



Financial Report

Second Quarter - Fiscal Year 2022

April 30, 2022

VALEO PHARMA INC.

Management's Discussion and Analysis for the three and six-month periods ended April 30, 2022

MANAGEMENT'S RESPONSIBILITY FOR FINANCIAL REPORTING

The following is Management's Discussion and Analysis ("MD&A") of the financial condition and operating results of Valeo Pharma Inc. ("Valeo" or the "Corporation") for the quarter ended April 30, 2022. This document should be read in conjunction with the unaudited interim condensed consolidated financial statements and notes thereto for the quarter ended April 30, 2022, which have been prepared in accordance with *International Financial Reporting Standards*. All amounts herein are expressed in thousands of Canadian dollars (unless otherwise indicated) except for share and per share amounts. All other currencies are in thousands. This discussion and analysis document was prepared by management from information available as at June 14, 2022. Further information about Valeo Pharma Inc., including the Annual Information Form, is available online on SEDAR at www.sedar.com.

Non-IFRS Financial Measures

The non-IFRS measures included in this MD&A are not recognized measures under IFRS and do not have a standardized meaning prescribed by IFRS and may not be comparable to similar measures presented by other issuers. When used, these measures are defined in such terms as to allow the reconciliation to the closest IFRS measure. These measures are provided as additional information to complement those IFRS measures by providing further understanding of our results of operations from our perspective. Accordingly, they should not be considered in isolation nor as a substitute for analysis of our financial information reported under IFRS. Despite the importance of these measures to management in goal setting and performance measurement, we stress that these are non-IFRS measures that may have limits in their usefulness to investors.

We use non-IFRS measures, such as EBITDA and Adjusted EBITDA to provide investors with a supplemental measure of our operating performance and thus highlight trends in our core business that may not otherwise be apparent when relying solely on IFRS financial measures. We also believe that securities analysts, investors and other interested parties frequently use non-IFRS measures in the valuation of issuers. We also use non-IFRS measures in order to facilitate operating performance comparisons from period to period, prepare annual operating budgets, and to assess our ability to meet our future debt service, capital expenditure and working capital requirements.

The definition and reconciliation of EBITDA and Adjusted EBITDA used and presented by the Corporation to the most directly comparable IFRS measures follow below:

EBITDA is defined as net (loss)/income adjusted for income tax, depreciation of property and equipment, amortization of right of use asset, amortization of intangible assets, interest on short and long-term debt and other financing costs, interest income, licensing revenue and changes in fair values of derivative financial instruments. Management uses EBITDA to assess the Corporation's operating performance.

Adjusted EBITDA is defined as EBITDA adjusted for, as applicable, 1) share based compensation and other warrants or options issuance costs, 2) settlement for contract terminations such as severance for executives, or penalties for early termination of multi-year contracts, 3) impairment of intangible asset, 4) charges related to product recalls or contractual inventory returns not related to product shelf life, 5) listing fees not related to share issuance, 6) non-recurrent product launches staff recruitment fees and 7) specific material non-recurrent special provisions. We use Adjusted EBITDA as a key metric in assessing our business performance when we compare results to budgets, forecasts and prior years. Management believes Adjusted EBITDA is a more accurate measure of cash flow generation than, for example, cash flow from operations, particularly because it removes cash flow fluctuations caused by unusual changes in working capital. A reconciliation of net (loss)/income to EBITDA (and Adjusted EBITDA) are presented later in this document.

Use of Estimates and Judgements

The preparation of unaudited interim condensed consolidated financial statements requires management to undertake several judgements, estimates and assumptions about recognition and measurement of assets, liabilities, income and expenses. The actual results may differ from these judgements and estimates. These estimates and judgements are based on management's best knowledge of the events or circumstances and actions the Corporation may take in the future. The estimates are reviewed on an ongoing basis. Information about the significant judgements, estimates and assumptions that have the most significant effect on the recognition and measurement of assets, liabilities, income and expenses are discussed in Note 3 of the Corporation's 2021 audited annual consolidated financial statements and are still applicable for the six-month period ended April 30, 2022.

Cautionary note regarding forward-looking statements

This MD&A may contain some forward-looking information as defined under applicable Canadian securities laws. Forward looking information can generally be identified using forward-looking terminology such as "may", "anticipate", "expect", "intend", "estimate", "continue" or similar terminology. Forward looking information is subject to various known and unknown risks and uncertainties, many of which are beyond the ability of the Corporation to control or predict, that may cause the Corporation's actual results or performance to be materially different from actual results and are developed based on assumptions about such risks and other factors set out herein.

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GLOSSARY TERMS

Calendar & Financial		Corporate & Operations	
COGS	Cost of Goods Sold (or Cost of Sales)	Biosimilar	Biologic drug that is highly similar to a biologic drug.
G&A	General and Administrative	BU	Business Unit defined as Commercial Unit focussing on specific therapeutic areas
HO	Head Office	COVID-19	Mild to severe respiratory illness caused by a coronavirus
IR	Investors Relation	CTA	Clinical Trial Application with Health Canada
S&M	Sales and Marketing	DIN	Drug Identification Number
SBC	Share-Based Compensation	FDA	United States Food and Drug Administration
MA & Reg	Medical Affairs, Quality Assurance and Regulatory	FSE	Frankfurt Stock Exchange
FY-22	Fiscal Year 2022	GDUFA	Generic Drug User Fee Act in the USA
FY-21	Fiscal Year 2021	GPO	Group Purchase Organization
Q2-22	Second quarter FY-22	HC	Health Canada
Q1-22	First quarter FY-22	ICS	Inhaled Corticosteroid
Q4-21	Fourth quarter FY-21	INESSS	Quebec's Institut National d'Excellence en Santé et Services Sociaux
Q3-21	Third quarter FY-21	KAM	Key Account Manager
Q2-21	Second quarter FY-21	KOL	Key Opinion Leader
Q1-21	First quarter FY-21	LABA	Long-Acting Beta2 Agonist
Q4-20	Fourth quarter FY-20	LAMA	Long-Acting Muscarinic Antagonist
Q3-20	Third quarter FY-20	LMWH	Low Molecular Weight Heparin
QoQ	Current year quarterly results vs last year's quarterly results	MHI	Montreal Heart Institute
YE-21	Year-end 2021, October 31, 2021	NBRx	New to Brand Prescriptions
YTD	Year to date	NDS	New Drug Submission with Health Canada
YoY	Current FY results vs last FY results	OTCQB	U.S. over-the-counter venture market
W/C	Working Capital, defined as short-term assets less short-term liabilities	pCPA	pan-Canadian Pharmaceutical Alliance
		PD	Parkinson's Disease
		PLA	Product listing agreement
		PMPRB	Patented Medicine Prices Review Board
		RAMQ	Régie de l'assurance maladie du Québec
		TSX	Toronto Stock Exchange
		SKU's	Stock Keeping Units
		VPI	Wholly owned subsidiary of Valeo focussed on the commercialization of generic products

OVERVIEW OF THE BUSINESS AND BUSINESS STRATEGY

The Corporation is a specialty pharmaceutical company which sources, acquires or in-licenses branded products for sale in Canada. Preferences in acquisition are for innovative products, already approved in other territories and addressing major unmet medical needs. Valeo's business model consists of providing all the services required to register, to reimburse and to commercialize the acquired or in-licensed pharmaceutical products in Canada. Within this kind of in-licensing agreements, products may require up-front, regulatory and/or commercial stage milestone payments and all require regulatory approval from *Health Canada* prior to commercialization.

Valeo's business objective is to become a leading Canadian healthcare Corporation by focusing on the commercialization of innovative products in predefined strategic therapeutic areas. The Corporation has two wholly owned subsidiaries: VPI Pharmaceuticals Inc., located in Kirkland, Québec, specialized in hospital generic products and Valeo Pharma Corp. located in the United States.

In March 2021, Valeo closed a significant agreement with Novartis Pharmaceutical Canada Inc. ("Novartis") (See "Corporate Highlights") for the Canadian rights to Enerzair®Breezhaler® ("Enerzair") and Aetectura®Breezhaler® ("Aetectura"), two innovative asthma products. This material event did trigger a major transformation of the Corporation and its commercial activities into (2) distinct Business Units for our branded products, while generic products mainly positioned on hospital markets were put under the umbrella of an hospital generics division, all supported by head office functions. The first BU focuses on the Respiratory therapeutic area with the commercialization of the licenced asthma products, while the second BU focuses mainly on Thrombosis, Neurology, Oncology, with the commercialization of Redesca™, Onstryv®, Yondelis® and M-Eslon® as its main brands. Therapeutic areas are selected based on market potential (size and growth prospects), competitive landscape, and resource requirements needed to reach the target audience and execute our commercialization strategy.

As of the date of this document, the Corporation has about 100 full time employees including a team of 70 pharmaceutical representatives, sales professionals, and medical science liaisons staff. While expanding the field team, several key executive positions were also filled in order to strengthen the leadership team. A SVP Scientific and Medical Affairs, a VP HR & Talent Management, as well as new Business Unit

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heads, coming all from multinational companies joined Valeo. They all bring years of experience within the pharmaceutical industry, either in commercial, medical, and digital transformation leadership roles.

Valeo is well equipped to distribute on its own all our products. With a 20,767 square feet dedicated warehousing space located in Kirkland, Quebec we can handle all the inventory requirements for Canada. Valeo's facility includes warehouse space, three licensed narcotics vaults, the capability to handle cold chain requirements and shipping needs. Valeo also operates a sophisticated SAP enterprise resource planning system and possesses the in-house expertise to handle all activities associated with regulatory, quality control, supply chain, medical information, and pharmacovigilance. The ability to handle such a broad range of activities has been a key factor in our successful in-licensing activities and acquisition of third-party product rights for Canada.

Strategic product launches were performed by Valeo in 2021, with Redesca™ the first LMWH Biosimilar introduced in April 2021, and of Enerzair and Aetectura two innovative asthma drugs introduced in June 2021. Since then, the Valeo's team has tirelessly pursued an objective to execute on our strategic plan with the rigour, the focus, and the reactivity necessary to efficiently position each of these products on their respective market. Assembling the high performing team, negotiating the private and public coverage of all three drugs, launching effective marketing campaign, seeking medical collaboration with top clinicians in the country have been amongst the key activities developed in support of these launches. We expect the Respiratory and Specialty products BU to materially impact our financial performance over the coming months. The significant revenue growth seen in 2022 fiscal year is a clear testament of the transformative impact these products have on the corporation, and they all three will contribute for many years ahead until they reach their full potential.

At the end of Q2-22, Valeo's product portfolio included eleven (11) commercial stage products.

BRANDS	Indications	Partners	Regulatory, Commercial Status, and other important information
Respiratory Business Unit			
Enerzair® Breezhaler® (Commercial Agreement)	LABA/LAMA/ICS fixed triple dose asthma drug.	Novartis Pharmaceuticals Canada Inc. ("Novartis")	<ul style="list-style-type: none"> Commercialization & Supply Agreement in Q2-21. Public reimbursement secured across Canada, with the last process currently ongoing for British Columbia. Private insurance coverage exceeds 90%.
Aetectura® Breezhaler® (Commercial Agreement)	LABA/ICS dual combination asthma drug.		<ul style="list-style-type: none"> Canadian maintenance asthma market estimated at \$630M and expected to grow annually by 2-3% (Source: IQVIA, 2021). Commercial launch in June 2021 by a dedicated team of 60 sales professionals. Both products are selling in all provinces and have generated 8,247 Rx from 447 Medical doctors.
Specialty Products Business Unit			
Redesca™ (Distribution)	LMWH – Anticoagulant biosimilar used to treat and prevent deep vein thrombosis and pulmonary embolism.	Shenzhen Techdow Pharmaceuticals Co., Ltd.	<ul style="list-style-type: none"> Commercialized since April 2021 and supported by a dedicated team of key account managers across the country. Canadian annual LMWH market estimated at \$180M (Source: IQVIA) The product has 8+ years of proven in-market safety internationally and more than 150 million patient days treated in Europe alone. Provincial reimbursement secured in all Canadian provinces. Private insurance coverage exceeds 90%. Awarded several hospital contracts in AB and QC while the originator has started to be delisted. 35% of the Enoxaparin has been transferred to biosimilars and Redesca is owning 54% of this market.
Onstryv® (License)	Idiopathic PD as an add-on for patients on stable dose of Levodopa (L-dopa) alone or in combination with other drugs, to help with "off" episodes.	Zambon S.p.A.	<ul style="list-style-type: none"> Marketed since Q3-19. INESSS positive recommendation granted in February 2020. Ongoing engagement process with pCPA to negotiate the public reimbursement in Quebec.
M-Esolon (Distribution)	Extended-release morphine sulphate used for pain management.	Ethypharm Inc.	<ul style="list-style-type: none"> The Corporation is distributing the product and is recording sales on a gross basis.
Yondelis® (License)	Soft tissue sarcoma	PharmaMar S.A.	<ul style="list-style-type: none"> Marketed since August 2020.
Hesperco™	Bioflavonoid antioxidant used for immune support	Co-developed with Ingenew Pharma Inc. ("Ingenew")	<ul style="list-style-type: none"> Marketed since October 2020 on-line and available on Amazon Canada and in Loblaw's retail pharmacies. Results of a clinical trial conducted by The MHI has confirmed the merits of Hesperco for helping reduce Covid-19 related symptoms.
Ametop™ Gel 4%	For skin Anesthesia	Alliance Pharma	<ul style="list-style-type: none"> Marketed since Q4-21. Used mainly in hospital prior to venepuncture or venous cannulation.

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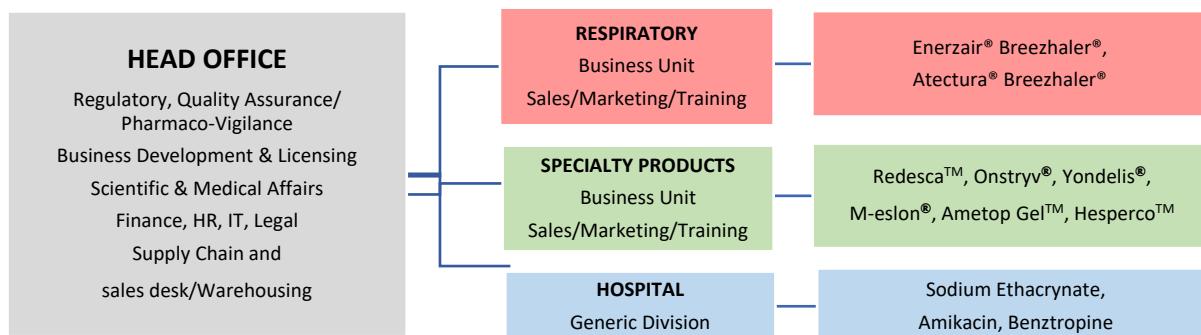
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Hospital Generic Division			
Benztropine (Distribution)	VPI-Anticholinergic agent used for the treatment of Parkinson disease	Asia/Pacific Generic Manufacturer	<ul style="list-style-type: none"> Marketed in Canada since Q4-18.
Ethacrynat Sodium	Loop diuretic for high blood pressure and associated swelling	Owned by Valeo worldwide except for Italy	<ul style="list-style-type: none"> Marketed in Canada since Q3-18 and in the United States since Q4-21 via a US-based distribution partner.
Amikacin	Injectable Antibiotic	European Generic Manufacturer	<ul style="list-style-type: none"> Approved by Health Canada in 2020. Commercialization has started in Q3-21.

Valeo continues to search for innovative products within its targeted areas of focus and maintains active business development activities to achieve this goal. Our experienced management team has a long and proven track record of successfully sourcing, registering, and commercializing drugs in a variety of therapeutic areas at all stages of their life cycle in Canada.

The recent creation of the two Business Units ("BU") and the ongoing integration of a dedicated sales team to support the respective commercial efforts of key products within our portfolio will create significant operating leverage over the coming years as we aim to add other strategic assets to each BU and take full advantage of our new corporate structure and commercial platform. We also equipped both BU and the Medical team with a digital platform (CRM) enabling them to reach out to customers remotely either to perform e-detailing, webinar, lunch & learns or even to provide remotely samples and training kits to health care practitioners ("HCP's"). While this technology has helped enrich and personalize the customer relationship, Valeo has favored face-to-face interactions with healthcare professionals especially since the Covid-10 restrictions have been progressively removed.

The following presents a summary of our new corporate and commercial structure which has been fully operational since the later part of FY-21.



Respiratory Business Unit

The Respiratory BU has been created to take full advantage of market opportunities for two innovative asthma therapies, Enerzair® and Aetectura®, licensed-in from Novartis in March 2021. Both products offer compelling therapeutic benefits over the current standard of care and are now available across all Canadian provinces and territories. Enerzair® and Aetectura® have helped established Valeo as a key player in the large, established, and growing asthma market. Our Respiratory BU is operational since the end of Q4-21 and is composed of a BU head, regional sales directors, Specialist and Primary Care representatives visiting and detailing Enerzair and Aetectura on a core target of Respiratory specialists and General practitioners representing more than 80% of the total scripts in Asthma. On top of that, a dedicated medical team composed of Medical Science Liaisons and a Medical advisor is supporting key interactions with Asthma Key opinion leaders across the country and helping to grow our products awareness.

Close to 4 million Canadians are living with asthma, a serious health issue affecting all age groups and 39% of asthma patients remain uncontrolled, despite available medications. This is primarily due to low adherence, treatment misuse and poor inhaler technique. The market opportunities for innovative medicines in asthma are significant and Valeo is well positioned to take full advantage of the favorable market dynamics. Over the last two years, the Covid-19 pandemic has dramatically impacted the way Asthma is currently managed by HCP's. With the number of in-person medical visit to patients being substantially reduced, much less opportunities to assess the level of asthma control, have presented themselves to physicians. As a result of that, less patients have been subject to treatment review and adaptation. We expect now, with restrictions being relieved, to see how much the number of uncontrolled asthma patients has increased during this period of time.

The acquisition of market data -both sales and prescriptions- to support and monitor our commercialization performance as well as to identify market opportunities, set the stage for monitoring significant quarterly sequential market gains in FY-22 and beyond. Our Q1-22 results were showing good progress over the prior Q4-21 quarter and significant positive variance over last year's Q1-21 results. Our Q2-22 results confirm this trend.

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Specialty Products Business Unit

The Specialty Product BU has been created to help Valeo derive maximum benefits from the commercialization of Redesca™ and other branded products.

REDESCA™ – a transformative product for Valeo.

Following the HC approval of Redesca™ in December 2020, Valeo has successfully launched the product in Q2-21. Due to the size of the commercial opportunity, the growing experience of our dedicated KAM team and the innovative approach to GPO's tenders, we have experienced rapid and growing demand for Redesca™ and a meaningful contribution to our quarterly results. Redesca™ is now largely covered by private insurance companies as well as by all provincial public jurisdictions, including BC who started to cover as of March 22, 2022.

Following a solid start in Q3-21 which included significant pipeline fill across Canada by several retailers, we expect rapid market share gains for Redesca™ as many hospitals adopt LMWH biosimilars as an alternative to more expensive innovator products.

The LMWH Canadian market is estimated at \$180 M and includes 3 major molecules.

- The Enoxaparin market (the “Primary Market”) is estimated at \$60M annually and includes 4 players (Lovenox – and 3 biosimilars to Lovenox, including Redesca™).
- The rest of the market (the “Secondary Market”) is composed of 2 other molecules – Dalteparin and Tinzaparin together representing sales of \$120 million annually. No biosimilar have been approved for these molecules and none are expected to enter the LMWH market over the next several years).

Redesca Market Share

Market data (IQVIA) have shown, as of April 2022, that biosimilars already eroded 35% of the Primary market with Redesca representing 54% of the overall biosimilar sales.

Over the coming months we expect the following:

- ➔ Enoxaparin Biosimilars to become dominant players in the LMWH enoxaparin market, as provinces and hospital exit past agreements and GPO tenders elect biosimilars as their products of choice.
- ➔ Provinces to de-list innovator drugs (already started in Quebec/New Brunswick/British Columbia) to prioritize enoxaparin biosimilar products over the innovator in the retail channel. At least one other major province is expected to de-list in the coming months.
- ➔ Enoxaparin biosimilars to start eroding the Secondary Market. This second wave of GPO/Provincial contract reviews will trigger significant opportunity for enoxaparin biosimilar such as Redesca™.

We believe Redesca™ and Valeo's team are well positioned to take advantage of the above market trends.

ONSTRYV®/YONDELIS®

Both products support a strategic position of Valeo in these key therapeutic areas.

Onstryv® could benefit from an improved market access and a patient support program enabling to assist certain patients without private health coverage to help with the cost of this medication which helps control the symptoms of PD. We recently engage with pCPA and active discussions are ongoing and could lead to a Letter of Intent (“LoI”) to support public reimbursement in some provinces.

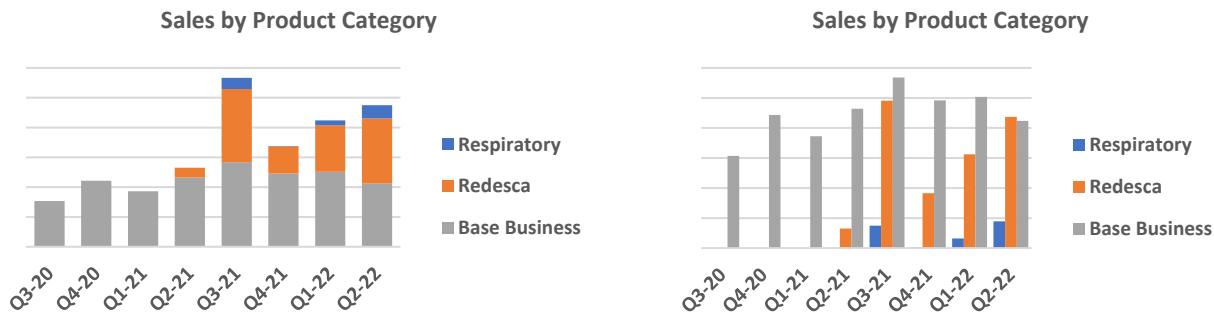
Yondelis® faces challenges due to a lack of public reimbursement and the difficulties associated with the need for 24-hour infusion. A patient support program aiming at navigating the health care system to provide coverage of the drug for cancer patients suffering from soft tissue sarcoma as well as providing support for infusion capabilities will be set up in the first half of FY-22. This should help patients to get broader access to Yondelis.

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Q2-22 Results Overview (industry data - Source – IQVIA April and May 2022)

Our Q2-22 and YTD-22 results reflect the added revenue and margin contributions of Redesca, Enerzair and Aetectura, three (3) transformative products launched in FY-21. Our base business has contributed lower than expected revenues and margins during Q2-22 due to timing issues. However, the continued strong sequential revenue growth of our three transformative products, Redesca, Enerzair and Aetectura, has contributed to expand our margins and improve our operating results during Q2-22 over prior quarters.

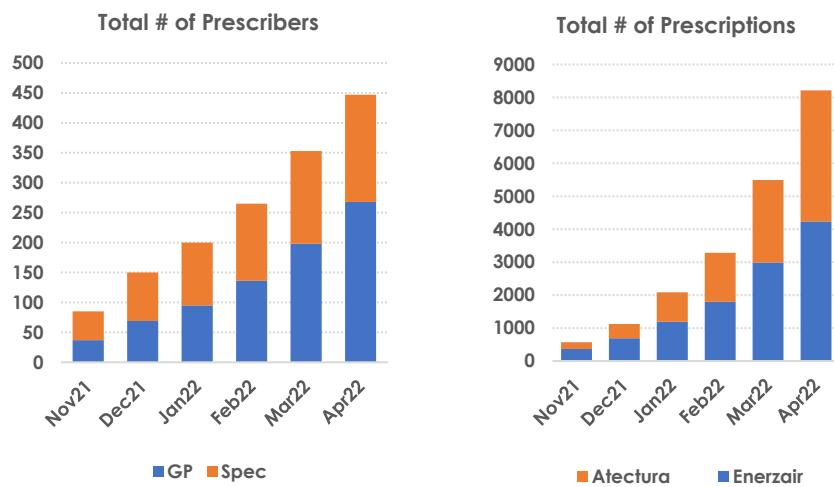


Following its launch in April 2021, Redesca sales in FY-21 and Q2-22 have been boosted by the need for hospitals to look for reliable and more economical alternatives to the existing originator product manufacturer which was experiencing COVID-19 related supply issues. This underlines the opportunity for Valeo to secure market shares for its LMWH Biosimilar, considering that Redesca is manufactured by the world's leading heparin producer, Techdow Hepalink.

As predicted, our FY-22 sales of Redesca are continuing to grow and are showing QoQ growth after a softer Q4-21 caused by the strong Q3-21 pipeline fill. (See Graph above). Our Q2-22 results have been positively impacted by stronger Redesca sales driven by recurrent demand from some key hospitals as well as the adjudication of new hospital market like in Quebec.

In addition to the growing contribution of Redesca on our overall revenues, our recent quarterly results are also showing the growing impact of Enerzair® and Aetectura® launched in June 2021, as well as the recurrent contribution from the rest of our commercial portfolio.

Enerzair and Aetectura have been launched in Q3-21, but the deployment of our full commercial team only took place at the end of FY-21. Since then, revenues for these two chronic innovative asthma products are growing monthly and are fuelled by the sequential addition of new prescribing practitioners, new patients, as well as the expansion of private and public reimbursement coverage that took place across Canada earlier in 2022. (See new prescribers and patients below)



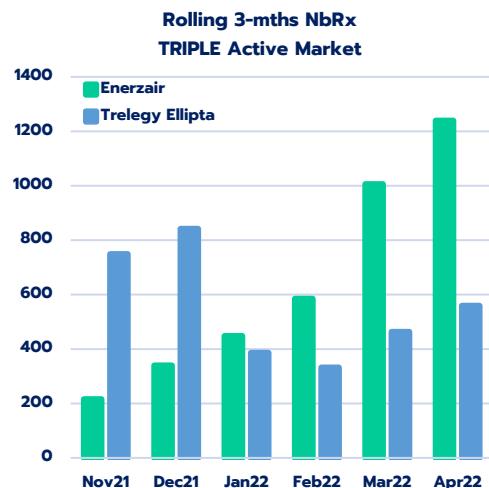
NBRx (New to Brand prescriptions - Existing Asthma patients “switching” to Enerzair and Aetectura)

NBRx is a key indicator of success for each of Enerzair (Triple Active Therapy) and Aetectura (Double Active Therapy). In addition to the sequential addition of new prescribing practitioners, new patients (See prior section), the success of any asthma drugs can be projected based on the number of patients switching from existing brands (New to Brand prescriptions or NBRx). With new asthma patients typically being initiated on single active therapies, NBRx represents patients switching from single to double active drugs (Aetectura and others), from double to triple active drugs (Enerzair and Trilogy) or switching from an existing double or triple active treatments to another similar treatments (Double -> Double, or Triple -> Triple).

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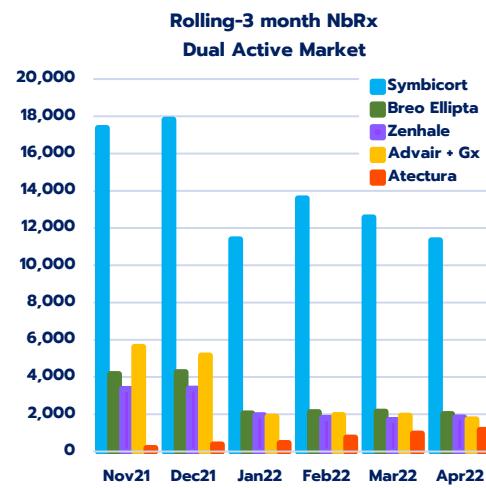
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Our historical sales data have only been reported since the October launch, but already we see strong Enerzair and Aetectura NbRx data (See Graph #1 and #2 below) which confirms the successful execution of our launch strategy and the rapidly growing market shares of each product within their respective TRIPLE and DUAL active segments.



Graph 1

Demonstrates the strong growth of the “TRIPLE” active segment of the Asthma market and Enerzair’s performance compared to the only other existing Triple therapy.



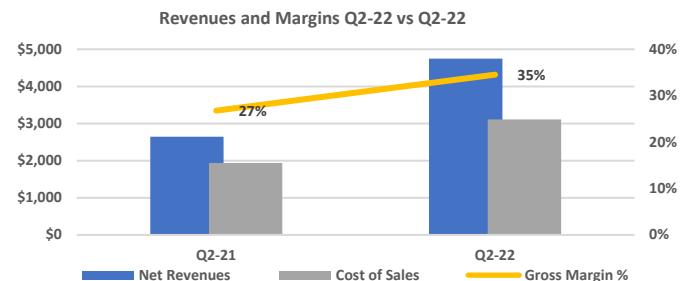
Graph 2

Illustrates Aetectura’s performance as a fast growing therapy within the large DOUBLE active segment of the Asthma market

Q2-22 Financial Results

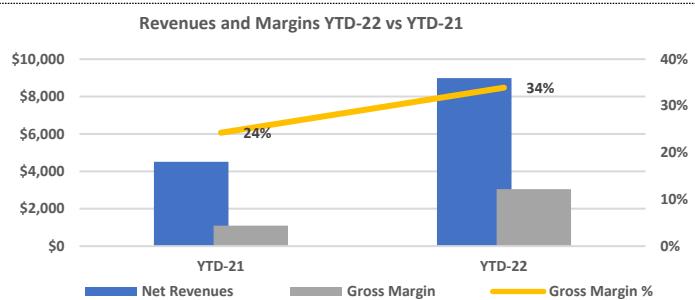
Q2-22 vs Q2-21 Performance

- Q2-22 Revenues grew 80% compared to Q2-21 at \$4.8 million compared to \$2.6 million.
- Q2-22 Gross Margin grew 134% compared to Q2-21.
- Net loss for Q2-22 was \$5.1 million.
- EBITDA loss for Q2-22 stood at \$3.6 million.
- Adjusted EBITDA loss for Q2-22 was \$3.8 million.



YTD-22 vs YTD-21 Performance

- YTD-22 Revenues grew 100% compared to YTD-21.
- YTD-22 Gross Margin grew 180% compared to YTD-21.
- Net loss for YTD-22 was \$11.0 million.
- EBITDA loss for YTD-22 stood at \$8.3 million.
- Adjusted EBITDA loss for YTD-22 was \$8.2 million.
- \$25M convertible debt secured during FY-22

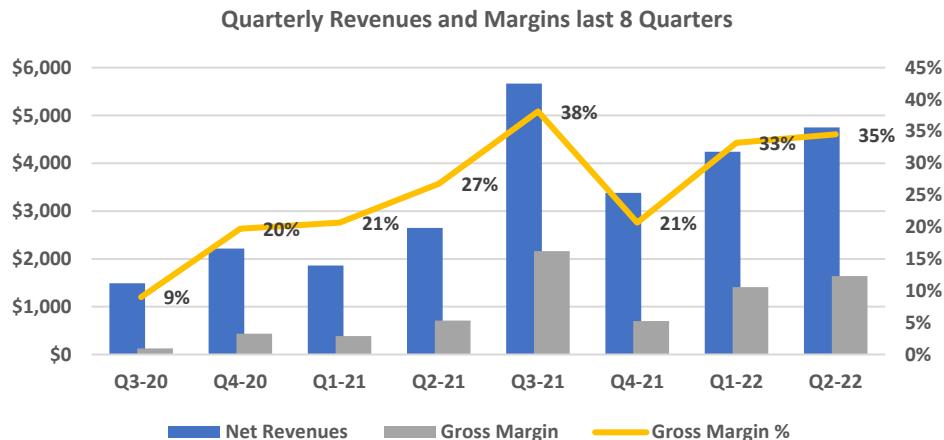


Q2-22 vs the prior quarter (Q1-22)

- Q2-22 Revenues grew 12% compared to Q1-22.
- Q2-22 Gross Margin grew 18% compared to Q1-22.
- Net loss for Q2-22 decreased by 13% compared to Q1-22.
- EBITDA loss for Q2-22 improved 22% over Q1-22
- Adjusted EBITDA loss for Q2-22 improved 14% over Q1-22.

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The above graph illustrates Valeo's performance over the last 8 quarters and the sequential quarterly growth of revenues and margins since the launch of Enerzair, Aetectura and Redesca in FY-21. Considering the \$180M+ peak sales potential of our existing product portfolio and our fixed costs infrastructure, the continued quarterly revenue growth of our portfolio will drive expanded margins and lead Valeo to profitability. (See "Liquidities" section of this MD&A).

During the course of last year, Valeo's results have been impacted by recurrent and non-recurrent costs related to setting up the new organizational structure and commercial team. This was required to take full advantage of the significant market opportunities for Redesca, Enerzair® and Aetectura®. Already, our FY-22 results are indicative of the progress made towards achieving profitability. We are expecting that the sequential increase of our revenues and margins – largely derived from the growth of Redesca, Enerzair and Aetectura sales will contribute to expand our operating margins and lead the company to profitability in the near future.

Also, our financial results show the full impact of the \$25 million convertible financing completed in December 2021. This financing has significantly strengthened our balance sheet and provided the capital required to support our operations and working capital requirements for the coming year. It is expected that our improved margins derived from the sequential quarterly growth of our revenues will help Valeo achieve its financial objective of becoming cash flow positive by the end of the current fiscal year.

Q2-22 Products Highlights

- On February 24, 2022, the Corporation announced the listing and public reimbursement of Enerzair and Aetectura in Ontario, Manitoba, New Brunswick, and by the NIHB and VAC federal programs.
- On March 22, 2022, the Corporation announced the listing and public reimbursement of Redesca and Redesca HP, in British Columbia. The Company also announces that Enerzair and Aetectura, have also been accepted for public reimbursement in Saskatchewan and in Prince Edward as of March 28, 2022.

Corporate & Financings

- On March 4, 2022, \$1,04 million of convertible debenture maturing February 27, 2023 plus accrued interest where converted into 2,600,419 common shares of the Corporation.
- On April 19th, 2022, the 12% Convertible Unsecured Subordinated Debentures issued pursuant to the \$15.0 million bought deal private placement closed on December 9, 2021, were approved for listing on the TSX under the symbol "VPH.DB" and begin trading.

Events Subsequent to Q2-22

- On May 2nd, 2022, the remaining non-convertible debentures issued July 2020 representing \$338 were reimbursed.
- On May 14th, 2022, the Corporation was informed by the respective provincial authorities that Enerzair and Aetectura, have been accepted for public reimbursement in British Columbia and Newfoundland.

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SELECTED FINANCIAL DATA

The following table sets forth financial information relating to the periods indicated and should be read in conjunction with the April 30, 2022, unaudited interim condensed consolidated financial statements.

Consolidated Statements of Loss

	Q2-22	Q2-21	Change		YTD-22	YTD-21	Change	
			\$ ¹	% ²			\$ ¹	% ²
Net Revenues	4,768	2,647	2,121	80%	9,009	4,508	4,501	100%
Cost of Sales	3,109	1,938	1,171	60%	5,941	3,414	2,527	74%
Gross Margin	1,659	709	950	134%	3,068	1,094	1,974	180%
Gross margin % to net sales	35%	27%		8%	34%	24%		10%
Expenses								
Sales and Marketing	3,539	949	2,590	273%	7,331	1,595	5,736	360%
General and Administrative	964	881	83	9%	2,229	1,825	404	22%
Medical affairs, QA & regulatory	845	257	588	229%	1,859	524	1,335	255%
Share Based Compensation	222	309	(87)	-28%	444	414	30	7%
Profit Sharing	32	1	31	3100%	43	1	42	4200%
Total Operating Expenses	5,602	2,397	3,205	134%	11,906	4,359	7,547	173%
Operating Loss	(3,943)	(1,688)	(2,255)	134%	(8,838)	(3,265)	(5,573)	171%
Other Expenses (income)								
Financial expense	1,178	213	965	453%	2,174	406	1,768	435%
Other income	(40)	(34)	(6)	18%	(70)	(78)	8	-10%
Unrealized loss on derivative warrant liability	17	-	17	100%	19	-	19	100%
Total Other Expenses	1,155	179	976	545%	2,123	328	1,795	547%
Net loss for the period	(5,098)	(1,867)	(3,231)	173%	(10,961)	(3,593)	(7,368)	205%
Other comprehensive loss								
Exchange differences on translating foreign operations	(6)	6	(12)	-200%	(4)	11	(15)	-136%
Defined benefit plan, net actuarial loss	74	93	(19)	-20%	74	90	(19)	-20
Total comprehensive loss	(5,030)	(1,768)	(3,262)	185%	(10,891)	(3,489)	(7,402)	212%
Loss per share								
Basic and diluted	(0.06)	(0.03)	(0.03)	122%	(0.14)	(0.06)	(0.08)	149%
Weighted average number of shares outstanding	80,661,530	65,565,241	15,096,289	23%	79,721,375	65,039,982	14,681,393	23%

1. A positive variance represents a positive impact to net income and a negative variance represents a negative impact to net income

2. Percentage change is presented in relative values

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EBITDA(L) Reconciliation (See "Management's Responsibility for Financial Reporting" – "Non-IFRS Financial Measures")

The following table provides a reconciliation of net loss to EBITDA(L) for Q2-22 as compared to Q2-21 as well as YTD-22 vs YTD-21

	Q2-22	Q2-21	Change		YTD-22	YTD-21	Change	
			\$ ¹	% ²			\$ ¹	% ²
Net Loss	(5,098)	(1,867)	(3,231)	173%	(10,961)	(3,593)	(7,368)	205%
Adjustments								
Interest Expense	1,198	190	1,008	531%	2,164	356	1,808	508%
Unrealized loss on derivative warrant liability	17	-	17	0%	19	-	19	0%
Depreciation	60	28	32	114%	119	55	64	116%
Amortization	187	122	65	53%	387	238	149	63%
EBITDA Loss	(3,634)	(1,527)	(2,107)	138%	(8,270)	(2,944)	(5,326)	181%
Other Adjustments								
Share-Based Compensation	222	309	(87)	-28%	444	414	30	7%
Recruitment costs - new product launch	-	50	(50)	-100%	-	175	(175)	-100%
Other warrants/ options costs	-	17	(17)	-100%	-	98	(98)	-100%
Inventory Write-off	7	14	(7)	-50%	-	17	(17)	-100%
Other provision	(370)	-	(370)	100%	(349)	-	(349)	100%
Adjusted EBITDA Loss	(3,775)	(1,137)	(2,638)	232%	(8,175)	(2,240)	(5,935)	265%

1. A positive variance represents a positive impact to net income and a negative variance represents a negative impact to net income

2. Percentage change is presented in relative values

	Q2-22 vs Q2-21, and YTD-22 vs YTD-21
Net Revenues	<ul style="list-style-type: none"> Net revenues represent sales of products based on Valeo's list price less chargebacks, price adjustments or other deductions related to provincial PLA's, GPO's agreements, early payment cash discounts, product returns or others. Such chargebacks and price deductions vary on a product-by-product basis. Consequently, the mix of product sales will greatly influence net revenues and ultimately our profitability. Our revenues are trending upwards due to the sequential addition of new products as well as market share gains for our lead products (Redesca, Enerzair and Atectura) which are benefiting from the recurrent contract wins (Redesca) or addition of new prescribers and patients requiring chronic treatments. <p>Net revenues in Q2-22 increased significantly over Q2-21 at \$4.8 million compared to \$2.6 million representing a 80% increase. For the YTD-22 period, net revenues increased significantly over YTD-21 at \$9.0 million compared to \$4.5 million representing a 100% increase. The QoQ and YTD increases resulted mainly from the strong contribution of Redesca in Q2-22 and YTD-22 which contributed to the full periods in FY-22 compared to less than 1 month in Q2-21 and YTD-21. The increase also reflected the continued commercial progress of Enerzair and Atectura which were formally launched in the later part of Q4-21 following the creation of our respiratory business unit (June-August 2021). Since Q4-21, sales of Enerzair and Atectura are growing monthly and are now having a material impact on our results. With private reimbursement now exceeding 90%, and public coverage being secured in most provinces, demand for these products is accelerating rapidly and fueled by a growing number of patients switching from other asthma therapies to Atectura and Enerzair. As more prescribers and patients adopt our drugs as their treatment of choice, this growing pool of patients provides a strong base of revenues that will help drive continued QoQ revenue growth and margin expansion.</p>
Gross Margin \$ and Gross Margin ratio %	<ul style="list-style-type: none"> As we launch new products and the commercial performance of our "Branded" product portfolio grows, we are set to see an improvement in our product mix, resulting in a significant expansion of our gross margin. This will directly impact our overall profitability. In addition to the transfer price for our products, our cost of goods also takes into consideration the amortization of product rights. These costs have increased in the second half of FY-21 following the license agreement with Novartis which is being amortized quarterly starting Q3-21. Our gross margin contribution in Q2-22 more than doubled over the Q2-21 period at \$1.7 million compared to \$0.7 million representing a 134% increase. Also, our gross margin contribution for YTD-22 increased significantly over the YTD-21 period at \$3.1 million compared to \$1.1 million representing a 180% increase. Our gross margin ratio for the same periods increased in FY-22 due to the improvement of our product mix. Gross margin ratio in Q2-22 compared to Q2-21 increased from 27% to 35%, while our gross margin ratio for YTD-22 stood at 34% vs 24% for YTD-21.

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	<ul style="list-style-type: none"> Amortization of product rights which includes amortization of the Novartis license fees since Q3-21 represented 3% of our net revenues for each of Q2-22 and FY-22 as compared to 2% and 3% of revenues for each of Q1-21 and FY-21
S&M expenses	<ul style="list-style-type: none"> As indicated earlier, Valeo commercializes Branded products that require S&M support, as well as hospital products such as M-Eslon, which require limited S&M commitments. Because S&M staff costs represents the bulk of the S&M expenses, those expenses have increased following the expansion of our commercial team and the creation of our respiratory business unit. Going forward we expect S&M expenses as a % of revenues to decrease over time. S&M expenses for Q2-22 were \$3.6 million compared to \$0.9 million for Q2-21. S&M expenses for YTD-22 were \$7.3 million compared to \$1.6 million for YTD-21. The increase between the reported periods resulted from the creation of our Respiratory BU and hiring of a dedicated sales team to support Redesca, Enerzair and Aetectura, which was required to capture the significant market opportunity for these products. We are on track to achieve peak sales of \$30-35 million for Redesca by FY-24 and expect to reach peak sales of \$125 million for Enerzair and Aetectura by FY-25/26. Most of the FY-22 increases were due to the addition of our salesforce which came as a result of our HO and commercial platform expansion during the second half of FY-21. The balance was related to promotion and marketing activities and costs for sampling, marketing material and programs, and to a lesser extent field activities. Over time we expect costs for samples, marketing materials and other S&M expenses to be more representative of recurrent spending and to trend downward as a % of revenues.
G&A expenses	<ul style="list-style-type: none"> Valeo's G&A expenses consist primarily of staff costs for our non-S&M management team such as administration, finance and accounting, business development, legal, and supply chain personnel. G&A expenses also include IR expenses which can fluctuate significantly between quarters as the Company implements various IR initiatives. G&A expenses for Q2-22 were \$1.0 million as compared to \$0.9 million for Q2-21 representing a 9% increase. G&A expenses for YTD-22 were \$2.2 million as compared to \$1.8 million for YTD-21 representing a 22% increase. The increase in G&A expenses resulted from the addition of HO personnel required to support our growth. Following the creation of our new corporate structure (See "Overview of the Business") we have created several key additional HO positions to support our new business model. The new structure, which includes two newly created Respiratory and Specialty products BU, was completed in the second half of FY-21 and will provide significant leverage over the coming years. Consequently, as expected our G&A expenses as a % of net revenues are trending downward compared to prior quarter. Such expenses represented 20% of our revenues in Q2-22 compared to 56% in Q4-21. Our Q2-22 and YTD-22 G&A expenses also benefited from a \$370 gain following the recovery of part of the loss incurred in Q4-21 associated with a bank fraud (more details can be found in our Q4-21 MD&A).
Medical Affairs, Quality Assurance and Regulatory ("MA & Reg")	<ul style="list-style-type: none"> MA & Reg expenses for Q2-22 were \$0.8 million compared to \$0.3 million for Q2-21. MA & Reg expenses for YTD-22 were \$1.9 million compared to \$0.5 million for Q2-21. In order to support our fast-growing branded product portfolio, we have expanded our MA, QA and Regulatory team and activities over the past year. Over time, we expect these expenses to trend downward as a % of revenues as we take full advantage of the market opportunities for our branded product portfolio.
SBC expenses	<ul style="list-style-type: none"> SBC expenses represent the costs relating to the issuance of stock options and RSUs to new staff and board members and the vesting of same over time. SBC expenses were \$0.2 million in Q2-22 as compared to \$0.3 million for Q2-21 representing a \$0.1 million decrease between the two periods. SBC expenses for YTD-22 and YTD-21 were stable at \$0.4 million for each period.
Profit Sharing	<ul style="list-style-type: none"> Profit sharing arrangements represent agreements with our partners to share net contribution from the sale of products. The increase in FY-22 profit share amounts is associated with the growth of our Redesca revenues and margins which drive profit share remittance to our partners.
Total Operating Expenses ("Total OPEX")	<ul style="list-style-type: none"> Total operating expenses stood at \$5.6 million and \$11.9 million in Q2-22 and for the YTD-22 period, compared to \$2.3 million in Q2-21 and \$4.3 million for the YTD-21 period. Our Total OPEX have increased in the later part of FY-21 to support the growth of our commercial platform and HO infrastructure. Since then, our ratio of total OPEX to revenues is declining as we take full advantage of this operational leverage. For Q2-22 the ratio of total OPEX to revenues stood at 117% compared to 149% for the prior Q1-22 period and 90% for Q2-21. We expect the ratio of Total OPEX to revenues to decline sequentially over the coming quarters as we continue to execute our commercial initiatives and take full advantage of the market opportunity for our lead products.
Financial expenses	<ul style="list-style-type: none"> Financial expenses reflect the capital structure of the Corporation and include costs for issuing interest bearing debentures in lieu of shares to finance our operations. The financial expenses also capture the costs for non-recurrent use of our operating line of credit, supplier financing, other financial charges and bank fees. Financial expenses also capture FX gain or loss, as well as lease interest. Our financial expenses were \$1.2 million in Q2-22 compared to \$0.2 million in Q2-21 representing a \$1.0 million increase. Financial expenses for Q2-22 included \$0.3 million as effective interest on the debentures.

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	<ul style="list-style-type: none"> Financial expenses for YTD-22 were \$2.2 million as compared to \$0.4 million in FY-21. Financial expenses for Q2-22 included \$0.6 million as effective interest on the debentures. The increases for each of Q2-22 and YTD-22 was due to a series of debenture financings closed over the past year. Valeo implemented a \$25 million convertible debenture financing on December 9, 2021. Despite the conversion and repayment of the April 2021 Bridge and some prior-existing debentures that took place as a result of this financing, the net impact of the new debentures increased our financial expenses in Q2-22 and YTD-22 as compared to the corresponding periods in FY-21 The increase between the two reported quarters also included incremental lease interest charges which resulted from the expansion and extension of our HO lease as well as an increase in the effective interest cost for the various debentures outstanding. The effective interest costs capture the cost relative to the issuance of warrants as a mean of reducing the actual interest in such instruments.
Other income	<ul style="list-style-type: none"> Nominal variations between the periods. The Corporation continues to provide back-office, accounting, regulatory and other consulting services as a means of leveraging its staff's expertise.
Unrealized loss on derivative warrant liability	<ul style="list-style-type: none"> Following the April 2021 bridge financing, warrants issued as part of the transaction resulted in the creation of an embedded derivative warrant liability. Going forward and until the April 2021 warrants are converted or expire, the change in fair value of the derivative instrument between the end of each reported period will be expensed on our Statement of Loss. For the Q2-22 period, the impact of the re-evaluation of the embedded derivative was \$17 compared to nil in Q2-21. The YTD-22 unrealized loss on derivative warrant liability totaled \$19 for the YTD-22 compared to nil for YTD-21.
Net loss for the period	<ul style="list-style-type: none"> In Q2-22, the growth of revenues and margins have contributed to reduce our quarterly loss compared to the prior Q1-22 period. Despite strong commercial gains, our net loss in Q2-22 was \$5.1 million compared to \$1.9 million in Q2-21. Our net loss in Q2-22 decrease 13% compared to the prior Q1-22 performance. Our net loss for YTD-22 was \$11.0 million as compared to \$3.6 million for YTD-21. Our Net loss for Q2-22 and YTD-22 reflect the incremental costs involved in the creation of the 2 BUs, as well as expansion of Valeo's commercial, medical and HO teams in the second part of FY-21. These initiatives were required to capture the significant market opportunities for Redesca, Enerzair and Aetectura as well as to accelerate the growth of other existing products. As demonstrated by the progress achieved over the last quarter, our new corporate and commercial infrastructure will help accelerate our growth and improve our profitability. Considering the bulk of our G&A expenses remains flat as a % of revenues, we expect that the continued revenue growth of Redesca, Enerzair and Aetectura will drive QoQ sequential margin expansion and help eliminate our operating loss over the near future.
EBITDA (Loss)	<ul style="list-style-type: none"> Management believes our EBITDA performance is more indicative of the commercial progress achieved by the Corporation as it eliminates the financial costs associated with our financial structure and the amortization of prior investments in our product portfolio such as license fees and regulatory filings. (See "Management's Responsibility for Financial Reporting" – "Non-IFRS Financial Measures") EBITDA loss in Q2-22 was \$3.7 million compared to \$1.5 million in Q2-21. EBITDA loss for YTD-22 was \$8.3 million compared to \$2.9 million for YTD-21. Same as for our net loss analysis, our EBITDA loss for each of Q2-22 and YTD-22 reflected the net impact of the creation of our new commercial and corporate structure in FY-21. Our EBITDA loss was up in Q2-22 at \$2.1 million but was down \$1.0 million compared to Q1-22, a 22% improvement which is indicative of our progress made towards our objective of achieving EBITDA profitability by leveraging our corporate infrastructure and the commercial potential of our existing commercial pipeline.
Adjusted EBITDA (L)	<ul style="list-style-type: none"> (See "Management's Responsibility for Financial Reporting" – "Non-IFRS Financial Measures") Our Adjusted EBITDA loss in Q2-22 and FY-22 includes adjustments such as Share-Based Compensation, as well as other non-recurrent adjustments to our net loss. For Q2-22, our Adjusted EBITDA loss also reflected the adjustment for the \$370 partial recovery of the bank fraud booked in Q4-21 as a positive adjustment to our Adjusted EBITDA loss. Following such adjustments, our Adjusted EBITDA loss in Q2-22 was \$3.8 million compared to \$1.1 million in Q2-21, representing a \$2.7 million increase, but down \$0.6 million or 14% compared to Q1-22. Adjusted EBITDA loss for YTD-22 was \$8.2 million compared to \$2.2 million for YTD-21.

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Consolidated Balance Sheet Highlights

As at	April 30, 22	Oct 31, 21	Change	
			\$ ¹	% ²
Cash	5,226	2,043	3,183	156%
Trade and other receivables	2,295	1,798	497	28%
Inventory	7,810	7,675	135	2%
Total current assets	16,147	12,350	3,797	31%
Property and equipment	1,354	1,174	180	15%
Right of use asset	924	967	(43)	-4%
Intangible assets	5,989	6,539	(550)	-8%
Total assets	24,414	21,030	3,384	16%
Trade accounts payable	3,197	7,320	(4,123)	-56%
Other accounts payable and accrued liabilities	1,329	2,635	(1,306)	-50%
Accrued interest on debentures	270	266	4	2%
Provisions	29	214	(185)	-86%
Convertible debentures (short-term)	709	0	709	100%
Non-convertible debentures (short-term)	338	4,854	(4,516)	-93%
Derivative warrant liability	601	-	601	100%
Total current liabilities	6,521	15,334	(8,813)	-57%
Convertible debentures	19,724	1,605	18,119	1129%
Lease liabilities	1,140	1,165	(25)	-2%
Defined benefit obligations	199	291	(92)	-32%
Derivative warrant liability	-	582	(582)	-100%
Total liabilities	27,584	18,977	8,607	45%
Share capital	25,720	24,616	1,104	4%
Warrants	3,778	3,769	9	0%
Contributed surplus	7,252	2,697	4,555	169%
Deficit	(39,671)	(28,710)	(10,961)	38%

1. A positive variance represents a positive impact the balance sheet and a negative variance represents a negative impact to the balance sheet
 2. Percentage change is presented in relative values

	Q2-22 vs YE-21
Cash and liquidities	<ul style="list-style-type: none"> Our cash balance at the end of Q2-22 was \$5.2 million compared \$2.0 million at YE-21 representing a \$3.2 million increase. The increase included the net impact of our \$25 million convertible financing closed in December 2021 less cash required for working capital and operating requirements for the first part of FY-22.
Trade and other receivables	<ul style="list-style-type: none"> Typically, our trade receivables average aging ranges between 35-40 days and tend to be collected rapidly due to the early payment cash discounts offered to clients and distributors. Early payment cash discounts are customary throughout the pharma industry, and they facilitate a fast conversion of receivables into cash. Our trade and other receivables increased by \$0.5 million between YE-21 and Q2-22 which is indicative of the commercial progress made between the 2 reported periods.
Inventory	<ul style="list-style-type: none"> Our inventory will fluctuate between periods to reflect sales of products and the requirements to support revenue growth and product launches. Typical shelf life for pharmaceutical products is 18-36 months and for that reason, product requirements for new product launches can often last more than one year and will tend to negatively impact short term cash flows and working capital requirements. Our inventory levels have remained stable between YE-21 and Q2-22 as new inventory purchased was offset by COGS used during the quarter.
Total current assets	<ul style="list-style-type: none"> Current assets have increased by \$3.8 million or 31% between the 2 periods mainly because of the net impact of the December 2021 financing on our cash position (See "Cash and liquidities" above).
Property and Equipment	<ul style="list-style-type: none"> Property and equipment represent investment in our HO and warehouse shelving, vaults and other equipment. Following the addition of three transformational assets over the last year (Redesca, Aetectura and Enerzair) we have made significant investment to expand our warehousing capabilities.

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	<ul style="list-style-type: none"> Between YE-21 and Q2-22 our property and equipment has increased from \$1.2 million to \$1.4 million, representing investments for increasing warehousing capabilities as well as additions in IT assets. Unlike other specialty pharmaceuticals companies that rely on 3PL ("Third Party Logistics") suppliers, Valeo's warehousing capabilities offer significant operational savings by eliminating 3PL costs.
Right of Use Asset ("ROU asset")	<ul style="list-style-type: none"> The right-of-use asset represents Valeo's right to use its leased facility over the life of a lease and is amortized over the term of the lease. During FY-21 we have renewed our lease for an additional 8 years and have expanded our lease area which translated in a net increase in our ROU assets. Concurrent to the increase of our ROU assets, our lease obligations have also increased. (See "Lease Liabilities" below). Between YE-21 and Q2-22, right-of-use assets have decreased slightly due to amortization charges.
Intangible assets	<ul style="list-style-type: none"> Intangible assets represent investments made in order to build our product pipeline. For assets owned by Valeo, these assets include formulation, R&D costs, regulatory and filings expenses. For other products, intangible assets include license fees to acquire product rights, regulatory fees and expenses as well as expenses to improve market access. Intangible assets are amortized using the straight-line method, over the remaining useful life of the asset (or license) starting when the product is ready for commercialization – typically when Valeo receives marketing approval and its first commercial product lot. Intangible assets are tested quarterly for impairments as per IFRS Standards (IAS 38) to ensure that the recoverable value of each assets exceeds its book-value. Our intangible assets have decreased by \$0.6 million in Q2-22 compared to YE-21 due to amortization charges. The amortization of license fees and transaction costs related to the Novartis license has commenced during Q3-21.
Total assets	<ul style="list-style-type: none"> Total assets increased by \$3.4 million between YE-21 and Q2-22, mainly as a result of the net impact of the December 2021 financing. (See "Cash and liquidities" above).
Accounts payables	<ul style="list-style-type: none"> Our trade accounts payables have decreased by \$4.1 million between YE-21 and Q2-22 representing a 56% decrease. The YE-21 trade accounts payables included the impact of a large shipment of Redesca products which arrived prior to end of Q4-21. The cost of this shipment was reflected in our trade payables at YE-21 and was settled during the first quarter of FY-22.
Other payables and accrued liabilities	<ul style="list-style-type: none"> Other payables and accrued liabilities decreased by \$1.3 million between YE-21 and Q2-22. The YE-21 levels included a non-recurrent \$0.5 million accrual for hiring fees relating to the creation of our Respiratory commercial team, as well as accruals for bonuses and other staff charges most of which were settled during the first quarter of FY-22
Provisions	<ul style="list-style-type: none"> Provisions include price accruals for price rebate and chargebacks resulting from GPO and PLA agreements not yet invoiced, as well as accruals for product returns. Provisions required at the end of Q2-22 have decreased by \$0.2 million as the bulk of our revenues during the quarter were made at list price or based on GPO contract prices and thus did not require provisions for gross to net sales adjustments.
Short term portion of Convertible Debentures	<ul style="list-style-type: none"> Convertible debentures issued in February and March 2020 ("2020 Debentures") will mature in Q2-22 and now appear as short-term liabilities. The amount of 2020 Debentures was reduced during the last quarter as \$1.0 million debentures plus accrued interest were converted into common shares.
Short term portion of Non-Convertible Debentures	<ul style="list-style-type: none"> At the end of Q2-22, Valeo had \$0.3 million of non-convertible debentures due over the next 12 months. This amount was down \$4.5 million compared to YE-21. The \$0.3 million balance was repaid at the start of Q3-22.
Derivative warrant liability	<ul style="list-style-type: none"> Following the April 2021 bridge financing, warrants issued as part of the transaction resulted in the creation of an embedded derivative warrant liability. Because the April 2021 bridge financing warrants expire during Q2-23, such liability is now reported as short-term.
Total current liabilities	<ul style="list-style-type: none"> Our current liabilities have decreased by \$8.8 million between YE-21 and the end of Q2-22. The reduction was due to the strong reduction of our trade payables and accrued liabilities as well as the conversion or repayment of debenture maturing over the coming year, but partly offset by the derivative warrant liability which is now reported as a short-term liability.
Convertible debentures	<ul style="list-style-type: none"> During the Q1-22 quarter, the Corporation completed a \$25 million convertible debentures financing. After netting the \$4.4 million allocation of the conversion features of the debenture to our contributed surplus and taking into account existing debentures converted into the financing or repaid subsequent to the transaction, the net impact of issuing those debentures represented an increase of \$18.1 million as at the end of Q2-22 compared to our YE-21 balance.
Lease liability (long-term portion)	<ul style="list-style-type: none"> The lease liability (long-term portion) represents the present value of Valeo's non-current lease payments less the ROU asset (See above). There was nominal variance between the two reported periods.
Defined Benefit obligations	<ul style="list-style-type: none"> The Defined benefit obligations represents Valeo's obligations towards the pension fund in excess of the pension fund assets. During Q2-22 such obligations have decreased by 32% following changes to our obligations as compared to our pension fund assets.

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Total liabilities	<ul style="list-style-type: none"> The \$8.6 million increase between YE-21 and Q2-22 reflects the issuance of the \$25 million debentures less the reduction of trade payables and accrued liability (see comments above), as well as the \$4.5 million reduction in non-convertible debentures.
Share Capital	<ul style="list-style-type: none"> The \$1.1 million increase between YE-21 and Q2-22 reflected the conversion of convertible debentures in Q2-22 less the issue costs related to the issuance of convertible debentures in Q1-22.
Warrants	<ul style="list-style-type: none"> No material changes between the two reported periods
Contributed Surplus	<ul style="list-style-type: none"> The \$4.6 million increase between YE-21 and Q2-22 included the \$4.4 million allocation of the conversion features of the debenture issued during Q2-22, as well as \$0.4 million for share-based compensation, less a \$0.3 million recovery for expired options/warrants and debentures converted.
Deficit	<ul style="list-style-type: none"> Increase reflects the performance of the Corporation during the year – Statement of Loss

SELECTED QUARTERLY FINANCIAL INFORMATION

	Q2-22	Q1-22	Q4-21	Q3-21	Q2-21	Q1-21	Q4-20	Q3-20
Net Revenues	4,768	4,241	3,382	5,667	2,647	1,861	2,215	1,490
Cost of Sales	3,109	2,832	2,682	3,506	1,938	1,476	1,778	1,363
Gross Margin	1,659	1,409	700	2,161	709	385	437	127
<i>Gross Margin % to net sales</i>	<i>35%</i>	33%	21%	38%	<i>27%</i>	21%	20%	9%
Expenses								
Sales and Marketing	3,539	3,792	4,183	2,399	949	646	333	401
General and Administrative	964	1,265	1,897	1,721	880	945	627	725
Medical affairs, QA & regulatory	845	1,014	1,258	432	257	267	288	226
Share Based Compensation	222	222	409	173	309	105	232	162
Profit Sharing	32	11	9	55	1	-	(9)	23
Total Operating Expenses	5,602	6,304	7,756	4,780	2,397	1,963	1,471	1,537
<i>Total OPEX to Revenue ratio %</i>	<i>117%</i>	149%	229%	84%	<i>91%</i>	105%	66%	103%
Operating Loss	(3,943)	(4,895)	(7,056)	(2,619)	(1,687)	(1,578)	(1,034)	(1,410)
Other expenses (income)								
Financial expense	1,178	996	496	375	213	193	176	249
Other income	(40)	(30)	(21)	(25)	(34)	(44)	(34)	(44)
Unrealized loss on derivative warrant liability	17	2	130	10	-	-	-	-
Total Other Expenses	1,155	968	605	360	179	149	142	205
Net loss for the period	(5,098)	(5,863)	(7,661)	(2,979)	(1,867)	(1,727)	(1,176)	(1,615)
EBITDA (Loss)	(3,634)	(4,636)	(6,719)	(2,332)	(1,526)	(1,417)	(880)	(1,271)
Adjusted EBITDA (Loss)	(3,775)	(4,400)	(5,436)	(836)	(1,136)	(1,103)	(486)	(705)

(See "Management's Responsibility for Financial Reporting" – "Non-IFRS Financial Measures")

Notes	Valuable information
Revenues	<ul style="list-style-type: none"> Our revenues in Q2-22 were up 12% compared to the prior Q1-22 quarter which is indicative of the continued commercial progress made by Redesca, Enerzair and Aetectura. Our Q4-21 revenues were down compared to Q3-21 as the impact of the Q3-21 pipeline fill was absorbed and led to softer sales of Redesca for that quarter. Our Q3-21 results included the strong pipeline fill that followed the launch of Resdesca. Redesca sales started in Q2-21 and have been material since launch. This explains the growth of our revenues and margins after Q1-21.
Cost of Sales and Gross Margin	<ul style="list-style-type: none"> Fluctuates with revenues as well as the mix of product sold. The continued improvement of our product mix and the growing contribution of higher margin products such as Redesca, Enerzair and Aetectura has contributed to stronger margins since Q2-21, except for Q4-21 which was impacted by reduced sales following the Q3-21 Redesca pipeline fill. Cost of Sales also includes amortization of product rights previously capitalized as intangible assets. Such amortization starts upon the launch of the respective products.

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S&M expenses	<ul style="list-style-type: none"> Our S&M expenses have decreased by 7% in Q2-22 as compared to the prior quarter. This followed an increase in S&M expenses in Q3-21 and Q4-21. The addition of 54 sales professional during FY-21 and the increased S&M activities to support the commercialization of Redesca, Enerzair, and Aetectura impacted our S&M expenses starting Q3-21. Our salesforce is now fully operational and can support several new products, and this should facilitate an improvement of our net results following the addition of new branded products.
G&A expenses	<ul style="list-style-type: none"> G&A expenses were down 24% in Q2-22 compared to Q1-22 and 33% in Q2-22 compared to Q4-21. Similar to S&M expenses, our G&A expenses increased in Q3-21 ad Q4-21 following the creation of our new commercial infrastructure and expansion of HO activities to support the expansion of our commercial pipeline. As expected, G&A expenses are trending down as a % of revenues at 20% in Q2-22 compared to 56% in Q4-21. G&A expenses over the last 4 quarters were impacted by a bank fraud leading to \$548, \$371, and \$21 provisions and expenses net of recovery for each of Q3-21, Q4-21 and Q1-22 as well as a \$371 recovery for Q2-22.
Medical Affairs, Quality Assurance and Regulatory ("MA, QA & Reg")	<ul style="list-style-type: none"> Our MA & Reg costs have increased in Q4-21 reflecting the costs of the expanded MA department, which is required to support the commercialization of Redesca, Enerzair and Aetectura. MA and Reg costs also reflect the increase in PSP (Patient support Programs) and the increase in advisory board meetings with our expanding network of KOL's and opinion leaders. The 19% and 17% respective QoQ decreases in Q1-22 and Q2-22 compared to prior quarter is due to timing of MA and Reg activities.
SBC expenses	<ul style="list-style-type: none"> Represents the costs of issuing stock options and RSUs. Fluctuation between quarters is due to the hiring of staff, the addition of Board members and the vesting associated with issued options and RSUs. The issuance and vesting of a large number of options issued to new staff over the recent quarters impacted the SBC expenses for those quarters.
Profit Sharing	<ul style="list-style-type: none"> Starting Q3-20 the Corporation started accruing and paying amounts under profit-sharing arrangements. Such arrangements are meant to reduce the transfer price to be paid by Valeo and have the licensee and licensor share the commercial success of the products.
Total Operating Expenses ("Total OPEX")	<ul style="list-style-type: none"> Our Total OPEX have increased in Q4-21 to support the growth of our commercial platform and HO infrastructure thus providing significant leverage to grow our revenues and add key products to commercial portfolio. Since then, our ratio of total OPEX to revenues is declining as we take full advantage of this operational leverage. The ratio of total OPEX to revenues is 117% for Q2-22, compared to 149% in Q1-22, and 229% in Q-21 following the expansion of our operations. We expect the ratio of Total OPEX to revenues to decline sequentially over the coming quarters as we continue to execute our commercial initiatives and take full advantage of the market opportunity for our lead products.
Financial expenses	<ul style="list-style-type: none"> Our financial expenses fluctuate between quarters depending on the level of short term and long-term borrowing required to fund our operations. Our Financial expenses increased in Q1-22 following the implementation of the \$25 million convertible financing. Financial expenses increased in Q3-21 following the closing of our \$6.6 million non-convertible financing.
Other (Income) expenses	<ul style="list-style-type: none"> Fluctuates between periods based on the level of services rendered. The Corporation continues to provide back-office, regulatory and other consulting services as a mean of leveraging its staff's expertise.
Net loss	<ul style="list-style-type: none"> Our Net loss in Q2-22 has decreased by 13% as compared to Q1-22 due to the growth of our revenues and margins as well as control over our expenses and the fraud recovery. Our Net loss had increased in Q4-21 compared to Q3-21 due to the respective increase in S&M, G&A, and financial expenses explained earlier. We expect our net loss to be eliminated over the coming year as we continue experiencing revenues growth and secure the benefits of incremental market shares from Redesca, Enerzair, and Aetectura as well as other products in our portfolio. We believe that in order to eliminate the impact of our debentures and several non-cash items, that the EBITDA (L) and Adjusted EBITDA(L) metrics to be more representative of our quarterly performance. (See EBITDA (L) and Adjusted EBITDA (L) below.)
EBITDA (Loss)	<ul style="list-style-type: none"> EBITDA Loss (See "Management's Responsibility for Financial Reporting" – "Non-IFRS Financial Measures") eliminates the impact of the CDU, ITC and other financings which reflect the Corporation's financing strategy adopted to attract the required capital to fund its operations. Similar to our net operating loss, over the last year our EBITDA loss has also been impacted by staff additions and expenses required to support the growth of our organization, the creation of our new corporate and sales structure and the launch of new products. Our EBITDA loss for Q2-22 was down 22% compared to the prior quarter due to improved operating margins and a reduction in our operating expenses. Our contribution margins increased 16% between the two quarters and our operating expenses were down 11% in Q2-22 compared to Q1-22. The improvement in EBITDA loss is indicative of our progress made toward achieving EBITDA profitability over the coming year.

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Adjusted EBITDA (Loss)	<ul style="list-style-type: none"> Our Adjusted EBITDA (Loss) is a much better indicator of our progress over the last year as it eliminates the impact of non-recurrent expenses required to execute our business plan and achieve fast growth objectives. Our Adjusted EBITDA (loss) in Q2-22 improved 14% in Q2-22 compared to Q1-22. Our Adjusted EBITDA (loss) had increased in Q4-21 compared to Q3-21 following the implementation of our new commercial and HO structure and incremental costs required to support the launch of Aetectura, Enerzair and Redesca. Similar to our net loss and EBITDA (Loss), we expect our Adjusted EBITDA performance to trend upward over the coming quarters as the sales growth of Redesca, Enerzair, and Aetectura, as well as other products in our portfolio translate into incremental operating margins, hence contributing to reduce/eliminate our Adjusted EBITDA loss.
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LIQUIDITIES AND CAPITAL RESOURCES

	Q2-22	Q2-21	Change		YTD-22	YTD-21	Change	
			\$	%			\$	%
Net loss from operations	(5,098)	(1,867)	(3,231)	173%	(10,961)	(3,593)	(7,368)	205%
Other Items not affecting cash	581	581	-	0%	1,841	984	857	87%
Changes in non-cash working capital	(1,678)	(307)	(1,371)	447%	(6,128)	(1,680)	(4,448)	265%
Cash used in operations	(6,195)	(1,593)	(4,602)	289%	(15,248)	(4,289)	(10,959)	256%
Investing activities								
Cash used by investing activities	(132)	(2,074)	1,942	-94%	(289)	(2,190)	1,901	-87%
Financing Activities								
Cash (used) provided by financing activities	(563)	6,685	(7,248)	-108%	18,674	7,398	11,276	152%
Foreign exchange loss (gain) on cash	23	(85)	108	-127%	46	(105)	151	-144%
Increase (decrease) in cash	(6,867)	2,933	(9,800)	-334%	3,183	814	2,369	291%
Cash, beginning of the period	12,093	717	11,376	1587%	2,043	2,836	-793	-28%
Cash, end of period	5,226	3,650	1,576	43%	5,226	3,650	1,576	43%

1. A positive variance represents a positive impact to the cash flow and a negative variance represents a negative impact to the cash flow
2. Percentage change is presented in relative values

	Q2-22 vs Q2-21	YTD-22 vs YTD-21
Cash used in operations	<ul style="list-style-type: none"> Cash used in operations represents cash flows from operations, excluding income and expenses not affecting cash. Cash used in operations for Q2-22 was \$6.2 million compared to \$1.6 million in YTD-21. The \$4.6 million increase came from a \$3.2 million increase in net loss, and a \$1.4 million increase in non-cash working capital due mainly to a reduction of trade payables and accrued liabilities. There were no material changes in other items not affecting cash between the 2 reported periods. 	<ul style="list-style-type: none"> Cash used in operations was \$15.4 million in YTD-22 compared to \$4.3 million in YTD-21. The \$11.2 million increase came from a \$7.4 million increase in net loss, and a \$4.6 million increase in non-cash working capital. During YTD-22, trade payables and accrued liabilities decreased by \$5.5 million while inventory only increased by \$0.1 million. During YTD-21, trade payables and accrued liabilities generated cash by increasing by \$3.5 million but offset by a \$4.6 million use of cash to support an increase in inventory. The net cash used for non-cash working capital in YTD-22 was partially offset by the increase in items not affecting cash for \$1.8 million including \$0.8 million for interest expenses, and \$0.4 million and \$0.5 million for each of share-based compensation and depreciation/amortization charges.
Cash used in investing activities	<ul style="list-style-type: none"> Cash used by investing activities during Q2-22 was \$0.1 million compared to \$2.1 million in Q1-21. Cash used by investing activities was \$0.3 million in YTD-22 compared to \$2.2 million in YTD-21. The QoQ and YTD variances reflects addition to property and equipment during FY-22 as compared to investments in products rights in FY-21 as a result of the license fee paid to Novartis and related cost to acquire rights to Enerzair and Aetectura. 	
Cash provided by financing activities	<ul style="list-style-type: none"> During Q2-22 we used \$0.6 million to repay part of the remaining amounts due under the non-convertible debentures issued in July 2020. 	<ul style="list-style-type: none"> During YTD-22, financing activities provided cash of \$18.7 million compared to \$7.4 million for the YTD-21 period. During YTD-22, Valeo secured \$23.5 million from the net proceeds of the convertible debenture financing closed in December 2021, less \$4.8 million representing repayments and conversion of prior existing debentures. During YTD-21, the Corporation secured a

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	• During Q1-21, we secured \$6.7 million of funds mainly as a result of the \$6.6 million bridge financing secured in April 2021.	\$6.6 million bridge financing as well as \$1.0 from the issuance of shares following the conversion of warrants and to a lesser extend from the exercise of options.
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Related Party Transactions

The following table presents the related party transactions for the respective periods:

	Three months ended April 30,		Six months ended April 30,	
	2022	2021	2022	2021
Key management salary and benefits	367	295	962	520
Directors and employee stock option compensation	222	309	444	414
Consulting fees paid to a company controlled by an officer	59	46	146	91
Service income	31	34	55	77
	679	684	1,607	1,102

The following table represents the related party transactions as at:

	April 30, 2022	October 31, 2021
Amounts owed to key management, officers and directors		
Consulting fees	-	11
Convertible debentures	538	231
Accrued interest on convertible debentures	8	5
Non-convertible debentures	15	436
Accrued interest on non-convertibles debentures	1	14
Amounts owed to Manitex, a shareholder of the Corporation		
Non-convertible debentures	-	15
Accrued interest on non-convertible debentures	-	1
Amounts owed to 100079 Canada Inc., a shareholder of the Corporation		
Convertible debentures	1,285	955
Accrued interest on convertible debentures	15	24
Non-convertible debentures	-	2,041
Accrued interest on non-convertible	-	100

Going Concern

This MD&A have been prepared on a going-concern basis, which implies that the Corporation will continue realizing its assets and discharging liabilities in the normal course of business for the foreseeable future. As reflected in the quarterly unaudited financial statements, the Corporation is in the process of ramping up its activities and has not yet achieved profitability. During the three-month period ended on April 30, 2022, the Corporation incurred a net loss of \$5.1 million, and used cash in operations of \$6.2 million. Despite the positive working capital of \$9.6 million at the end of Q2-22, this raises significant doubt about the Corporation's ability to continue as a going concern.

Accordingly, the ability of the Corporation to realize the carrying value of its assets and continue operations as a going concern is dependent upon its ability to obtain additional financing and ultimately on generating future profitable operations. Management anticipates that the commercialization of new products will provide incremental cash flow that could contribute to working capital requirements. There are no assurances that any of these initiatives will be successful. Factors within and outside the Corporation's control could have a significant bearing on its ability to obtain additional financing or on the generation of additional revenues.

These quarterly consolidated financial statements do not include any adjustments related to the carrying values and classifications of assets and liabilities that would be necessary should the Corporation be unable to continue as a going concern.

Liquidities

As at,	30-Apr-22	31-Oct-21	Change	
			\$ ¹	% ²
Cash	5,226	2,043	3,183	156%
Trade and other receivables	2,295	1,798	497	28%
Inventory	7,810	7,675	135	2%
Trade accounts payables	3,197	7,320	(4,123)	-56%
Working Capital	9,626	(2,984)	12,610	423%

1. A positive variance represents a positive impact, and a negative variance represents a negative impact to the balance sheet items
2. Percentage change is presented in relative values

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Cash and liquidities at the end of Q2-22 stood at \$5.2M as compared to \$2 million at the start of the year representing a \$3.2 million increase. Our working capital as at the end of Q2-22 stood at \$9.6 million as compared to a \$3 million deficit as at YE-21 representing a \$12.6 million improvement.

Following a series of successful financing in FY-21 and FY-22 we have secured significant capital to strengthen our balance sheet and our cash position and provide liquidities to support the growth of our new Respiriology franchise and support the costs related to our new corporate and sales structure (See "Business Overview") aimed at capturing the significant market opportunities for Redesca, Enerzair and Aetectura. As evidenced by our recent quarterly performance, the contribution of these products will materially impact the Corporation's revenues and gross margins going forward, and consequently Valeo is determined on leveraging the commercial potential of its current product portfolio and especially the \$150+ million combined peak sales potential of Redesca, Enerzair and Aetectura. Leveraging our commercial assets, as well as securing other business development opportunities that can 1) contribute immediately to our results, and 2) allow Valeo to reach EBITDA profitability over the coming year, is of the upmost importance for Valeo's management.

Entering into in-licensing agreements with pharmaceutical companies necessitates the payment of up-front amounts, milestone payments as well as all the costs normally associated with preparing for the launch of a pharmaceutical product. Going forward, Valeo intends to fund these in-licensing agreements with a combination of cash, cash from operations, equity provided by current and new shareholders, as well as convertible or non-convertible debt if required. As funding requirements to acquire product rights vary widely depending upon the nature and potential of the product and the in-licensing agreement, the Corporation intends to seek funding on a project-by-project basis and to prioritize non-dilutive financing instruments as a mean of funding new product acquisition. Funding requirements for products under discussion vary from \$nil to \$20 million.

The Corporation anticipates that the licensing of additional products rights and/or the commencement of additional product distribution agreements currently under advanced negotiations would materially impact Valeo's results. Should Valeo be successful in completing such transactions, which is still uncertain at this time, they would significantly increase our existing revenues and margins, as well as provide material operational synergies by leveraging our existing HO and commercial platform. These initiatives would contribute to accelerate our profitability. Historically, Valeo has been very successful in entering into licensing agreements and securing product rights by limiting funding requirements for such transactions. The existing and projected profitability of products rights currently being considered provides significant flexibility for deal structuring and to use licensing terms as a mean of funding deal economics and covering the bulk of the licensors expected financial returns.

Financial Risk Factors

(a) Market risk

(i) Currency risk

The Corporation is exposed to financial risks that arise from fluctuations in foreign exchange rates and the degree of volatility of these rates. The Corporation has an investment in a U.S. subsidiary however this subsidiary is currently not active. The Corporation does not hold financial derivatives to manage the fluctuation of these risks. As at April 30, 2022, a 5% increase/decrease in the USD/CAD would have a \$162 impact on net loss and equity (\$262 as at October 31, 2021). The following presents the accounts that are exposed to foreign exchange volatility:

As at,	April 30, 2022		October 31, 2021	
	Foreign Currency	CDN equivalent	Foreign Currency	CDN equivalent
Cash – USD	3,358	4,295	612	759
Accounts receivables and other assets – USD	102	130	-	-
Accounts payable and accrued liabilities – USD	11	14	2,455	3,040

OCI would not be materially impacted in the above situation.

(ii) Cash flow and fair value interest rate risk

The Corporation is exposed to fluctuation in its future cash flows arising from changes in interest rates through its variable rate exposure under its operating line of credit. Convertible and non-convertible debentures or long-term loans negotiated at a fixed rate expose the Corporation to fair value interest rate risk.

(b) Credit Risk

The Corporation considers its maximum credit risk to be based on the following financial assets: cash, trade and other receivables. Credit-risk arises from cash and deposits with banks and financial institutions. The Corporation reduces this risk by dealing with creditworthy financial institutions. Credit risk also results from the possibility that a loss may occur from the failure of another party to adhere to payment terms. To lower this risk, the Corporation's extension of credit is based on an evaluation of each customer's financial condition. Management reviews the aging of trade accounts receivable and other factors relating to the risk that customer accounts may not be paid in full and, when appropriate, reduces the carrying value to provide for possible loss. No loss has been charged to earnings in the last two fiscal years.

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The Corporation sells its products through a small number of wholesalers and retail pharmacy chains. The Corporation has collection terms that vary based on the nature of products sold. Accordingly, when determining the percentage of receivables which are current, the Corporation considers all receivables under 30 days for Valeo Pharma Inc. and all receivables under 90 days for VPI Pharma Inc. As at April 30, 2022, 83% of trade accounts receivables were current (82% as at October 31, 2021). As at April 30, 2022, three customers accounted for 83% of the trade receivables (84% as at October 31, 2021). The Corporation believes that there is no unusual exposure associated with the collection of these receivables.

(c) Liquidity risk

Liquidity risk is the risk that the Corporation will not be able to meet its obligations as they fall due. The following are the contractual maturities of financial liabilities.

As at April 30, 2022	30 days to		3 months to		Total
	Less than 30 days	3 months	12 months	More than 12 months	
Accounts payable and accrued liabilities	4,046	189	320	-	4,555
Lease liability	16	31	125	2,203	2,375
Convertible debentures	-	750	2,294	31,027	34,071
Non-convertible debenture	350	-	-	-	350
	4,412	970	2,739	33,230	41,351

As at October 31, 2021	30 days to		3 months to		Total
	Less than 30 days	3 months	12 months	More than 12 months	
Accounts payable and accrued liabilities	8,369	580	1,006	-	9,955
Lease liability	16	31	125	2,297	2,469
Convertible debentures	-	-	213	1,879	2,092
Non-convertible debenture	-	3,754	1,802	-	5,556
	8,385	4,365	3,146	4,176	20,072

(d) Capital Structure Financial Policy

The Corporation's objectives for managing capital are: (i) to maintain a flexible capital structure which optimizes the cost/risk equation; and (ii) to manage capital in a manner which maximizes the interests of our shareholders.

The Corporation manages the capital structure and adjusts it considering changes in economic conditions and the risk characteristics of the underlying assets. The Corporation's capital structure is managed in conjunction with the capital structure and financial needs of the day-to-day operations. The Corporation currently funds the working capital requirements out of its internally generated cash flows and the use of credit facilities. To maintain or adjust the capital structure, the Corporation will work to secure new debt from its shareholders and expand the shareholder base with new participation that would make additional funds available. As at April 30, 2022 the Corporation is not subject to any externally imposed capital requirements.

Covid-19 Risk

An outbreak of a novel strain of coronavirus, identified as "COVID-19", was declared a global pandemic by the World Health Organization on March 11, 2020. In response, many countries have required entities to limit or suspend business operations and implemented travel restrictions and quarantine measures. These measures have disrupted the activities of many entities and have led to significant volatility in the global markets.

The Corporation's business may be negatively impacted by the COVID-19 pandemic, which has created, and continues to create, significant societal and economic disruptions.

The changing and rapidly-evolving effects of the COVID-19 pandemic – the duration, extent and severity of which are currently unknown – on investors, businesses, the economy, government bodies, society and the financial markets could, among other things, add volatility to the global stock markets and impact interest rate environments.

The COVID-19 pandemic and measures to prevent its spread may negatively impact the Corporation, its customers, counterparties, employees, third-party service providers and other stakeholders, as applicable, in a number of ways, including, but not limited to, by: (i) adversely affecting the business operations of the Corporation, including access to its products by patients, the Corporation's planned sales and marketing processes for its approved products and the Corporation's ability to source, evaluate and pursue acquisition opportunities; (ii) disrupting the Corporation's supply chain, including the manufacture and/or delivery of its products by third-party manufacturers on which the Corporation relies; (iii) adversely affecting local, national or international economies and employment levels; (iv) causing business interruptions, including as a result of steps taken by the Corporation in compliance with government recommendations and orders, such as requiring employee to work remotely, which may cause strain on such existing resources as information technology systems, and

VALEO PHARMA INC.

Management's Discussion and Analysis for the three and six-month periods ended April 30, 2022

suspension of all non-essential travel; (v) disrupting public and private infrastructure, including communications and financial services, which could disrupt the Corporation's normal business operations; (vi) adversely affecting the Corporation's ability to comply with the covenants in its credit facility or requiring modifications to such covenants, for which there can be no assurance that such modifications would be provided; (vii) disrupting health care delivery; (viii) disrupting operations at Health Canada, which may result in delays in reviews and approvals, including with respect to products for which the Corporation has made or may make new drug submissions; (ix) disrupting operations at public or private payors and related agencies, such as CADTH, PMPRB, pCPA, which may result in delays in gaining access or reimbursement with respect to products for which the Corporation has made or may make submissions.

Risk Factors

For a detailed discussion of additional risk factors, please refer to the Company's latest Annual Information Form on SEDAR at www.sedar.com

VALEO PHARMA INC.

Management's Discussion and Analysis for the three and six-month periods ended April 30, 2022

Interim Condensed Consolidated Financial Statements (Unaudited)

Valeo Pharma Inc.

April 30, 2022
Second quarter fiscal year 2022

NOTICE OF NO AUDITOR REVIEW OF INTERIM FINANCIAL STATEMENTS Under National Instrument 51-102, Part 4, subsection 4.3(3)(a), if an auditor has not performed a review of the interim financial statements, the statements must be accompanied by a notice indicating that the financial statements have not been reviewed by an auditor. The accompanying unaudited interim financial statements of the Corporation have been prepared by management and are the responsibility of the Corporation's management. The Corporation's independent auditor has not performed a review or an audit of these interim financial statements.

Valeo Pharma Inc.

Interim Condensed Consolidated Statements of Financial Position

(Unaudited)

(All amounts in thousands of Canadian dollars)

<u>As at,</u>	<u>Notes</u>	<u>April 30, 2022</u>	<u>October 31, 2021</u>
ASSETS			
Current			
Cash		5,226	2,043
Trade and other receivables	4	2,295	1,798
Inventory	5	7,810	7,675
Prepaid expenses and deposits		816	834
Total current assets		16,147	12,350
Property and equipment	6	1,354	1,174
Right of use asset	7	924	967
Intangible assets	8	5,989	6,539
Total assets		24,414	21,030
LIABILITIES AND SHAREHOLDERS' EQUITY (DEFICIT)			
Current			
Trade accounts payable	10	3,197	7,320
Other accounts payable and accrued liabilities	10	1,329	2,635
Accrued interest on debentures		270	266
Provisions	11	29	214
Lease liability	12	48	45
Convertible debentures	13	709	-
Non-convertible debentures	14	338	4,854
Derivative warrant liability	15	601	-
Total current liabilities		6,521	15,334
Lease liability	12	1,140	1,165
Convertible debentures	13	19,724	1,605
Derivative warrant liability	15	-	582
Defined benefit obligations		199	291
Total liabilities		27,584	18,977
SHAREHOLDERS' EQUITY (DEFICIT)			
Share capital	16	25,720	24,616
Warrants	16	3,778	3,769
Contributed surplus		7,252	2,697
Accumulated other comprehensive loss		(249)	(319)
Deficit		(39,671)	(28,710)
Total shareholders' equity (deficit)		(3,170)	2,053
Total liabilities and shareholders' equity (deficit)		24,414	21,030

Going concern (note 1); Related Party Transactions (note 24); Commitments (note 27); Subsequent events (note 28)

/s/ "Steven Saviuk" _____, Director

/s/ "Richard Mackay" _____, Director

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

Valeo Pharma Inc.

Interim Condensed Consolidated Statements of Loss and Comprehensive Loss (Unaudited)

*(All amounts in thousands of Canadian dollars, except for share and per share amounts)
For the three- and six-month periods ended April 30, 2022 and 2021*

Notes	Three months ended April 30		Six months ended April 30	
	2022	2021	2022	2021
Revenues	4,768	2,647	9,009	4,508
Cost of Goods Sold	18	3,109	1,938	5,941
Gross Profit	1,659	709	3,068	1,094
Expenses				
Sales and marketing	19	3,539	949	7,331
General and administrative	20	964	881	2,229
Medical affairs and regulatory	21	845	257	1,859
Share based compensation	16 b	222	309	444
Profit Sharing		32	1	43
Total operating expenses	5,602	2,397	11,906	4,359
Operating loss	(3,943)	(1,688)	(8,838)	(3,265)
Other expenses (income)				
Financial	22	1,178	213	2,174
Other income	23	(40)	(34)	(70)
Unrealized loss on derivative warrant liability	15	17	-	19
Total other expenses (income)	1,155	179	2,123	328
Net loss for the period	(5,098)	(1,867)	(10,961)	(3,593)
Other comprehensive income (loss)				
Exchange differences on translating foreign operations		(6)	6	(4)
Defined benefit plan, net actuarial gain		74	93	74
Total comprehensive loss for the period	(5,030)	(1,768)	(10,891)	(3,489)
Loss per share:				
Basic and diluted		(0.06)	(0.03)	(0.14)
Weighted average number of shares outstanding	80,661,530	65,565,241	79,721,375	65,039,982

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

Valeo Pharma Inc.

Interim Condensed Consolidated Statements of Changes in Shareholders' Equity (Deficit)

(All amounts in thousands of Canadian dollars)

For the six months ended April 30, 2022 and 2021

	Notes	Share Capital		Accumulated Other Comprehensive Loss			Deficit	Total
		Common Shares	Warrants	Contributed surplus	Defined benefit plan	Foreign exchange translation		
Balance as at October 31, 2020		15,024	1,333	1,611	(387)	(35)	(14,477)	3,069
Net loss		-	-	-	-	-	(3,593)	(3,593)
Other comprehensive income		-	-	-	93	11	-	104
Share based compensation	16 b	-	-	414	-	-	-	414
Stock options exercised	16 a	109	-	(39)	-	-	-	70
Equity instruments issued to consultants		34	-	64	-	-	-	98
Compensation options exercised		167	29	(64)	-	-	-	132
Warrants issued		-	531	-	-	-	-	531
Warrants exercised		874	(103)	-	-	-	-	771
Issue costs		(40)	(11)	-	-	-	-	(51)
Balance as at April 30, 2021		16,168	1,779	1,986	(294)	(24)	(18,070)	1,545
 Balance as at October 31, 2021		 24,616	 3,769	 2,697	 (294)	 (25)	 (28,710)	 2,053
Net loss		-	-	-	-	-	(10,961)	(10,961)
Other comprehensive income		-	-	-	74	(4)	-	70
Share based compensation	16 b	-	-	444	-	-	-	444
Stock options exercised	16 a	123	-	(43)	-	-	-	80
Equity instruments issued to consultants		34	-	-	-	-	-	34
Issue costs		(267)	-	-	-	-	-	(267)
Broker's compensation units expired		93	9	(102)	-	-	-	-
Convertible debentures issued	13	-	-	4,431	-	-	-	4,431
Convertible debentures converted	13	1,121	-	(175)	-	-	-	946
Balance as at April 30, 2022		25,720	3,778	7,252	(220)	(29)	(39,671)	(3,170)

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

Valeo Pharma Inc.

Interim Condensed Consolidated Statements of Cash Flow

(Unaudited)

(All amounts in thousands of Canadian dollars)

For the six months ended April 30, 2022 and 2021

	Notes	April 30, 2022	April 30, 2021
OPERATING ACTIVITIES:			
Net loss from operations		(10,961)	(3,593)
Adjustments:			
Depreciation and amortization	6,7,8	508	293
Share based compensation	16 b,c	444	414
Interest expense		807	144
Consulting fees paid by issuance of equity instruments		34	98
Defined benefit plan expense		(16)	(15)
Unrealized loss on foreign exchange		45	46
Unrealized loss on derivative warrant liability		19	-
Write down of inventory	18	-	3
Net change in non-cash operating working capital	17	(6,128)	(1,680)
Cash used by operating activities		(15,248)	(4,289)
INVESTING ACTIVITIES:			
Acquisition of property and equipment	6	(257)	(44)
Acquisition of intangible assets	8	(32)	(2,146)
Cash used by investing activities		(289)	(2,190)
FINANCING ACTIVITIES:			
Increase in convertible debentures	14	25,000	6,645
Repayment of non-convertible debentures	14	(4,807)	-
Payment of financing fees		(1,505)	(172)
Proceeds from issuance of shares	16	80	972
Principal repayment of lease liabilities	12	(94)	(47)
Cash provided by financing activities		18,674	7,398
Foreign exchange gain (loss) on cash		46	(105)
Increase in cash		3,183	814
Cash, beginning of period		2,043	2,836
Cash, end of period		5,226	3,650

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

Valeo Pharma Inc.

Notes to the Interim Condensed Consolidated Financial Statements

(Unaudited)

(All amounts in thousands of Canadian dollars, except for share, unit and per share amounts)

1. Presentation of Financial Statements and Going Concern

Description of the Business

Valeo Pharma Inc. (the “Corporation”) is a specialty pharmaceutical company that acquires or in-licenses brand and hospital specialty products for sale in Canada. Its head office, principal place of business and registered office is located at 16667 Hymus Blvd., Kirkland, Quebec, Canada. The Corporation’s wholly owned subsidiary VPI Pharmaceuticals Inc. (“VPI”) is located within the Corporation’s premises, and Valeo Pharma Corp (“Valeo USA”) is located in the United States (not active).

The Corporation is incorporated under the Canada Business Corporations Act. The Corporation’s shares, warrants and debentures trade on the Toronto Stock Exchange (TSX) under the symbol VPH, VPH.WT, VPH.WT.A and VPH.DB. The Corporation’s shares are also listed on the Frankfurt Stock Exchange (“FSE”) under the symbol VP2 and on the US OTCQB market under the symbol VPHIF.

Statement of Compliance

These unaudited interim condensed consolidated financial statements of the Corporation have been prepared for the six-month period ended April 30, 2022 in accordance with International Financial Reporting Standards (“IFRS”) as issued by the International Accounting Standards Board (“IASB”). These unaudited interim condensed consolidated financial statements have been prepared in accordance with those IFRS standards and interpretations of the International Financial Reporting Interpretations Committee issued and effective or issued and early adopted as at the time of preparing these statements. These unaudited interim condensed consolidated financial statements do not include all the information required for full disclosure in the annual financial statements and should be read in conjunction with the annual consolidated financial statements for the year ended October 31, 2021 as they follow the same accounting policies and methods of application.

These unaudited interim condensed consolidated financial statements were approved and authorized for issuance by the Board of Directors on June 13, 2022.

Going Concern

These unaudited interim condensed consolidated financial statements have been prepared on a going-concern basis, which implies that the Corporation will continue realizing its assets and discharging its liabilities in the normal course of business for the foreseeable future. As reflected in the unaudited interim condensed consolidated financial statements, the Corporation is in the process of ramping up its commercial activities and has not yet achieved profitability. During the six-month period ended April 30, 2022, the Corporation incurred a net loss of \$10,961 and used cash in operations of \$15,248. As at April 30, 2022, the Corporation had a working capital surplus of \$9,626. This raises significant doubt about the Corporation’s ability to continue as a going concern.

Accordingly, the ability of the Corporation to realize the carrying value of its assets and continue operations as a going concern is dependent upon its ability to obtain additional financing and ultimately on generating future profitable operations. Management anticipates that the generation of additional revenues out of its existing products and the commercialization of new products will enable the Corporation to achieve profitability. There is no assurance that any of these initiatives will be successful. Factors within and outside the Corporation’s control could have a significant bearing on its ability to obtain additional financing or on the generation of additional revenues.

These unaudited interim condensed consolidated financial statements do not include any adjustments related to the carrying values and classifications of assets and liabilities that would be necessary should the Corporation be unable to continue as a going concern. Such adjustments could be material.

Covid-19

An outbreak of a novel strain of coronavirus, identified as “COVID-19”, was declared a global pandemic by the World Health Organization on March 11, 2020. In response, many countries have required entities to limit or suspend business operations and implemented travel restrictions and quarantine measures. These measures have disrupted the activities of many entities and have led to significant volatility in the global markets.

The Corporation’s business may be negatively impacted by the COVID-19 pandemic, which has created, and continues to create, significant societal and economic disruptions.

The changing and rapidly-evolving effects of the COVID-19 pandemic – the duration, extent and severity of which are currently unknown – on investors, businesses, the economy, government bodies, society and the financial markets could, among other things, add volatility to the global stock markets and impact interest rate environments.

Valeo Pharma Inc.

Notes to the Interim Condensed Consolidated Financial Statements

(Unaudited)

(All amounts in thousands of Canadian dollars, except for share, unit and per share amounts)

1. Presentation of Financial Statements and Going Concern – cont'd

The COVID-19 pandemic and measures to prevent its spread may negatively impact the Corporation, its customers, counterparties, employees, third-party service providers and other stakeholders, as applicable, in a number of ways, including, but not limited to, by: (i) adversely affecting the business operations of the Corporation, including access to its products by patients, the Corporation's planned sales and marketing processes for its approved products and the Corporation's ability to source, evaluate and pursue acquisition opportunities; (ii) disrupting the Corporation's supply chain, including the manufacture and/or delivery of its products by third-party manufacturers on which the Corporation relies; (iii) adversely affecting local, national or international economies and employment levels; (iv) causing business interruptions, including as a result of steps taken by the Corporation in compliance with government recommendations and orders, such as requiring employee to work remotely, which may cause strain on such existing resources as information technology systems, and suspension of all non-essential travel; (v) disrupting public and private infrastructure, including communications and financial services, which could disrupt the Corporation's normal business operations; (vi) adversely affecting the Corporation's ability to comply with the covenants in its credit facility or requiring modifications to such covenants, for which there can be no assurance that such modifications would be provided; (vii) disrupting health care delivery; (viii) disrupting operations at Health Canada, which may result in delays in reviews and approvals, including with respect to products for which the Corporation has made or may make new drug submissions; (ix) disrupting operations at public or private payors and related agencies, such as CADTH, PMPRB, pCPA, which may result in delays in gaining access or reimbursement with respect to products for which the Corporation has made or may make submissions.

2. Summary of Significant Accounting Policies

Basis of consolidation

These unaudited interim consolidated financial statements consolidate those accounts of the Corporation and its wholly owned subsidiaries (the "Group"). All subsidiaries have an annual reporting date of October 31. All transactions and balances between Group companies are eliminated upon consolidation, including unrealized gains and losses on transactions between Group companies. Where unrealized losses on intra-group asset sales are reversed on consolidation, the underlying asset is also tested for impairment from a group perspective.

Amounts reported in the financial statements of subsidiaries have been adjusted where necessary to ensure consistency with the accounting policies adopted by the Corporation. Profit or loss and other comprehensive income ("OCI") of subsidiaries acquired or disposed of during the period are recognized from the effective date of acquisition, or up to the effective date of disposal, as applicable.

Basis of measurement

These unaudited interim condensed consolidated financial statements have been prepared on a going-concern basis, under the historical cost convention, except for the revaluation of certain financial assets and financial liabilities to fair value including the derivative warrant liability.

3. Use of Estimates and Judgements

The preparation of the unaudited interim condensed consolidated financial statements requires management to undertake several judgements, estimates and assumptions about recognition and measurement of assets, liabilities, income and expenses. The actual results may differ from these judgements and estimates. These estimates and judgements are based on management's best knowledge of the events or circumstances and actions the Corporation may take in the future. The estimates are reviewed on an ongoing basis. Information about the significant judgements, estimates and assumptions that have the most significant effect on the recognition and measurement of assets, liabilities, income and expenses are discussed in Note 3 of the Corporation's 2021 audited annual consolidated financial statements and are still applicable for the six-month period ended April 30, 2022.

4. Trade and Other Receivables

As at	April 30, 2022	October 31, 2021
Trade and other receivables	2,044	1,473
Receivables from a related party	-	1
Sales taxes receivables	251	324
	2,295	1,798

Valeo Pharma Inc.

Notes to the Interim Condensed Consolidated Financial Statements

(Unaudited)

(All amounts in thousands of Canadian dollars, except for share, unit and per share amounts)

5. Inventory

As at	April 30, 2022	October 31, 2021
Finished good	7,540	7,407
Raw material	270	268
	7,810	7,675

6. Property and Equipment

	Leasehold improvements	Computer equipment	Equipment and furniture	Security vault	Total
Cost as at October 31, 2021	710	339	491	196	1,736
Additions	24	213	20	-	257
Cost as at April 30, 2022	734	552	511	196	1,993
Accumulated depreciation as at October 31, 2021	94	267	150	51	562
Depreciation	33	11	30	3	77
Accumulated depreciation as at April 30, 2022	127	278	180	54	639
Net carrying value as at April 30, 2022	607	274	331	142	1,354

7. Right of Use Asset

	Cost	Depreciation	Carrying value
Balance as at October 31, 2021	1,003	(36)	967
Additions	-	(43)	(43)
Balance as at April 30, 2022	1,003	(79)	924

8. Intangible Assets

	Submission costs	License fee	Total
Balance as at October 31, 2021	2,810	3,729	6,539
Additions	32	-	32
Reclassification	(194)	-	(194)
Amortization	(141)	(247)	(388)
Balance as at April 30, 2022	2,507	3,482	5,989

9. Operating Loan

On April 5, 2022, the Corporation renewed its revolving demand credit facility with its present lender. At all times, borrowed amounts under the facility will not exceed the lesser of \$2,500 and the total of (a) assigned credit balances for the Corporation plus (b) 80% of Canadian and US based accounts receivables (85% for investment grade receivables) of the Corporation net of over 90-day accounts, related parties accounts and all other accounts not valued by the lender plus (c) 50% of the inventory value up to a maximum of \$1,250.

The lender will make the facility available by way of prime rate-based loans in CAD\$, United States base rate (“USBR”) loans in USD\$ and stand-by letters of guarantee in CAD\$. The interest rates for prime based loans are prime rate plus 0.75% per annum; and USBR plus 0.75% per annum for USBR loans. For letters of guarantee the rate applicable will be that set out in the letter of credit indemnity agreement applicable to the issued letter of guarantee. As at April 30, 2022 the operating loan was unused, and the Corporation had a \$2,200 letter of guarantee issued in favour of one of its licensors to cover any financial obligations under its supply agreement.

Valeo Pharma Inc.

Notes to the Interim Condensed Consolidated Financial Statements

(Unaudited)

(All amounts in thousands of Canadian dollars, except for share, unit and per share amounts)

10. Accounts Payable and Accrued Liabilities

As at	April 30, 2022	October 31, 2021
Trade accounts payable	3,197	7,320
Other accounts payable and accrued liabilities	1,228	2,535
Payables to related parties (i)	101	100
	4,526	9,955

(i) Included in Payables to related parties

Consulting fees owed to a company controlled by an officer	19	11
Expenses owed to officers, employees and consultants in the normal course of business	82	89

11. Provisions

The following table presents the changes in the provision for product returns, pricing rebates and chargebacks during the period:

	Total
Balance as at October 31, 2021	214
Charges	-
Utilization	(185)
Balance as at April 30, 2022	29

12. Lease Liability

The following table presents the changes in the lease liability during the periods:

	Six months ended April 30, 2022	Year ended October 31, 2021
Balance as at start of period	1,210	295
Lease modification	-	949
Interest expense	72	80
Lease payments	(94)	(114)
Balance as at end of period	1,188	1,210

Which consists of		
Current lease liability	48	45
Non-current lease liability	1,140	1,165

Valeo Pharma Inc.

Notes to the Interim Condensed Consolidated Financial Statements

(Unaudited)

(All amounts in thousands of Canadian dollars, except for share, unit and per share amounts)

13. Convertible debentures

	Note	Six months ended April 30, 2022	Year ended October 31, 2021
Balance as at start of period		1,605	1,504
Additions	a)	25,000	-
Fair value of conversion option allocated to equity	a)	(4,431)	-
Issue costs	a)	(1,240)	-
Transaction costs		-	-
Transaction costs amortization		109	9
Accretion expense	c)	334	92
Conversion into shares	b)	(944)	-
Balance as at close of period		20,433	1,605
Which consists of			
Current convertible debentures		709	-
Non-current convertible debentures		19,724	1,605

- a) During the first quarter of fiscal year 2022, the Corporation closed a bought deal private placement (the "Offering") of \$15.0 million aggregate principal amount of 12.0% convertible unsecured debentures (the "Debentures") due December 31, 2024 (the "Maturity Date") at a price of \$1,000 (the "Offering Price") per Debenture. The Corporation also closed a concurrent \$10.0 million private placement of convertible unsecured debentures issued on the same terms as those issuable pursuant to the Offering (the "Concurrent Private Placement"), resulting in gross proceeds from the Offering and Concurrent Private Placement of \$25.0 million. The Corporation issued a total of 25,000 Debentures accruing interest at the rate of 12% per annum payable quarterly beginning on March 31, 2022. At the holders' option, the Debentures may be converted into common shares of the Corporation at any time and from time to time, up to the Maturity Date, at a conversion price of \$1.15 per common share.

The Corporation valued the liability component of the debentures by calculating the present value of the principal and interest payments, discounted at a rate of 20%, being management's best estimate of the rate that a non-convertible debenture with similar terms would bear. The equity component consists of the conversion option. On initial recognition, the liability component was \$20,569, and the equity component (conversion options) was \$4,431. Transaction costs of \$1,240 were netted against the liability component and will be amortized using the effective interest method over the term of the debenture. A further \$260 in transaction costs, related to the equity component of the derivative liability, was capitalized to share issue costs.

- b) During the second quarter of fiscal year 2022, \$944 of convertible debentures issued in February 2020, \$175 of equity component and \$2 of interest payable were converted into \$1,121 of share capital.
- c) During the six-month period ended April 30, 2022, the debentures accrued interest of \$1,597 included in financial expense on the Statement of Loss. This amount includes an accretion expense of \$334. A total of \$258 is included in accrued interest on the Statement of Financial Position.

14. Non-convertible debentures

	Note	Six months ended April 30, 2022	Year ended October 31, 2021
Balance as at start of period		4,854	1,463
Additions		-	6,645
Rewards	a)	(4,807)	(3,200)
Derivative warrant liability		-	(442)
Transaction costs		27	(124)
Accretion expense	c)	264	512
Balance as at end of period		338	4,854

Valeo Pharma Inc.

Notes to the Interim Condensed Consolidated Financial Statements

(Unaudited)

(All amounts in thousands of Canadian dollars, except for share, unit and per share amounts)

14. Non-convertible debentures – cont'd

- a) Non-convertible debentures issued in April 2021 representing a total of \$2.86 million were exchanged for new convertible debentures in December 2021. The remaining non-convertible debentures issued in April 2021 representing \$585 were reimbursed in December 2021.
- Non-convertible debentures issued in July 2020 representing a total of \$805 were exchanged for new convertible debentures in December 2021. Non-convertible debentures issued July 2020 representing a total of \$557 were reimbursed in March 2022.
- b) The remaining non-convertible debentures issued July 2020 representing \$338 were reimbursed in May 2022 (see Subsequent Event note 28).
- c) During the six-month period ended April 30, 2022, the debentures accrued interest of \$326 included in financial expense on the Statement of Loss. This amount includes an accretion expense of \$264. A total of \$12 is included in accrued interest on the Statement of Financial Position.

15. Derivative warrant liability

The derivative warrant liability results of unsecured non-convertible debentures issued during the prior period. The balances represent the evaluation of the liability at the end of the respective periods.

The following table details the changes in the Corporation's derivatives warrant liability:

	Number	\$			
Balance at October 31, 2021	1,336,700	582			
Revaluation of derivative warrant liability	-	19			
Balance at April 30, 2022	1,336,700	601			
Classified as current liability	1,336,700	601			
Classified as long-term liability	-	-			
Number of Warrants	Issue date	Expiry date	Exercise price	Fair value of warrants	Remaining contractual life in years
1,336,700	April 26, 2021	April 26, 2023	1.25	0.45	0.99

The derivative warrants liability evaluation was performed using a Black-Scholes option pricing model with a risk-free rate of 2.53%; a volatility of 62.81%; an expected life of 2 years; an exercise price of \$1.25 with a nil expected dividend and forfeiture rate.

16. Share Capital and Other Equity Instruments

a) Share capital

The Authorized Share Capital is composed of an Unlimited number of Class "A" shares, voting, participating with no par value. The issued and outstanding share capital is as follows:

	Note	Number	\$
Balance as at October 31, 2020		64,055,359	15,024
Prospectus costs		-	4
Exercise of stock options	b)	385,810	110
Exercise of warrants	d)	1,284,575	873
Compensation options exercised	e)	273,875	167
Shares issued as compensation		30,731	34
Issue costs		-	(44)
Balance as at April 30, 2021		66,030,350	16,168

Valeo Pharma Inc.

Notes to the Interim Condensed Consolidated Financial Statements

(Unaudited)

(All amounts in thousands of Canadian dollars, except for share, unit and per share amounts)

16. Share Capital and Other Equity Instruments – *cont'd*

	Note	Number	\$
Balance as at October 31, 2021		78,800,174	24,616
Shares issue costs		-	(267)
Exercise of stock options	b)	200,000	123
Conversion of debentures		2,603,419	1,121
Compensation options expired		-	93
Shares issued as compensation		45,505	34
Balance as at April 30, 2022		81,649,098	25,720

b) Share option issuance and compensation expense

The Corporation has an equity-settled stock option incentive plan (the “Plan”) for directors, officers, employees, and consultants to enable the purchase of common shares of the Corporation. The exercise price for these shares cannot be less than the lowest price permitted by applicable regulatory authorities. Subject to the terms of the Plan, the Board of Directors may, from time to time, grant options to participants to purchase that number of shares that they determine, in their absolute discretion. The shares subject to each option shall become available for purchase by the participant from the Corporation on the date or dates determined by the Board of Directors when the option is granted. In the event of the termination of an optionee who is either an employee, director or officer, the Board of Directors may extend the period during which the optionee may exercise the options provided such period does not exceed three months from termination of employment or duties as director. The number of shares subject to an option shall not exceed 10% of the total issued and outstanding Common shares; and no one optionee shall be granted options that exceed 5%, 2% for any one consultant and 2% for any one person retained to provide investor relation services of the issued and outstanding common shares of the Corporation (on a non-diluted basis), during a 12-month period.

Changes in outstanding options were as follows:

	Six months ended April 30, 2022		Year ended October 31, 2021	
	Number	Weighted Average Exercise Price	Number	Weighted Average Exercise Price
Options outstanding, beginning of period	6,544,722	\$0.84	4,275,532	\$0.47
Granted	462,500	\$0.66	2,940,000	\$1.28
Forfeited	(250,000)	\$1.10	(50,000)	\$0.58
Cancelled/expired during the period	(118,750)	\$0.57	(180,000)	\$0.66
Exercised	(200,000)	\$0.40	(440,810)	\$0.22
Options outstanding, end of period	6,438,472	\$0.80	6,544,722	\$0.84
Options exercisable, end of period	3,595,142	\$0.59	3,870,000	\$0.57

The following options were granted in the six months ended April 30, 2022:

Number	Notes	Date of grant	Expiry date	Exercise price	Fair value
462,500	i	April 27, 2022	April 27, 2029	\$0.66	\$0.30

i) Vest 50% on each anniversary of the grant date

The fair values of the options granted during the third quarter were estimated using the Black-Scholes option pricing model, with the following assumptions:

Risk-free interest rate	2.64%
Volatility factor	62.87%
Expected life	7 years
Expected dividend rate	0%
Forfeiture rate	10%

Valeo Pharma Inc.

Notes to the Interim Condensed Consolidated Financial Statements

(Unaudited)

(All amounts in thousands of Canadian dollars, except for share, unit and per share amounts)

16. Share Capital and Other Equity Instruments - cont'd

The expected stock price volatility was estimated by using historical data from public companies in the same sector as the Corporation and over the period consistent with the duration of the award. The total share-based compensation for the six months ended April 30, 2022 was \$345 (2021 - \$414) recognized in contributed surplus reported in the Statement of Income.

c) Restricted stock units (RSUs)

On April 28, 2021, the Shareholders of the Corporation approved the implementation of an RSU equity incentive plan (the "RSU Plan"), which provides for the granting to directors, officers, employees and consultants of the Corporation ("Eligible Participants") non-transferable Restricted Share Units, Performance Share Units, Deferred Share Units, or Other Share-Based Awards, or any combination thereof (the "RSU Awards"). The purpose of this RSU Plan is to allow for certain discretionary bonuses and similar awards as an incentive and reward for selected Eligible Participants related to the achievement of long-term financial and strategic objectives of the Corporation and the resulting increases in shareholder value. This RSU Plan is intended to promote a greater alignment of interests between the shareholders of the Corporation and the selected Eligible Participants by providing an opportunity to acquire Shares as long-term investments and equity interests in the Corporation. The number of Shares reserved for issuance and which will be available for issuance pursuant to Awards granted under the RSU Plan will equal 5% of the issued and outstanding Shares of the Corporation from time to time, provided that the aggregate number of Shares available for issuance to Insider Participants under this RSU Plan, together with all other equity incentive plans of the Corporation (including its Share Option Plan), may not exceed 10% of the issued Shares at any given time. The total share-based compensation for the six months ended April 30, 2022 was \$99 (2021 - \$0) recognized in contributed surplus reported in the Statement of Income.

The following schedule presents the RSUs issued at the end of the respective periods:

	Number of shares	Market price at time of grant
Balance as at October 31, 2021	475,000	\$1.12
Issued during the period	175,926	\$0.54
Vested	-	-
Balance as at April 30, 2022	650,926	\$0.96

d) Warrants

The following schedule presents the common shares issuable on exercise of all warrants outstanding at the end of the respective periods:

	Number of shares	Weighted Average Exercise Price
Balance as at October 31, 2021	24,658,182	\$1.03
Issued during the period	-	-
Exercised	-	-
Balance as at April 30, 2022	24,658,182	\$1.03

e) Compensation Options

In connection with the issuance of units in both September 2020 and June 2021, the Corporation issued compensation units entitling the holder to purchase 1 share and ½ warrant and 1 share and 1 warrant, respectively, subject to the same terms and conditions as the original unit offering. The September 2020 compensation units expired on March 10, 2022.

The following schedule presents the common shares and warrants issuable on exercise of compensation options:

	Number of shares	Number of warrants	Weighted Average Exercise Price
Balance as at October 31, 2021	1,140,673	955,336	\$1.23
Expired	(370,673)	(185,336)	\$1.20
Balance as at April 30, 2022	770,000	770,000	\$1.25

Valeo Pharma Inc.

Notes to the Interim Condensed Consolidated Financial Statements

(Unaudited)

(All amounts in thousands of Canadian dollars, except for share, unit and per share amounts)

17. Other Cash Flow Information

	Six months ended April 30,	
	2022	2021
(Increase) decrease in non-cash assets related to operations		
trade receivables	(568)	(276)
other receivables	75	(289)
inventory	(135)	(4,558)
prepaid expenses	215	(75)
Increase (decrease) in non-cash liabilities related to operations		
accounts payable and accrued liabilities	(5,530)	3,465
provision for chargebacks and returns	(185)	53
	(6,128)	(1,680)

The amounts above exclude the foreign exchange gain or loss reported in the corresponding items of the Interim Condensed Consolidated Statements of Financial Position.

18. Cost of Goods Sold

	Three months ended April 30,		Six months ended April 30,	
	2022	2021	2022	2021
Finished goods	2,836	1,804	5,396	3,146
Freight, storage and distribution fees	142	71	298	139
Amortization of intangible assets	124	123	247	126
Write down of inventory	7	(60)	-	3
	3,109	1,938	5,941	3,414

19. Sales and Marketing Expenses

	Three months ended April 30,		Six months ended April 30,	
	2022	2021	2022	2021
Employee compensation	2,341	665	4,894	1,078
Sales expenses	343	154	540	326
Marketing expenses	855	130	1,897	191
	3,539	949	7,331	1,595

20. General and Administrative Expenses

	Three months ended April 30,		Six months ended April 30,	
	2022	2021	2022	2021
Employee compensation	537	416	1,148	720
Administrative expenses	311	342	829	643
Investor relations expenses	54	93	132	406
Depreciation of property and equipment	40	18	77	35
Depreciation of right of use asset	22	12	43	21
	964	881	2,229	1,825

Valeo Pharma Inc.

Notes to the Interim Condensed Consolidated Financial Statements

(Unaudited)

(All amounts in thousands of Canadian dollars, except for share, unit and per share amounts)

21. Medical Affairs and Regulatory Expenses

	Three months ended April 30,		Six months ended April 30,	
	2022	2021	2022	2021
Employee compensation	418	115	861	192
Patient support programs	138	80	275	211
Advisory boards and other expenses	225	3	582	8
Amortization of intangible assets	64	59	141	113
	845	257	1,859	524

22. Financial Expenses

	Three months ended April 30,		Six months ended April 30,	
	2022	2021	2022	2021
Interest on debentures	888	130	1,468	228
Effective interest on debentures	274	47	598	101
Lease interest	36	8	72	17
Bank and other interest	-	3	26	8
Bank charges	11	8	23	12
Foreign exchange fluctuation	(31)	17	(13)	40
	1,178	213	2,174	406

23. Other Income

	Three months ended April 30,		Six months ended April 30,	
	2022	2021	2022	2021
Service income	31	34	55	77
Interest income	9	-	15	1
	40	34	70	78

Service income represents quality control, legal and finance services charged to a related company renting office space at the Corporation's head office.

24. Related Party Transactions

The following table presents the related party transactions presented in the Statement of Loss for the respective periods:

	Three months ended April 30,		Six months ended April 30,	
	2022	2021	2022	2021
Key management salary and benefits	367	295	962	520
Directors and employee stock option compensation	222	309	444	414
Consulting fees paid to a company controlled by an officer	59	46	146	91
Service income	31	34	55	77
	679	684	1,607	1,102

Valeo Pharma Inc.

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(All amounts in thousands of Canadian dollars, except for share, unit and per share amounts)

24. Related Party Transactions – cont'd

The following table represents the related party transactions presented in the Statement of Financial Position as at:

As at,	April 30, 2022	October 31, 2021
Amounts owed to key management, officers and directors		
Consulting fees	-	11
Convertible debentures	538	231
Accrued interest on convertible debentures	8	5
Non-convertible debentures	15	436
Accrued interest on non-convertibles debentures	1	14
Amounts owed to Manitex, a shareholder of the Corporation		
Non-convertible debentures	-	15
Accrued interest on non-convertible debentures	-	1
Amounts owed to 100079 Canada Inc., a shareholder of the Corporation		
Convertible debentures	1,285	955
Accrued interest on convertible debentures	15	24
Non-convertible debentures	-	2,041
Accrued interest on non-convertible	-	100

25. Financial Instruments

Short term financial instruments, comprising trade receivables, other receivables, bank indebtedness, accounts payable and accrued liabilities, loans and non-convertible debentures are carried at amortized cost, which, due to their short-term nature, approximates their fair value. Long term financial instruments consisting of convertible debentures are accounted for at amortized cost using the effective interest rate method, which corresponds to the fair value.

The Corporation categorizes its financial assets and liabilities measured at the fair value into one of three different levels depending on the observation of the inputs used in the measurement. As at April 30, 2022, the Corporation carried derivative warrants defined as level 3 financial instruments (see note 15). There were no transfers between levels during the period.

The three levels are defined as follows:

Level 1: Fair value is based on unadjusted quoted prices for identical assets or liabilities in active markets.

Level 2: Fair value is based on inputs other than quoted prices included within Level 1 that are observable for the asset or liability, either directly (i.e. prices) or indirectly (i.e. derived from prices); and

Level 3: Fair value is based on valuation techniques that require one or more significant unobservable inputs.

The fair value of a financial instrument is approximated by the consideration that would be agreed to in an arm's length transaction between willing parties and through appropriate valuation methods, but considerable judgement is required for the Corporation to determine the value. The actual amount that could be realized in a current market exchange could be different than the estimated value. The Corporation's activities expose it to a variety of financial risks: market risk (including currency risk and cash flow and fair value interest rate risk); credit risk and liquidity risk. The Corporation's overall risk management program focuses on the unpredictability of the financial market and seeks to minimize potential adverse effects on the Corporation's financial performance. The Corporation does not use derivative financial instruments to hedge these risks.

Valeo Pharma Inc.

Notes to the Interim Condensed Consolidated Financial Statements

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(All amounts in thousands of Canadian dollars, except for share, unit and per share amounts)

26. Financial Risk Factors

(a) Market risk

(i) Currency risk

The Corporation is exposed to financial risks that arise from fluctuations in foreign exchange rates and the degree of volatility of these rates. The Corporation has an investment in a U.S. subsidiary however this subsidiary is currently not active. The Corporation does not hold financial derivatives to manage the fluctuation of these risks. As at April 30, 2022, a 5% increase/decrease in the USD/CAD would have a \$162 impact on net loss and equity (\$262 as at October 31, 2021).

The following presents the accounts that are exposed to foreign exchange volatility:

As at,	April 30, 2022		October 31, 2021	
	Foreign Currency	CDN equivalent	Foreign Currency	CDN equivalent
Cash – USD	3,358	4,295	612	759
Accounts receivables and other assets – USD	102	130	-	-
Accounts payable and accrued liabilities – USD	11	14	2,455	3,040

OCI would not be materially impacted in the above situation.

(ii) Cash flow and fair value interest rate risk

The Corporation is exposed to fluctuation in its future cash flows arising from changes in interest rates through its variable rate exposure under its operating line of credit. Convertible and non-convertible debentures or long-term loans negotiated at a fixed rate expose the Corporation to fair value interest rate risk.

(b) Credit Risk

The Corporation considers its maximum credit risk to be based on the following financial assets: cash, trade and other receivables. Credit-risk arises from cash and deposits with banks and financial institutions. The Corporation reduces this risk by dealing with creditworthy financial institutions. Credit risk also results from the possibility that a loss may occur from the failure of another party to adhere to payment terms. To lower this risk, the Corporation's extension of credit is based on an evaluation of each customer's financial condition. Management reviews the aging of trade accounts receivable and other factors relating to the risk that customer accounts may not be paid in full and, when appropriate, reduces the carrying value to provide for possible loss. No loss has been charged to earnings in the last two fiscal years.

The Corporation sells its products through a small number of wholesalers and retail pharmacy chains. The Corporation has collection terms that vary based on the nature of products sold. Accordingly, when determining the percentage of receivables which are current, the Corporation considers all receivables under 30 days for Valeo Pharma Inc. and all receivables under 90 days for VPI Pharma Inc. As at April 30, 2022, 83% of trade accounts receivables were current (82% as at October 31, 2021). As at April 30, 2022, three customers accounted for 83% of the trade receivables (84% as at October 31, 2021). The Corporation believes that there is no unusual exposure associated with the collection of these receivables.

(c) Liquidity risk

Liquidity risk is the risk that the Corporation will not be able to meet its obligations as they fall due. The following are the contractual maturities of financial liabilities.

As at April 30, 2022	30 days				Total
	Less than 30 days	3 months	to 12 months	More than 12 months	
Accounts payable and accrued liabilities	4,046	189	320	-	4,555
Lease liability	16	31	125	2,203	2,375
Convertible debentures	-	750	2,294	31,027	34,071
Non-convertible debenture	350	-	-	-	350
	4,412	970	2,739	33,230	41,351

Valeo Pharma Inc.

Notes to the Interim Condensed Consolidated Financial Statements

(Unaudited)

(All amounts in thousands of Canadian dollars, except for share, unit and per share amounts)

26. Financial Risk Factors – cont'd

As at October 31, 2021					Total
	Less than 30 days	30 days to 3 months	3 months to 12 months	More than 12 months	
Accounts payable and accrued liabilities	8,369	580	1,006	-	9,955
Lease liability	16	31	125	2,297	2,469
Convertible debentures	-	-	213	1,879	2,092
Non-convertible debenture	-	3,754	1,802	-	5,556
	8,385	4,365	3,146	4,176	20,072

(d) Capital Structure Financial Policy

The Corporation's objectives for managing capital are: (i) to maintain a flexible capital structure which optimizes the cost/risk equation; and (ii) to manage capital in a manner which maximizes the interests of our shareholders.

The Corporation manages the capital structure and adjusts it considering changes in economic conditions and the risk characteristics of the underlying assets. The Corporation's capital structure is managed in conjunction with the capital structure and financial needs of the day-to-day operations. The Corporation currently funds the working capital requirements out of its internally generated cash flows and the use of credit facilities. To maintain or adjust the capital structure, the Corporation will work to secure new debt from its shareholders and expand the shareholder base with new participation that would make additional funds available. As at April 2022 the Corporation is not subject to any externally imposed capital requirements.

27. Commitments

(i) Lease obligation

The Corporation leases its premises and is currently bound by an eight-year lease which was renewed in June 2021 and will expire in August 2029. The Corporation has an option to extend the lease up to August 2034. The Corporation is expecting to exercise its option.

The yearly contractual undiscounted lease obligation payments are as follows:

	\$
2022	83
2023 to 2029	166
2030	187
2031	220
2032	230
2033	240
2034	207
Total	2,329

(ii) Licensing agreements

Milestones:

Under certain agreements, the Corporation may have to pay additional consideration should it achieve certain sales volumes or if certain milestones are met, such as approval for provincial reimbursement.

Royalty and profit sharing:

Under certain agreements, the Corporation is required to pay royalties, included in cost of goods sold or general and administrative expenses, based on Net Sales specific to each agreement at rates of up to 10% in any given year based on aggregate Net Sales levels achieved during the year as defined in each respective agreement. Furthermore, certain agreements require the Corporation to make profit sharing payments ranging from 25% to 50% of net profits as defined in the respective agreement.

28. Subsequent events

On May 2nd, 2022, the remaining non-convertible debentures issued July 2020 representing \$338 were reimbursed.