growth	333%	1304%	407%	32%	2368%	526%	431%	25%
Raw Materials & Supplies	€ 0	€ 89	€ 89	€ 269	€ 736	€ 1,005	€ 90	€ 90
Other Purchases & Expenses	€ 3,946	€ 3,853	€ 7,799	€ 4,837	€ 7,466	€ 12,303	€ 22,200	€ 18,000
Taxes	€ 9	€ 2	€ 10	€ 12	€ 9	€ 20	€ 20	€ 20
Wages & Salaries	€ 916	€ 814	€ 1,730	€ 813	€ 719	€ 1,532	€ 2,000	€ 2,100
Social Security Charges	€ 660	€ 385	€ 1,045	€ 321	€ 475	€ 797	€ 1,200	€ 1,250
Depreciation & Provisions	€ 7	€ 299	€ 306	€ 5	€ 320	€ 325	€ 306	€ 310
Other Expenses	€ 71	€ 69	€ 140	€ 75	€ 62	€ 137	€ 140	€ 150
Income from operations	(€ 5,324)	(€ 5,436)	(€ 10,760)	(€ 5,955)	(€ 7,902)	(€ 13,858)	(€ 13,956)	(€ 6,920)
Operating Margin	-1870%	-7119%	-2979%	-1578%	-419%	-613%	-116%	-46%
Financial Income	€ 6	€ 5	€ 11	€ 3	€ 2	€ 6	€ 6	€ 6
Financial Expenses	€ 0	€ 0	€ 0	€ 5	€ 0	€ 11	€ 0	€ 0
Exceptional Items	€ 304	(€ 181)	€ 123	€ 17	€ 0	€ 178	€ 0	€ 0
Pre-Tax Income	(€ 5,014)	(€ 5,611)	(€ 10,626)	(€ 5,941)	(€ 7,900)	(€ 13,684)	(€ 13,950)	(€ 6,914)
Provision for Income Tax	(€ 886)	(€ 662)	(€ 1,547)	(€ 860)	(€ 1,185)	(€ 2,148)	(€ 2,093)	(€ 1,037)
Tax Rate	17.7%	11.8%	14.6%	15.0%	15.0%	15.7%	15.0%	15.0%
Net Income	(€ 4,129)	(€ 4,950)	(€ 9,078)	(€ 5,081)	(€ 6,715)	(€ 11,537)	(€ 11,858)	(€ 5,877)
Reported EPS	(€ 0.24)	(€ 0.27)	(€ 0.50)	(€ 0.25)	(€ 0.25)	(€ 0.43)	(€ 0.44)	(€ 0.20)
Basic Shares Outstanding	16,886	18,065	18,065	20,500	26,712	26,712	27,000	30,000

Source: Company Filing // Zacks Investment Research, Inc. Estimates

⁴ Historical financial statement information presents data as originally reported.

HISTORICAL STOCK PRICE

Quantum Genomics Corp. – Share Price Chart (€)⁵



⁵ Source: barchart.com

DISCLOSURES

The following disclosures relate to relationships between Zacks Small-Cap Research (“Zacks SCR”), a division of Zacks Investment Research (“ZIR”), and the issuers covered by the Zacks SCR Analysts in the Small-Cap Universe.

ANALYST DISCLOSURES

I, John Vandermosten, hereby certify that the views expressed in this research report accurately reflect my personal views about the subject securities and issuers. I also certify that no part of my compensation was, is, or will be, directly or indirectly, related to the recommendations or views expressed in this research report. I believe the information used for the creation of this report has been obtained from sources I considered to be reliable, but I can neither guarantee nor represent the completeness or accuracy of the information herewith. Such information and the opinions expressed are subject to change without notice.

INVESTMENT BANKING AND FEES FOR SERVICES

Zacks SCR does not provide investment banking services nor has it received compensation for investment banking services from the issuers of the securities covered in this report or article.

Zacks SCR has received compensation from the issuer directly or from an investor relations consulting firm engaged by the issuer for providing non-investment banking services to this issuer and expects to receive additional compensation for such non-investment banking services provided to this issuer. The non-investment banking services provided to the issuer includes the preparation of this report, investor relations services, investment software, financial database analysis, organization of non-deal road shows, and attendance fees for conferences sponsored or co-sponsored by Zacks SCR. The fees for these services vary on a per-client basis and are subject to the number and types of services contracted.

POLICY DISCLOSURES

This report provides an objective valuation of the issuer today and expected valuations of the issuer at various future dates based on applying standard investment valuation methodologies to the revenue and EPS forecasts made by the SCR Analyst of the issuer’s business. SCR Analysts are restricted from holding or trading securities in the issuers that they cover. ZIR and Zacks SCR do not make a market in any security followed by SCR nor do they act as dealers in these securities. Each Zacks SCR Analyst has full discretion over the valuation of the issuer included in this report based on his or her own due diligence. SCR Analysts are paid based on the number of companies they cover. SCR Analyst compensation is not, was not, nor will be, directly or indirectly, related to the specific valuations or views expressed in any report or article.

ADDITIONAL INFORMATION

Additional information is available upon request. Zacks SCR reports and articles are based on data obtained from sources that it believes to be reliable, but are not guaranteed to be accurate nor do they purport to be complete. Because of individual financial or investment objectives and/or financial circumstances, this report or article should not be construed as advice designed to meet the particular investment needs of any investor. Investing involves risk. Any opinions expressed by Zacks SCR Analysts are subject to change without notice. Reports or articles or tweets are not to be construed as an offer or solicitation of an offer to buy or sell the securities herein mentioned.