DRIHEALTHCARE

ADVANCING SCIENCE

in the Fast-Growing Pharmaceutical and Biotechnology Sector

March 2023

Disclaimer

This presentation has been prepared by DRI Healthcare Trust (the "Trust"). The Trust is an unincorporated open-ended fund trust governed by the laws of the Province of Ontario, Canada and is externally managed by DRI Capital Inc. ("DRI Capital" or the "Manager" and together with the Trust, "DRI Healthcare"). The Trust completed an initial public offering (the "IPO") on February 11, 2021 in which it acquired an initial portfolio of royalty assets from DRI Capital. The predecessor of the Manager was founded in 1989. Any references to employees or historical figures prior to the IPO refer to those of the Manager.

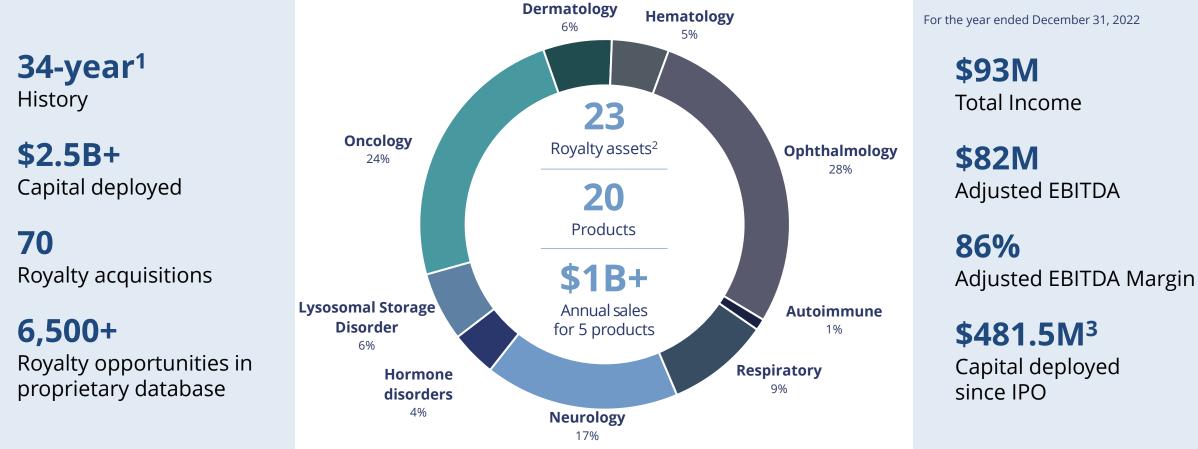
Certain statements made in this presentation, including responses to questions, may contain forward-looking statements within the meaning of the safe harbor provisions of Canadian provincial securities laws. Forward-looking statements involve risks and uncertainties, and undue reliance should not be placed on such statements. Certain material factors or assumptions are applied in making forward-looking statements and actual results may differ materially from those expressed or implied in such statements.

For additional information about factors that may cause actual results to differ materially from expectations, and about material factors or assumptions applied in making forward-looking statements, please consult the MD&A, the Risk Factors section of the Annual Information Form and DRI Healthcare Trust's other filings with Canadian securities regulators. DRI Healthcare Trust does not undertake to update any forward-looking statements; such statements speak only as of the date made.

This presentation also makes reference to certain non-GAAP financial measures including Total Cash Receipts and certain non-GAAP ratios including Adjusted EBITDA Margin and Adjusted Cash Earnings per Unit. These measures and ratios are not standardized measures under IFRS and are therefore unlikely to be comparable to similar financial measures disclosed by other issuers. Rather, these measures and ratios are provided as additional information to complement those IFRS measures by providing further understanding of DRI Healthcare Trust's financial performance from management's perspective. Accordingly, these measures should not be considered in isolation nor as a substitute for analysis of financial information reported under IFRS.

All dollar figures in this presentation are stated in US dollars.

Low risk exposure to rapid biopharma growth



Therapeutic area allocation based on net book value pro forma as at December 31, 2022, excluding Tzield royalty acquired in Q1 2023

Diverse portfolio with large pharmaceutical company characteristics

Historical data includes activities prior to establishment of DRI Healthcare Trust in February 2021

2. Excludes secured loan to CTI BioPharma

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3. Excludes \$176 million in potential additional deployment in milestones and options

Our competitive advantages

1 Seasoned team

of specialized investment professionals with life science backgrounds and advanced business and scientific degrees

2 Disciplined capital allocation

based on robust investment criteria that has resulted in 18% net IRRs over 16 years¹

3 Proactive sourcing

proprietary database tracking royalties on more than 2,000 drugs worldwide combined with deep industry relationships developed over our 30-year history

3 Exceptional execution

fundamental ground-up diligence on opportunities to execute high quality transactions

1. Based on results of funds managed by DRI Healthcare Trust's manage

Track record of delivering growth and value

| Drug Royalty I | Drug Royalty II | Drug Royalty III | DHT |
|-------------------------|----------------------|----------------------------|--|
| 2006 – 2008 | 2009 - 2013 | 2013 - 2018 | 2021 - present |
| 19 New Royalties | 27 New Royalties | 15 New Royalties | 9 New Royalties & 1 Loan |
| valued at | valued at | valued at | valued at up to |
| \$645M | \$730M ¹ | \$586M | \$658M ² |
| IRR | IRR | IRR | Current Yield ³ >5% |
| 19% | 18% | 20% | |
| Remicade Kolair | Simponi [®] | EYLEA SPINRAZA KEYTRUDA | VONJO [®] Tzield [®] OMIDRIA [®] |

Consistent track record of efficient capital deployment at high returns

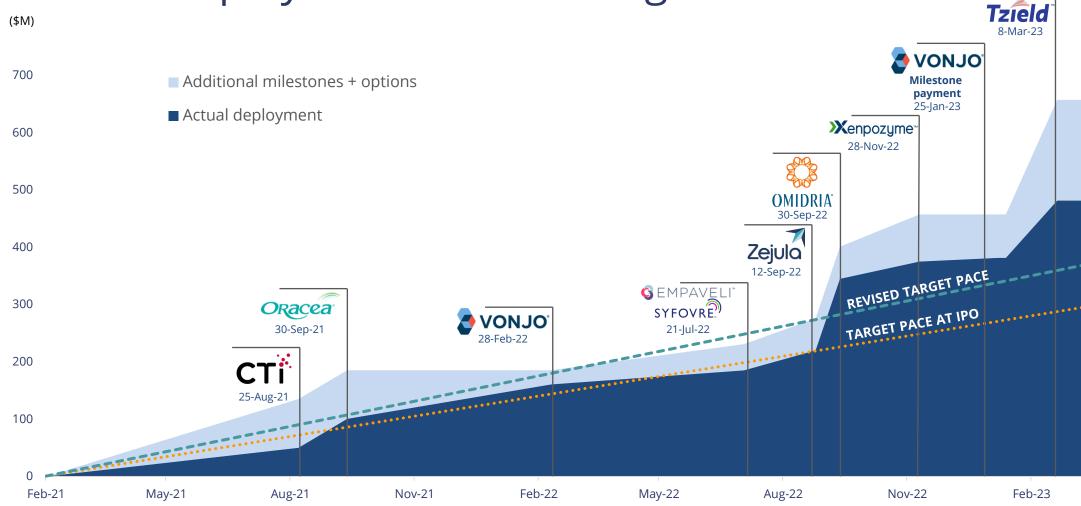
- 1. Includes \$82 million in capital deployed via co-investments through RMF 2 Co-Investment Fund
- 2. Includes up to \$18.5 million in milestone payments for Vonjo, a \$21 million option to increase our exposure to Empaveli, a \$10 million milestone payment for Zejula, \$26.5 million in milestone payments for Xenpozyme, and a \$100 million milestone payment for Tzield
- 3. Does not include annual special dividend

Delivering on our long-term strategy

| | At IPO | Today | 2025 target |
|--------------------------------|---|--|--|
| Capital deployment | \$650 – 750 million over 5 years | \$481.5 million deployed to date + \$176 million in milestones and options | Revised deployment target of \$850 – 900 million |
| Sustainable cash generation | Declining cash curve due to expected asset expiries | Flat to slightly growing cash receipts through 2030 without any new deals | 7% - 9% annual royalty receipt growth |
| Portfolio duration | 8 years | >10 years | >10 years |
| Capital resources | IPO proceeds and debt capacity | Attractive credit facilities with compounding effect of cash flows | Expanded credit facilities with compounding effect of cash flows |

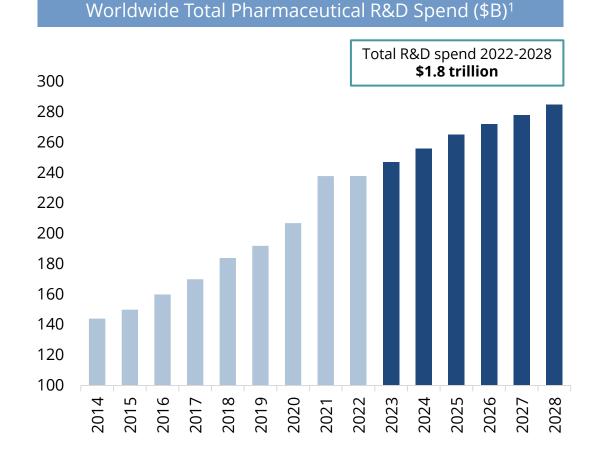
Focus on building long-term and sustainable strategic growth

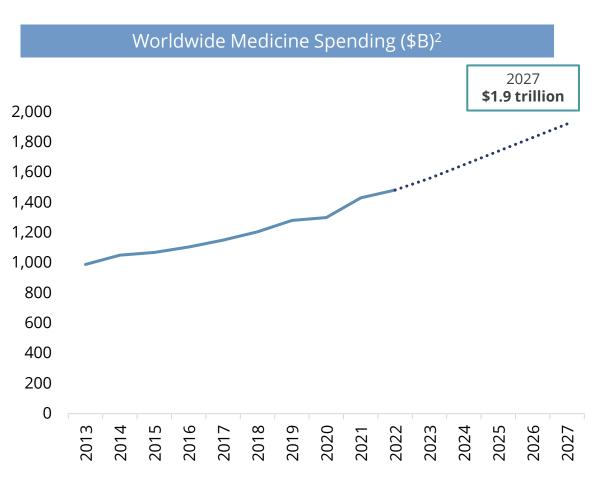
Current deployment exceeds targets



Deployment pace and need for capital by counterparties → 5 year deployment target increased to \$850 – 900 million

Industry R&D spending is propelling innovation



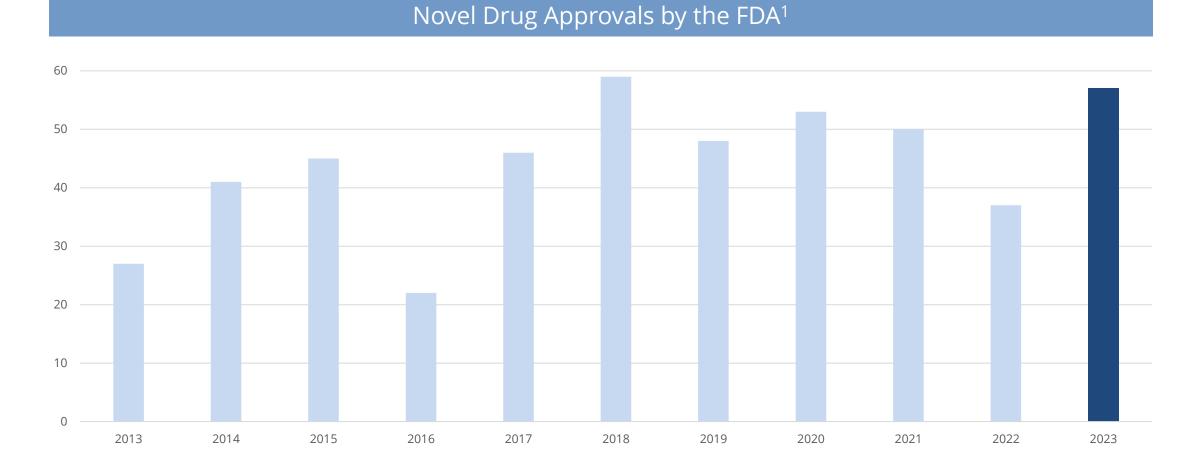


Growing R&D increases our future opportunity set, while current cash needs bolster our pipeline



Source: Evaluate Pharma World Preview 2022, Outlook to 2028 15th edition, October 2022 Source: IQVIA Global Use of Medicines 2023, Outlook to 2027, January 2023

R&D spending has resulted in many innovative products



FDA approval of novel drugs opens the door to abundant royalty acquisition opportunities

1. Source: Historic: FDA.gov, 2023 Forecast: Evaluate Vantage 2023 Preview, December 2022

Royalty transaction activity is expanding with strong industry tailwinds

Trailing three-year average number of transactions¹ Trailing three-year average value of transactions (\$B)¹ Aggregate royalty market -----Synthetic royalties Aggregate royalty market -----Synthetic royalties

Royalty financing's unique appeal has propelled it to mainstream adoption

Source: Internal database. Includes royalty related monetization transactions across the ecosystem (inventor, academic/non-profit, biotech and pharma)



State of the biotech market

Biotech IPOs¹ Biotech Equities Performance² 16 120 140 130 14 100 120 7.3% 12 110 80 Total Proceeds (\$B) 100 Number of IPOs 10 90 8.4% 8 80 11.3% 6 70 60 4 50 20 2 40 Jan-21 Jan-21 Mar-21 May-21 Jun-21 Jul-21 Jul-22 Sep-21 Jun-22 Jun-22 Jun-22 Jun-22 Jun-22 Sep-22 Sep-22 Sep-22 Sep-22 Jun-22 Ju Jan-23 Feb-23 0 1998 2000 2002 2004 2006 2008 2010 2012 2014 2016 2018 2020 2022 BTK Price Index S&P Biotech Select Industry Index SPX Index Total Proceeds — Number of IPOs

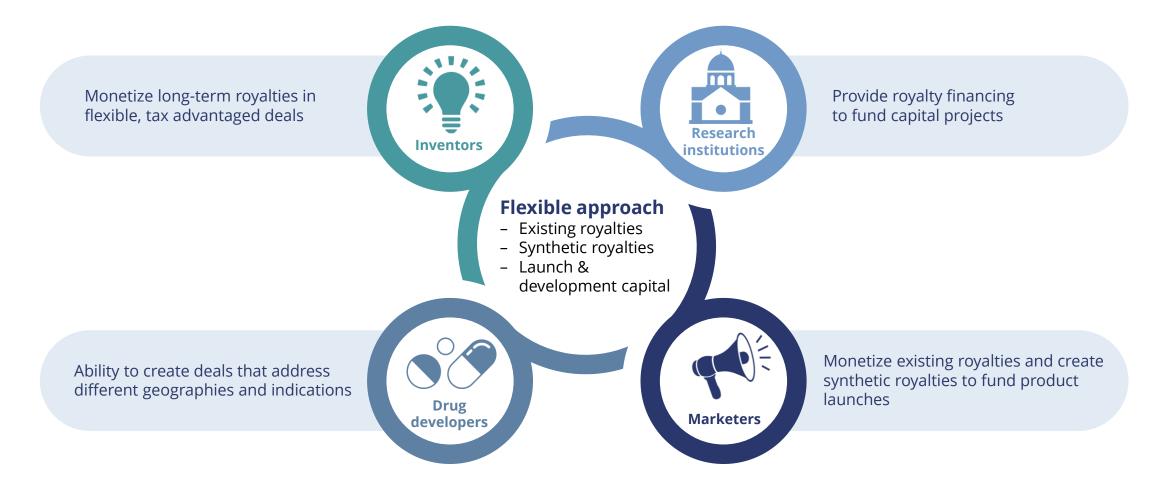
Rapid expansion of biotech market with >500 IPOs in last 10 years

Struggling equity capital markets for biotechs

Creates an environment where a large amount of capital is required, making DRI's royalty financing very attractive



Creating win-win deals for multiple counterparties



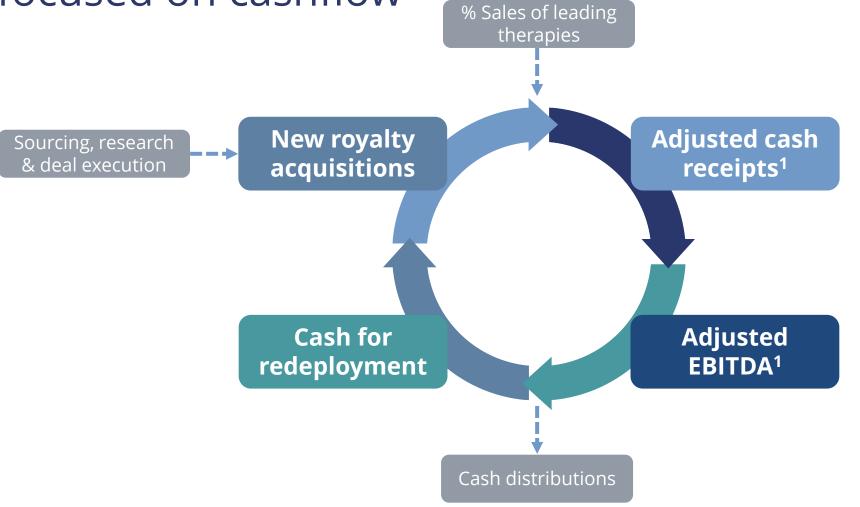
A proven and repeatable asset identification, selection and execution process

Proven track record of closing accretive transactions

| | Investment Thesis | Transaction Size | |
|--|---|--|--|
| CTi avonjo | High-quality oncology product with strong growth potential | \$116.5 million + \$18.5 million potential milestone | |
| Oracea | Dermatology product with existing commercial track record | \$50.5 million | |
| SEMPAVELI SYFOVRE | Hematology and ophthalmology product with long-term horizon and attractive growth prospects | \$24.5 million + \$21.0 million option | |
| Zejula | High-quality oncology product with multiple pipeline indications | \$35.0 million + \$10.0 million potential milestone | |
| OMIDRIA | Structured transaction on established product providing cash accretion | \$125 million | |
| Xenpozyme™ | Only approved product for ASMD with strong IP and long duration \$26.5 million in potential m | | |
| Tzield Newly approved Diabetes product with long-term cash flows and growth potential | | \$100.0 million + \$100 million in potential milestones | |

Completed seven transactions since IPO totaling up to \$657.5 million, with \$481.5 million deployed to date

Sustainable and efficient business model focused on cashflow

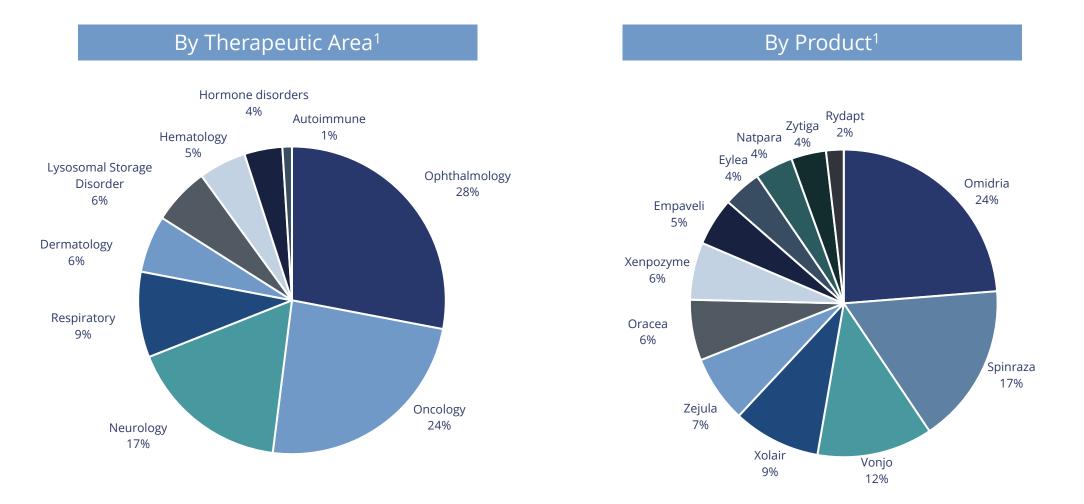


Diversified portfolio generates significant cash to deploy to new royalties and investors

1. Adjusted EBITDA and Adjusted Cash Receipts are non-GAAP measures



Robust diversified portfolio

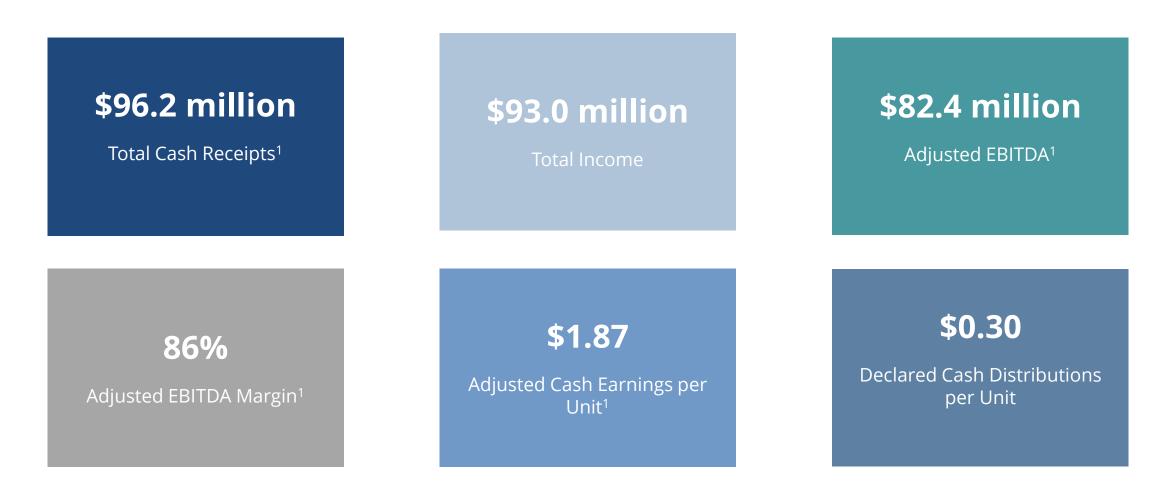


No individual product accounts for more than 24% of net book value

1. Based on net book value pro forma as at December 31, 2022 excluding Tzield royalty acquired in Q1 2023



2022 financial highlights

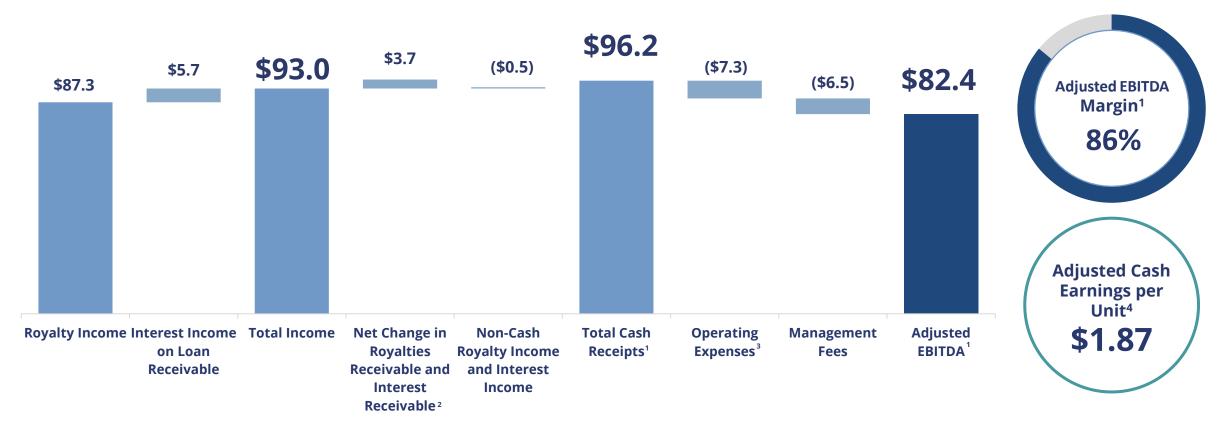


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Total Cash Receipts and Adjusted EBITDA are non-GAAP measures. Adjusted EBITDA Margin and Adjusted Cash Earnings per Unit is a non-GAAP ratio. See the following disclosure in "Financial Review: Non-GAAP Financial Measures" in the MD&A of DRI Healthcare Trust for the year ended December 31, 2022 (the "MD&A"), consisting of the first two paragraphs under the subheading "Total Cash Receipts and Total Cash Royalty Receipts" on page 15 of the MD&A, the reconciliation table for Total Cash Receipts and Total Cash Royalty Receipts on page 17 of the MD&A and the disclosure on pages 18 and 19 of the MD&A under the subheadings "Adjusted EBITDA", "Adjusted EBITDA Margin" and "Adjusted Cash Earnings Per Unit", which disclosure is incorporated by reference in this presentation. The MD&A is available on SEDAR at www.sedar.com

Strong cash generation

Adjusted EBITDA for the Year Ended December 31, 2022 (\$M)¹



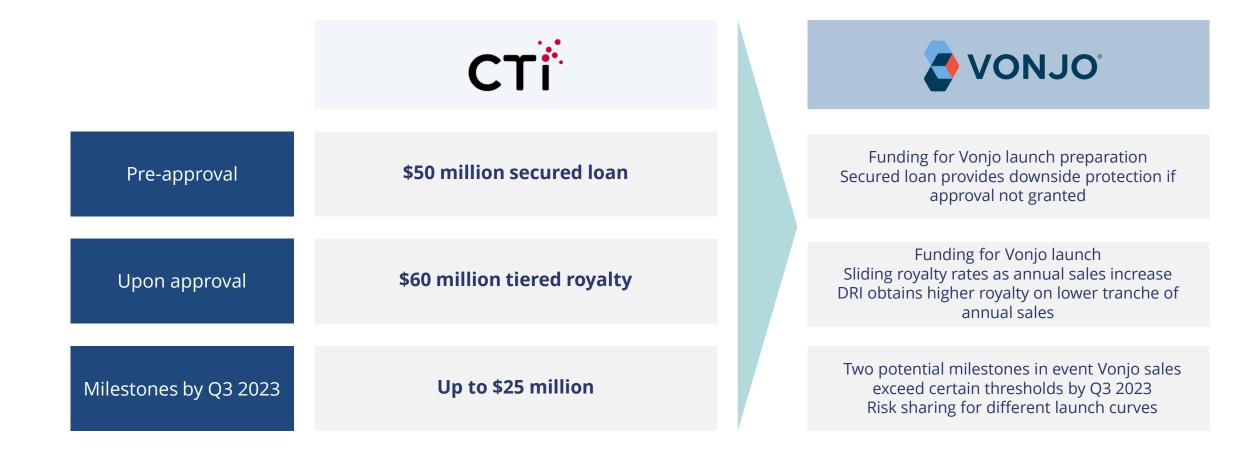
Cash available to drive portfolio growth and maintain distributions to unitholders

Adjusted EBITDA and Total Cash Receipts are non-GAAP financial measures. Adjusted EBITDA Margin is a non-GAAP ratio calculated as Adjusted EBITDA / Total Cash Receipts. See note 1 on page 4 of this presentation. The Net Change in Royalties Receivable and Interest Receivable represents royalties and interest receivable at the beginning of period, less royalties and interest receivable at the end of period, plus acquired royalties receivable and acquired cash royalty receipts included in the purchase price of the assets.

DRIHEALTHCARE 3. Operating expenses are net of \$0.3 million related to board of trustee unit-based compensation and \$0.3 million related to amortization of other current assets Adjusted Cash Farnings per Unit is a non-GAP ratio and is the sum of Adjusted Cash Farnings per Unit in each of the last four quarters calculated as per earning

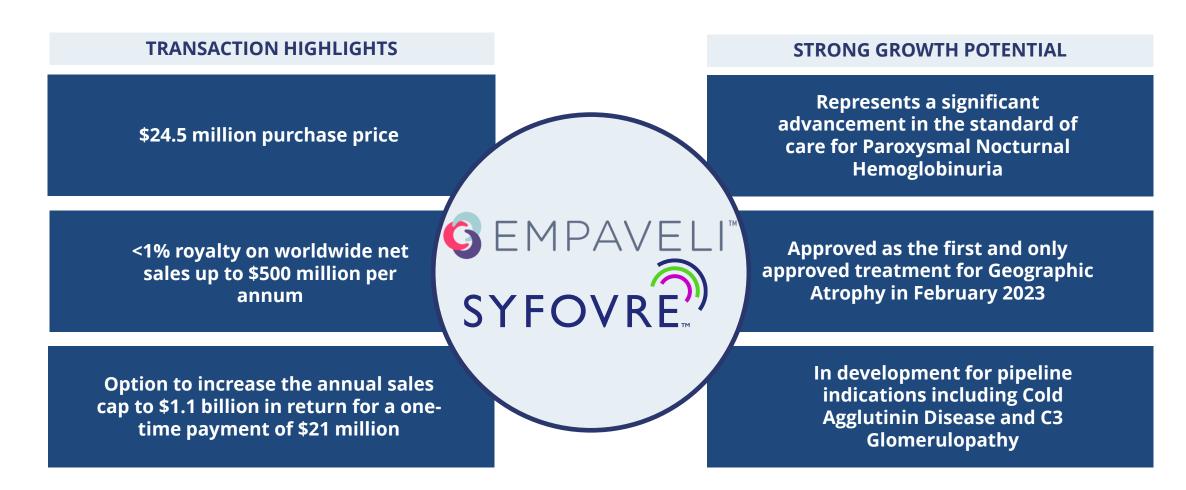
Adjusted Cash Earnings per Unit is a non-GAAP ratio, and is the sum of Adjusted Cash Earnings per Unit in each of the last four quarters, calculated as net earnings and other comprehensive earnings, plus: (i) amortization of royalty assets, (ii) impairment of royalty assets, (iii) amortization of how last four quarters, calculated as net earnings and other comprehensive earnings, plus: (i) amortization of royalty assets, (iii) impairment of royalty assets, (iii) amortization of by last four quarters, calculated as net earnings and other comprehensive earnings, plus: (i) amortization of royalty assets, (iii) impairment of royalty assets, (iii) amortization of by last four quarters, calculated as net earnings and other comprehensive earnings, plus: (i) amortization of royalty assets, (iii) non-cash royalty income, and (v) board of trustees unit-based compensation, and less: (i) net gain (loss) on foreign exchange derivatives, (iii) non-cash royalty income, and (v) non-cash interest income on loan receivable, divided by fully-diluted weighted average units outstanding. See note 1 on page 4 of this presentation.

Deal structure case study: CTI Biopharma / Vonjo



Proven ability to provide flexibility in deal structuring while managing risk

Empaveli royalty transaction



Long-term horizon and attractive growth prospects

Zejula royalty transaction

TRANSACTION HIGHLIGHTS

\$35 Million up front purchase price

0.5% net royalty on worldwide net sales by GSK

Royalties collected on 1-quarter lag

STRONG GROWTH POTENTIAL

\$10 million milestone payment if Zejula is approved by FDA for the treatment of endometrial cancer by December 21, 2025

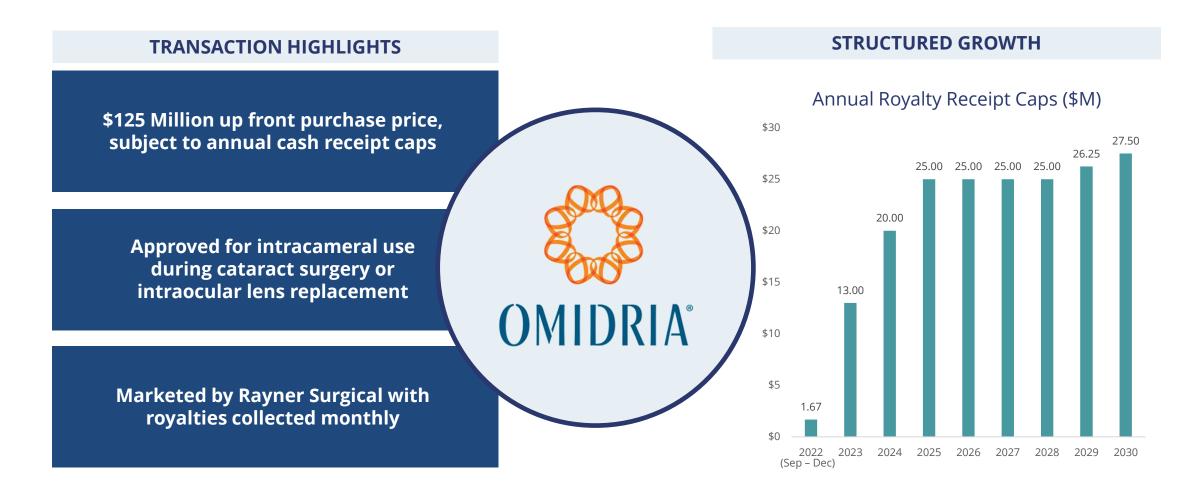
In development for metastatic castrate sensitive and resistant prostate cancer, endometrial cancer, HER2-breast cancer, and non-small cell lung cancer

Royalty term expected to continue for at least another 10 years world wide

Multiple indications in development represent a pipeline in a product

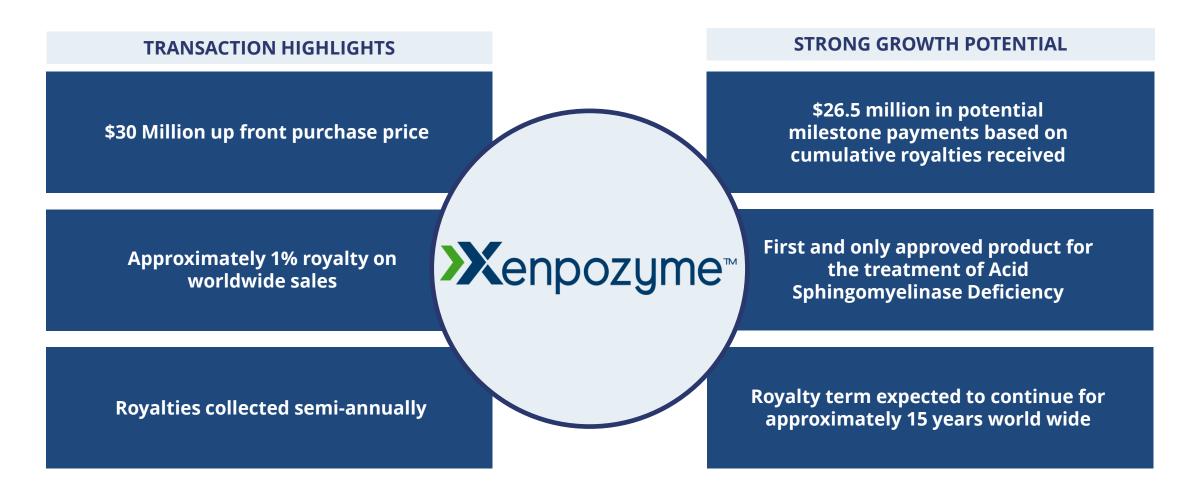
Zejula

Omidria royalty transaction



Substantial near-term cash flows with long-term structural growth

Xenpozyme royalty transaction



Long duration product with strong IP protection will generate high multiple on invested capital

Tzield royalty transaction

TRANSACTION HIGHLIGHTS

\$100 Million up front purchase price for a single digit royalty on Provention Bio's worldwide net sales

> Royalties based on sales from January 1, 2023 are collected on 1-quarter lag with first receipt in Q2 2023

Protected by patent and regulatory exclusivities for 12 year from first commercial sale

STRONG GROWTH POTENTIAL

Up to a \$50 million potential milestone tied to the successful advancement of treatment of newly diagnosed or recentonset Type 1 diabetes by 2028

> Additional \$50 million potential milestone payment based on exceeding certain sales thresholds

Transaction enhances our long-term cash flow profile and significantly increases the duration of our portfolio to >10 years

Newly approved product with long-term cash flows

Tzield

Portfolio performance

| (\$ thousands) Asset | Primany Markotor(c) | Thorapoutic Aroa | Total Cash Royalty Receipts ^{1,2} FY 2022 | Net Book Value 12/31/22 |
|--------------------------------|--|--|---|----------------------------|
| SEMPAVELI' SYFOVRE | Primary Marketer(s) Apellis 🔁 SOOI | Therapeutic Area Hematology / Ophthalmology | 269 | 23,437 |
| EYLEA | | Ophthalmology | 11,280 | 23,041 |
| FluMist.Quadrivalent | AstraZeneca | Vaccine | 2,952 | 2,096 |
| % Natpara | Takeda | Endocrinology | 2,625 | 19,457 |
| OMIDRIA* | ARayner | Ophthalmology | 1,670 | 121,782 |
| Oracea | SALDERMA | Dermatology | 7,662 | 32,144 |
| RYDAPT | U NOVARTIS | Oncology | 10,102 | 8,899 |
| SPINRAZA | Biogen | Spinal Muscular Atrophy | 16,466 | 86,587 |
| Stelara' Simponi' IL (RIS | Johnson Johnson 📀 MERCIK ひ NOVARTIS 🏾 🎸 Mitsubishi Tanabe | Autoimmune | 3,704 | 3,871 |
| Tzield | proventionbio | Diabetes | - | 100,000 ³ |
| SCROV 🚭 | СТЇ | Oncology | 3,155 | 62,876 |
| ≫ enpozyme [∞] | sanofi | ASMD | - | 31,293 |
| Xolair | Roche UNOVARTIS | Respiratory | 9,646 | 47,389 |
| Zejula | gsk | Oncology | 692 | 33,973 |
| O Zytiga [,] | Johmon Johmon AstraZeneca | Oncology | 18,059 | 18,465 |
| Various | | | 2,563 | 2,883 |
| Total | | | 90,845 | 518,193 |

DHT's assets have continued to show strong performance

 Does not include Xenpozyme royalty acquired in Q4 2022 or Tzield royalty acquired in Q1 2023 for which the first cash royalty receipts are expected to be received in Q2 2023
Total Cash Royalty Receipts is a non-GAAP measure
Represents purchase price **DRI**HEALTHCARE

Growth opportunities from existing assets

| | · · · · · | | | | |
|------------|---|--------------------------------------|--|--------------------------|---------|
| | | Phase 1 | Phase 2 | Phase 3 | Phase 4 |
| | | | RESPOND: Spinraza in patients who had re | ceived Zolgensma | |
| Spinraza | | ASCEND: Higher dose Spi | inraza in patients who had received Evrysdi | | |
| | | DEVO | TE: Higher dose Spinraza | | |
| Tzield | | PROTECT: Trea | tment for stage 3 Type 1 Diabetes | | |
| Vonjo | PACIFICA: Confirmatory trial in Myelofibrosis | | | | |
| Xolair | | OUTMATCH: Xolair monoth | nerapy or as adjunct therapy in food allergies | | |
| | | FIRST: 1L treatment of st | age III/IV Ovarian Cancer with Dostarlimab | | |
| | ZEST: tBRCA | m HER2–negative BC or tBRCAwt TNB | C who have detectable ctDNA after completi | on of definitive therapy | |
| Zejula | RUB | Y: maintenance treatment of Recurrer | nt or Primary Advanced Endometrial Cancer v | vith dostarlimab | |
| | | ZEAL: 1L maintenance therap | y in combination with pembrolizumab in NS | | |
| | | AMPLITUDE: Com | bination of Zytiga + Zejula in mHSPC | | |
| Zytiga | | MAGNITUDE: Com | nbination of Zytiga + Zejula in mCRPC | | |
| | | AMPLITUDE: Com | bination of Zytiga + Zejula in mHSPC | | |
| Empaveli / | | PLAUDIT: Treatment for wAIHA o | r CAD | | |
| Syfovre | DISCOVERY: 1 | reatment for IgA Nephropathy, Lupus | Nephritis, PNM, or C3G | | |
| Rydapt | | Rydapt + decitabine in unfit AML p | patients | | |
| | Rydapt + HDM201 in r, | /r AML with FLT mutation | | | |

Additional indications have potential to enhance royalty streams

Committed to best practices in ESG





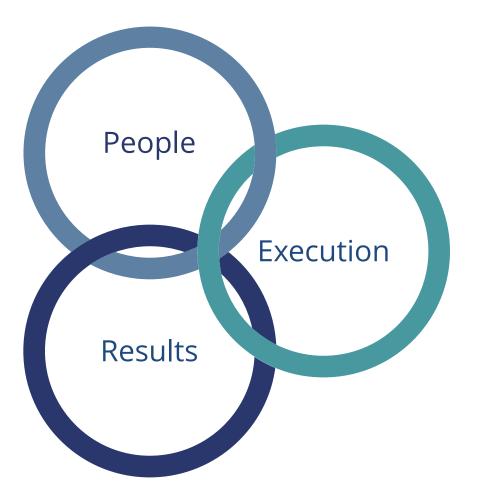
Governance

Valuing diversity and community support

- Best practice governance policies in place
- Diverse and majority independent Board
- Board oversight of ESG and risk management
- Active unitholder engagement
- Robust cybersecurity
- Whistleblower policy in place

Striving to deliver value to our stakeholders, our community, and society as a whole

Our key priorities



Invest in our people and build the industry leading royalty investment team

Execute on strong pipeline and operate at peak performance in all aspects of our business

Focus on long-term, sustainable growth generating strong unitholder returns

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Contact Us

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