



Adaptive
biotechnologies™

Third Quarter 2022
Earnings Conference Call

Safe Harbor

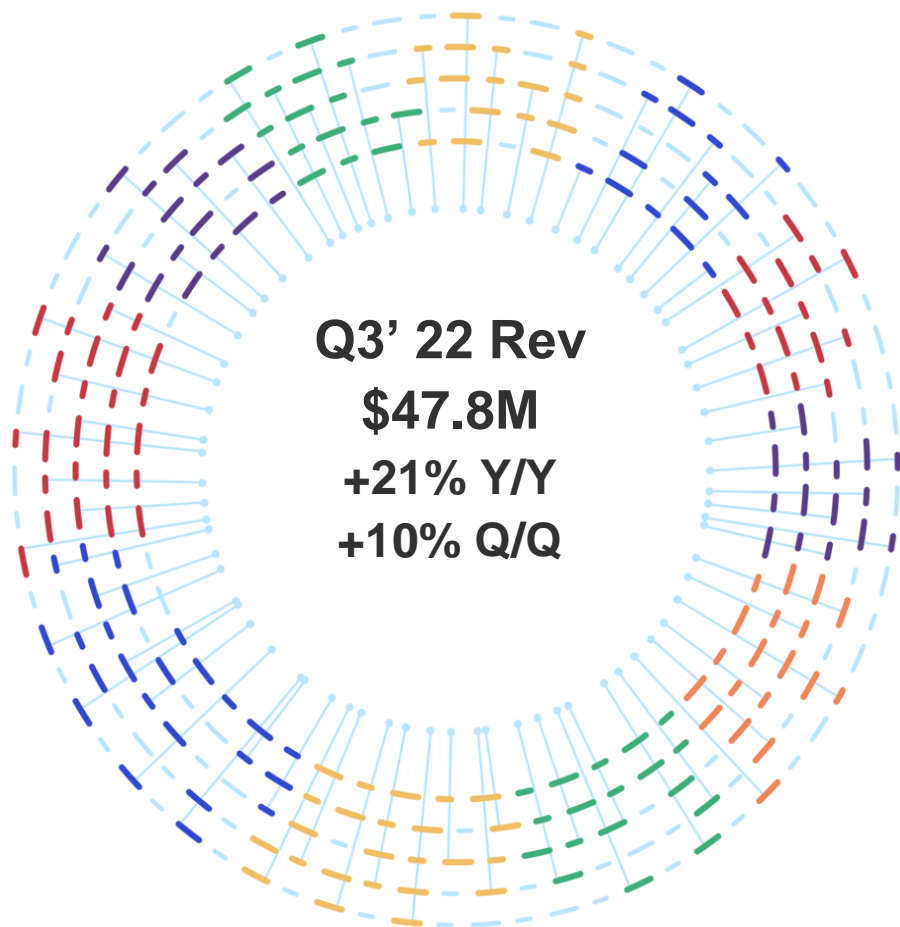
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This presentation contains “forward-looking statements” within the meaning of Section 27A of the Securities Act of 1933, as amended, and Section 21E of the Securities Exchange Act of 1934, as amended. These statements are intended to be covered by the “safe harbor” created by those sections. Forward-looking statements are neither historical facts nor assurances of future performance. Instead, they are based on our current beliefs, expectations and assumptions regarding the future of our business, future plans and strategies, our development plans, our preclinical and clinical results and other future conditions. In some cases, you can identify forward-looking statements by the following words: “may,” “will,” “could,” “would,” “should,” “expect,” “intend,” “plan,” “anticipate,” “believe,” “estimate,” “predict,” “project,” “potential,” “continue,” “ongoing” or the negative of these terms or other comparable terminology, although not all forward-looking statements contain these words. All statements, other than statements of historical facts, contained in this presentation are forward looking statements, including statements regarding the ability to map adaptive immune responses to target disease states, the ability to leverage any such findings to advance solutions to diagnose, treat and prevent infectious diseases; regarding our future financial or business performance, conditions, plans, prospects, trends or strategies and other financial and business matters; our current and prospective products and product candidates; FDA clearance or authorization of any products; planned non-IDE clinical studies, clinical trials and preclinical activities, research and development costs, current and prospective collaborations; the estimated size of the market for our products and product candidates; the timing and success of our development and commercialization of current products and product candidates; the availability of alternative therapies for our target markets; and the other risks and uncertainties described in our filings with the Securities and Exchange Commission including the Risk Factors and Management’s Discussion and Analysis of Financial Condition and Results of Operations sections of our most recently filed Quarterly Reports on Form 10-Q and our Annual Report on Form 10-K, including our most recent Annual Report on Form 10-K filed on February 15, 2022. Although we believe the expectations reflected in such forward-looking statements are reasonable, we can give no assurance that such expectations will prove to be correct. Risks and uncertainties could cause actual results to differ materially from those expressed in our forward-looking statements. Accordingly, readers are cautioned not to place undue reliance on these forward-looking statements. Except as required by applicable law, we do not plan to publicly update or revise any forward-looking statements contained herein.

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In addition, non-GAAP financial measures are included in this presentation. Please see table in appendix for reconciliation to the most directly comparable GAAP measure.

Q3 2022 Key Highlights



MRD Business

- Drove clonoSEQ test volume growth of 52% vs prior year
- Signed Epic integration agreement
- Signed agreement for a new primary-end point study in MM with existing partner

Immune Medicine Business

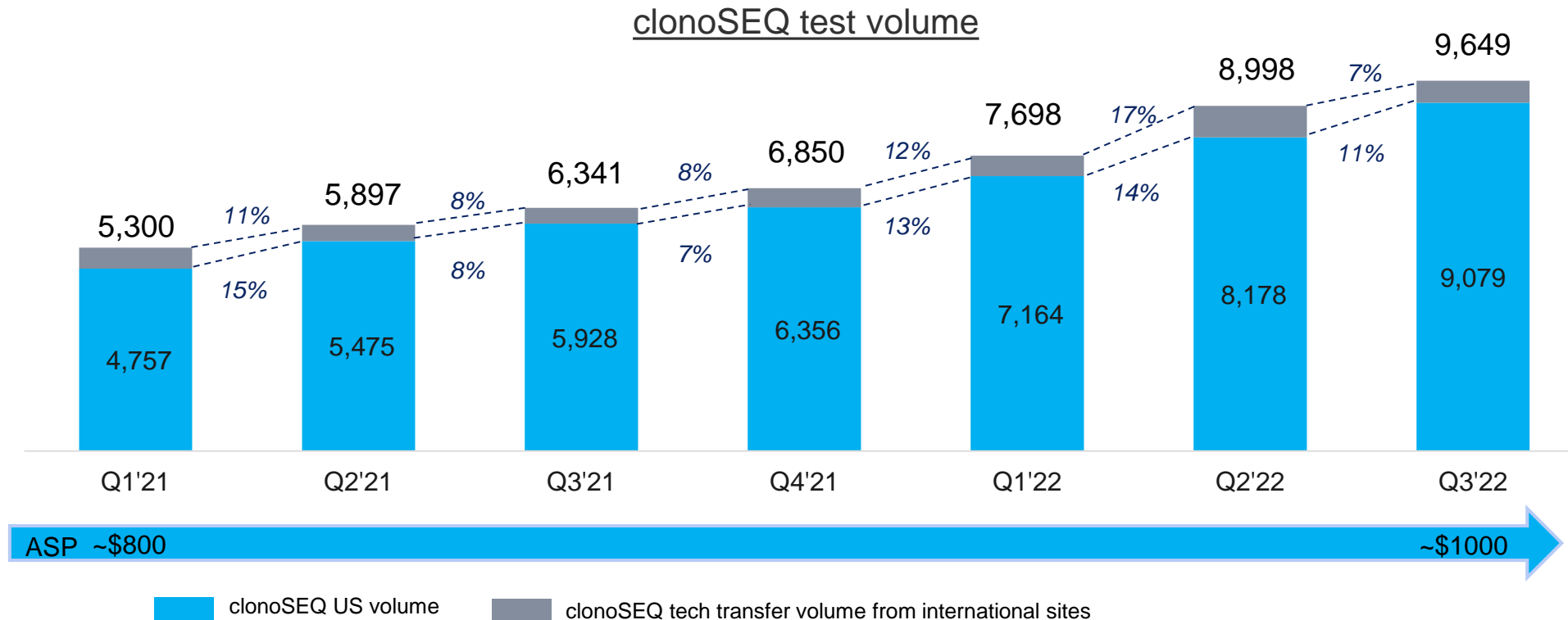
- IM Pharma revenue grew 23% vs prior year
- Increased services penetration in Ph1 and Ph2 clinical trials in multiple indications
- Genentech partnership on track with both shared and private products

Corporate

- Completed \$250M non-dilutive royalty financing with OrbiMed
- Continued to drive operating leverage
- Set path to profitability (positive adj. EBITDA 2025; cash flow break even 2026)

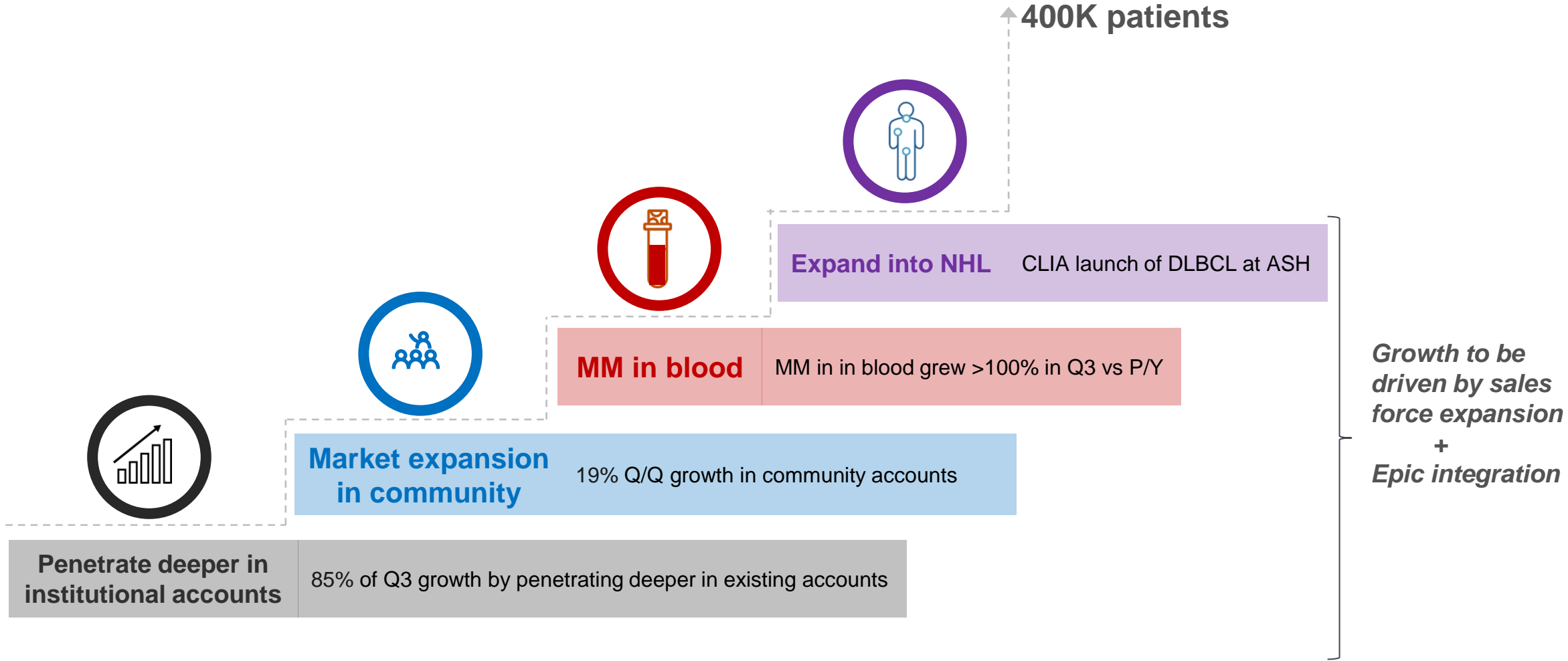
MRD business: clonoSEQ clinical testing

- Q3'22 test delivered volume +52% vs P/Y; +7% vs P/Q
 - 403 ordering accounts in Q3 (+53% vs P/Y)
 - 1,612 ordering HCPs in Q3 (+56% vs P/Y)
 - Unique patients tested increased (+62% vs P/Y)



ASP now expected to grow in the mid single digits annually

Advancing on our strategy to maintain leadership in lymphoid cancers

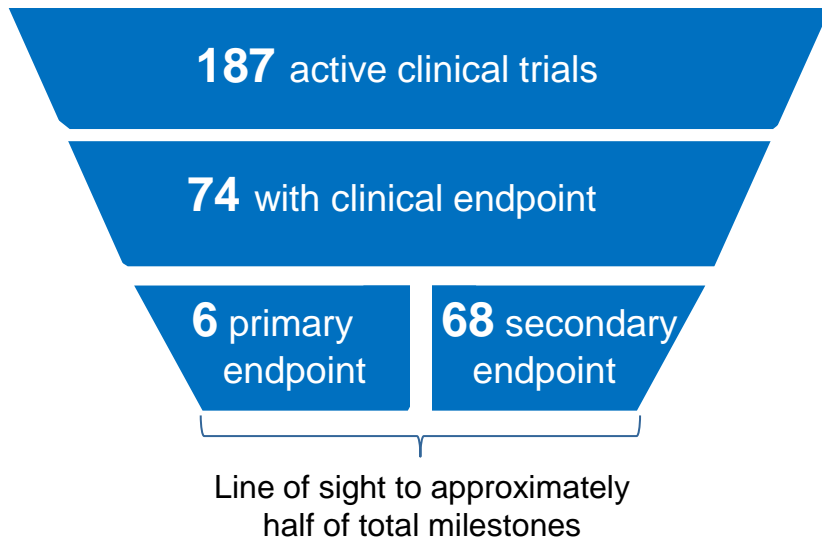


MRD pharma business: portfolio continues to increase

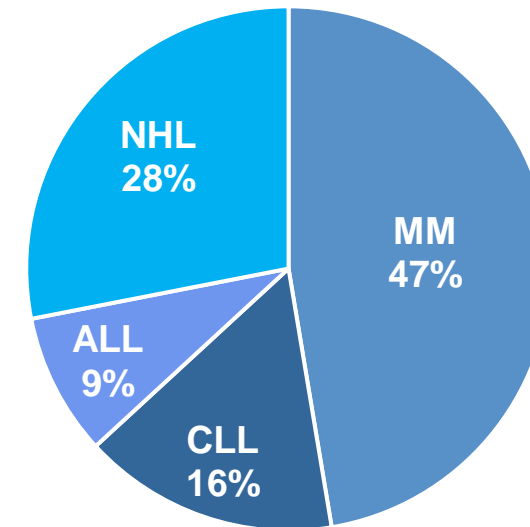
clonoSEQ MRD, gold standard in drug trials, growing use as an endpoint

Portfolio Overview

- >60 BioPharma partners
- Sequencing revenue plus regulatory milestones
- >\$370M in milestones from future & active trials



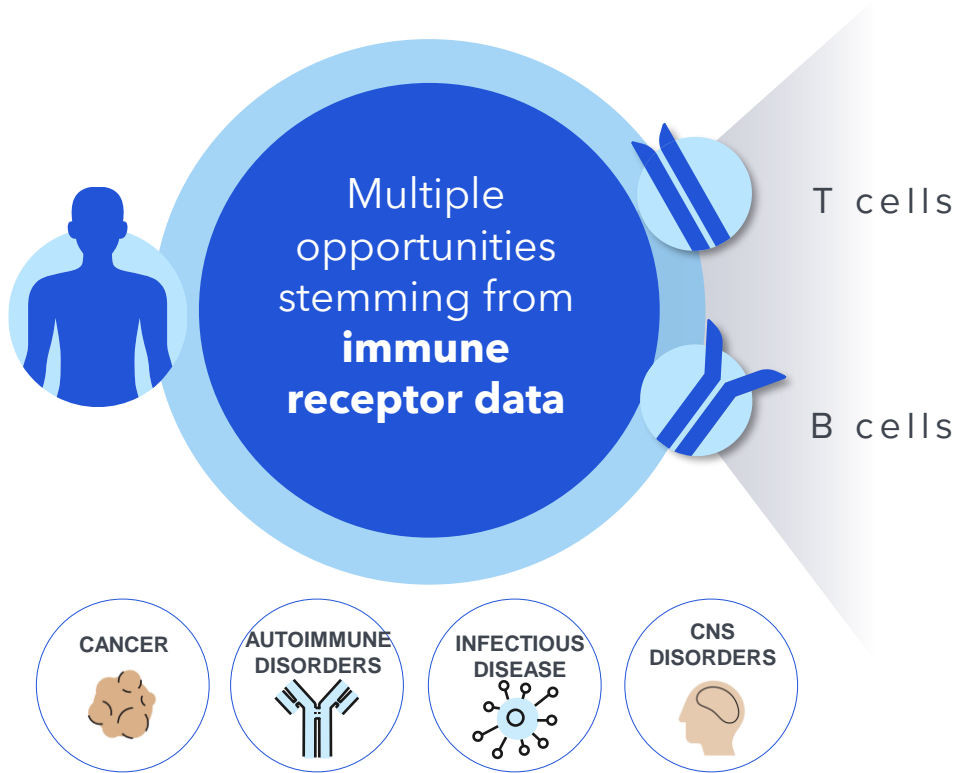
Portfolio Mix by Indication



50% of trials in phase 2 and phase 3

Immune Medicine Business

Immune Medicine Platform



Growth Areas

Multiple shots on goal to create value, grow and monetize our immune receptor data across clinical applications

Pharma Services

Immune receptor sequencing

Drug Discovery

Therapeutic TCRs, antibodies and targets

Pharma Services growing portfolio across multiple indications

4+

Major therapeutic areas

500+

Total studies to date

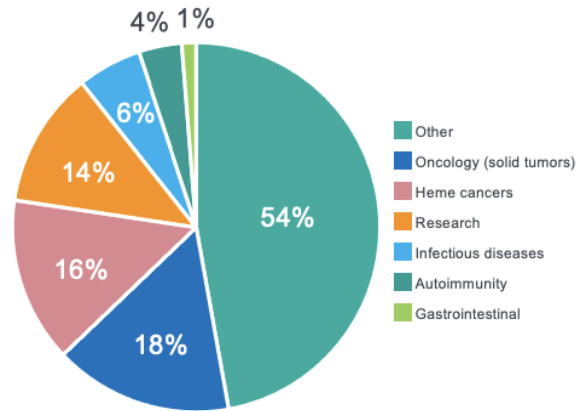
140+

Total active studies

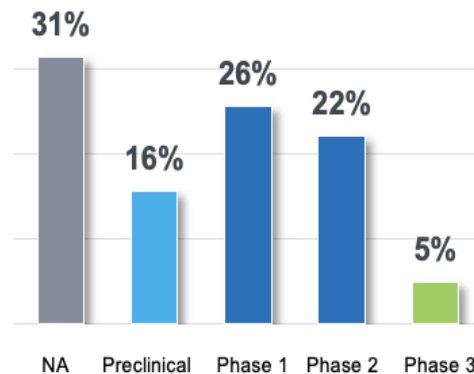
85+

Companies

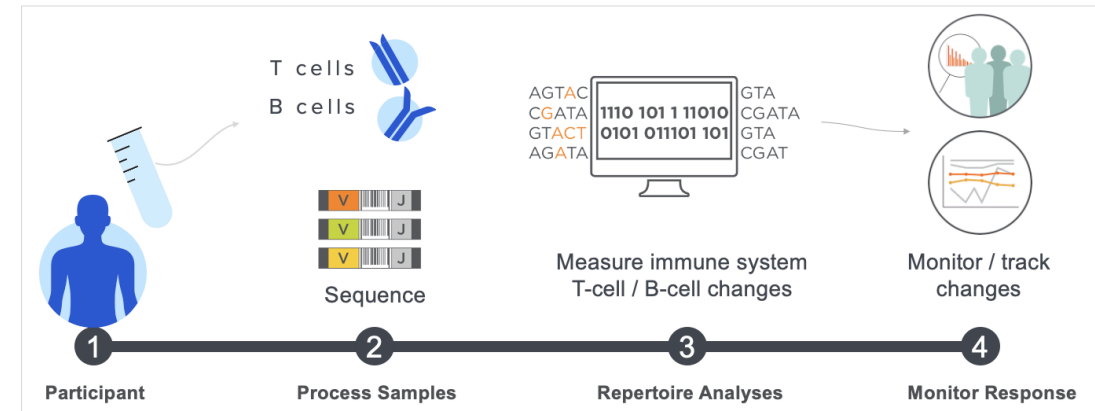
Portfolio mix by indication



Portfolio mix by study phase



Rich immune receptor biomarker data accelerates clinical trials



Growth drivers

23%
Q3 Y/Y revenue growth

- Scale companies / # of studies using sequencing
- Increase penetration in later stage trials and across indications

Drug Discovery unlocks the value of immune receptors as therapeutics

Partnered pipeline

Genentech
A Member of the Roche Group

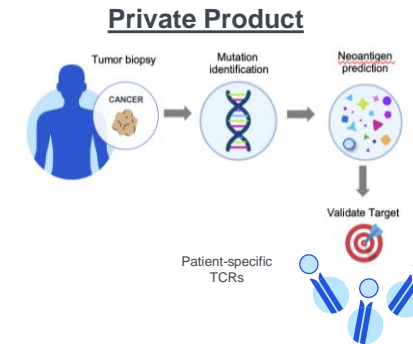
TCRs targeting shared cancer (neo)antigens

TCR candidate selected to progress towards IND
Deliver 2 add'l TCR data packages by YE

Build private product process

Establish private product specs and build data package
Begin steps toward early product dev

Shared Product



36%
Q3 Y/Y growth in GNE amortization

Adaptive pipeline

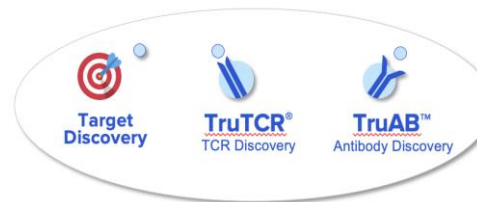
TCR Therapeutics

Establish POC data package(s)
Focus in areas of unmet clinical need

Antibody Therapeutics

Seek partner(s) for Ab discovery
Focus on key differentiators

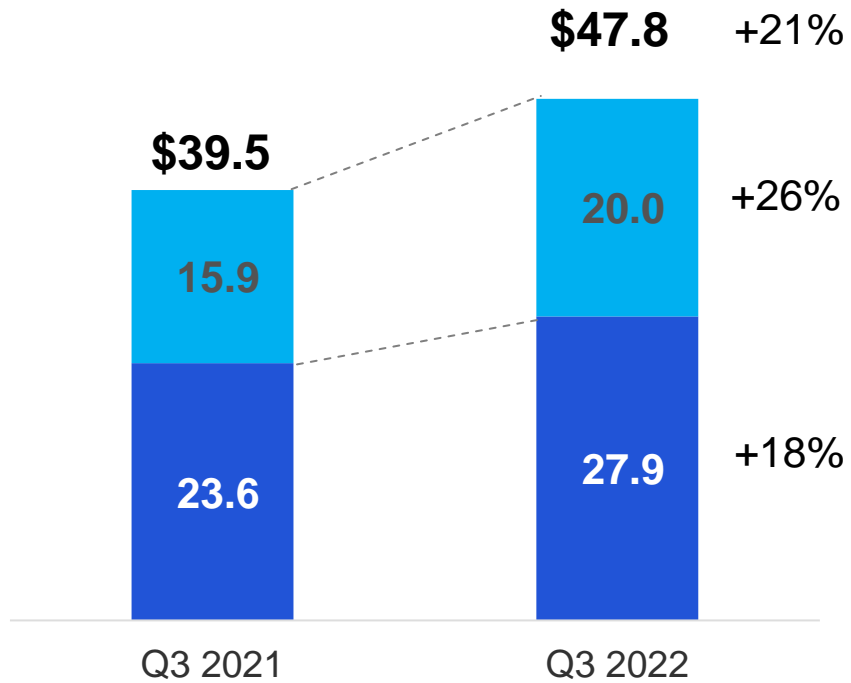
Growth drivers



Leverage core competencies (TruTCR, TruAB) to advance therapeutics directed against attractive targets

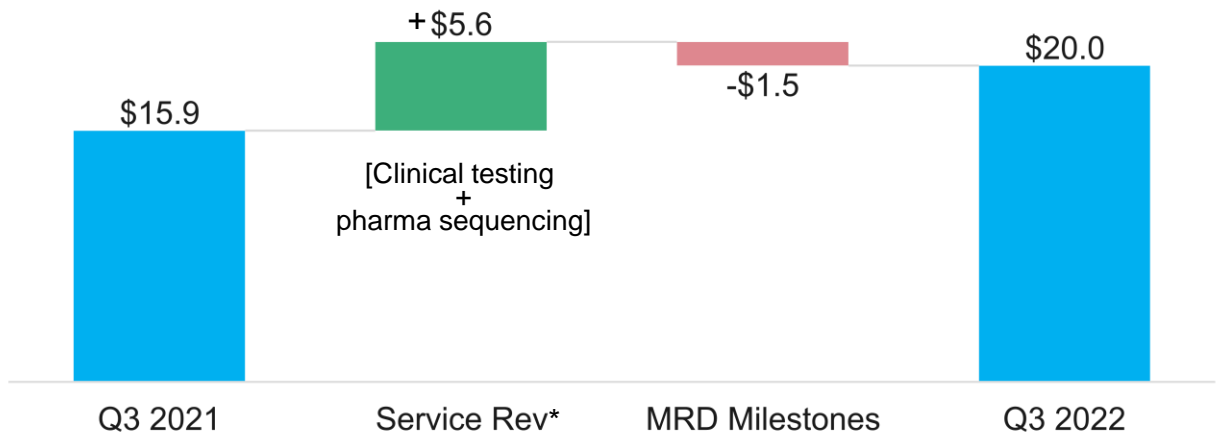
Q3 2022 key financial highlights

Total Revenue (\$M)

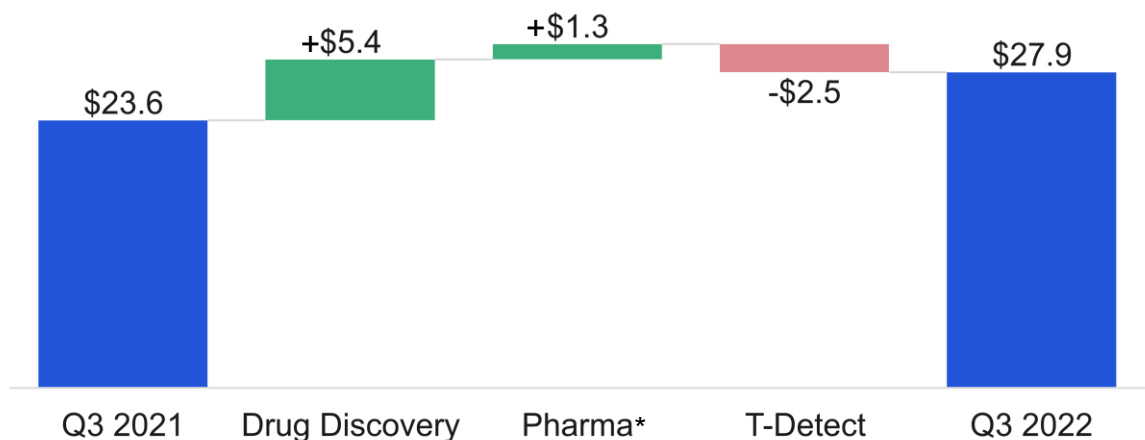


■ MRD Business
■ Immune Medicine Business

MRD Revenue (\$M)

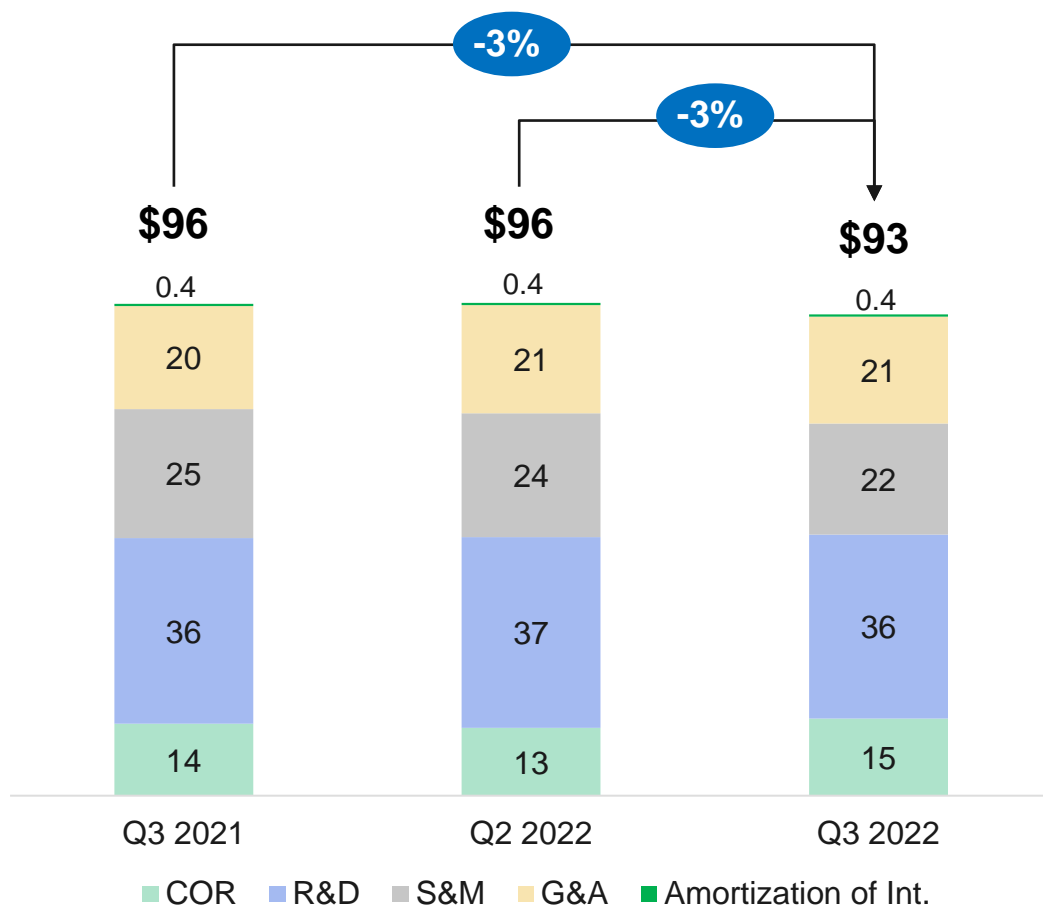


Immune Medicine Revenue (\$M)



Q3 2022 key financial highlights cont.

Operating Expenses (\$M)



Strong Balance Sheet

- ~\$528M in cash, cash equivalents and marketable securities as of 09/30/2022
- Q4 2022 expected cash burn <\$50M

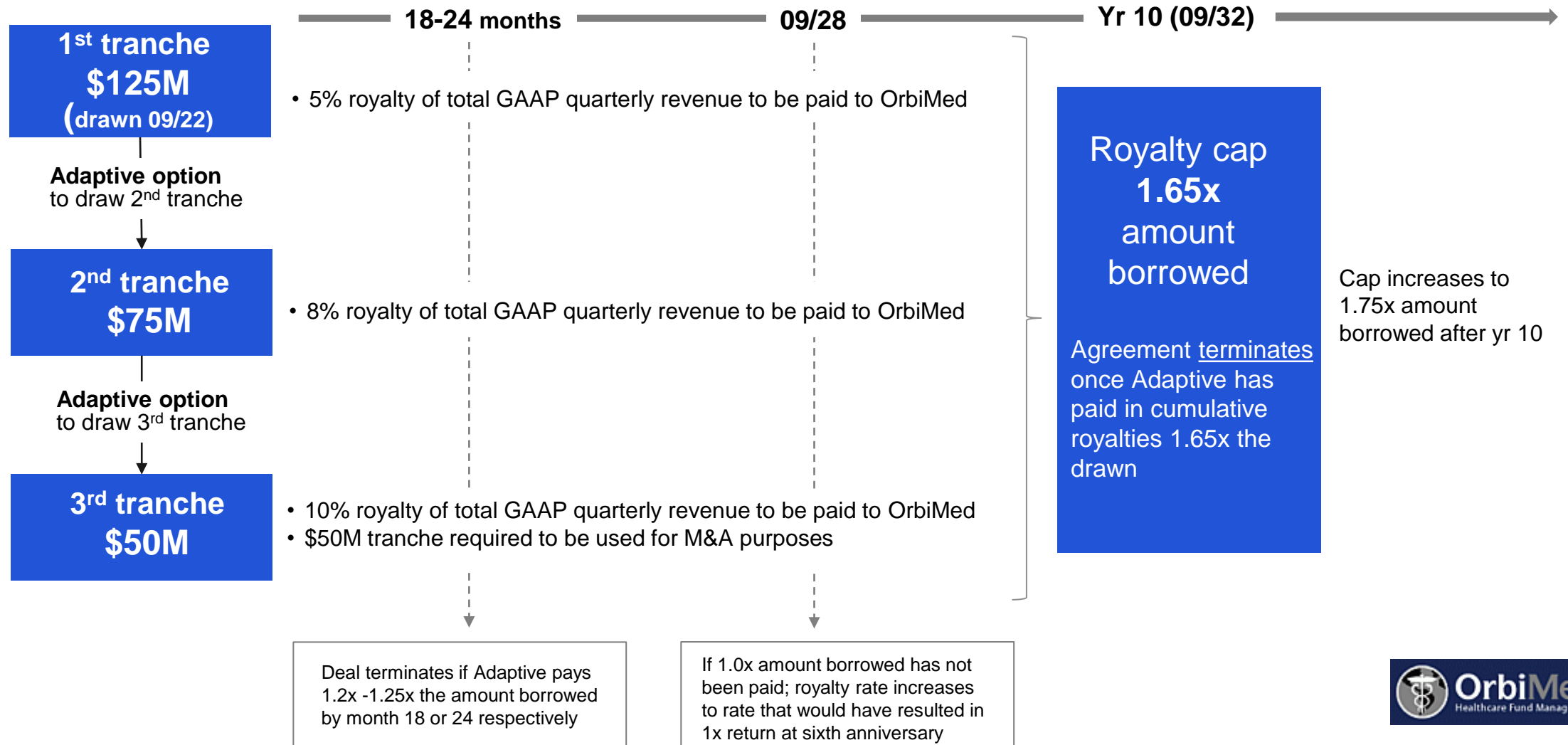
FY 2022 Revenue Guidance

- **Narrowing** range to \$185M-\$190M from \$185M-\$195M
 - MRD and Immune Medicine represents ~47% / 53% of total revenue at mid-point of range

FY 2022 Opex Guidance

- **Updated** FY to <\$400M vs. \$410M-\$415M previously

Royalty financing agreement for up to \$250M



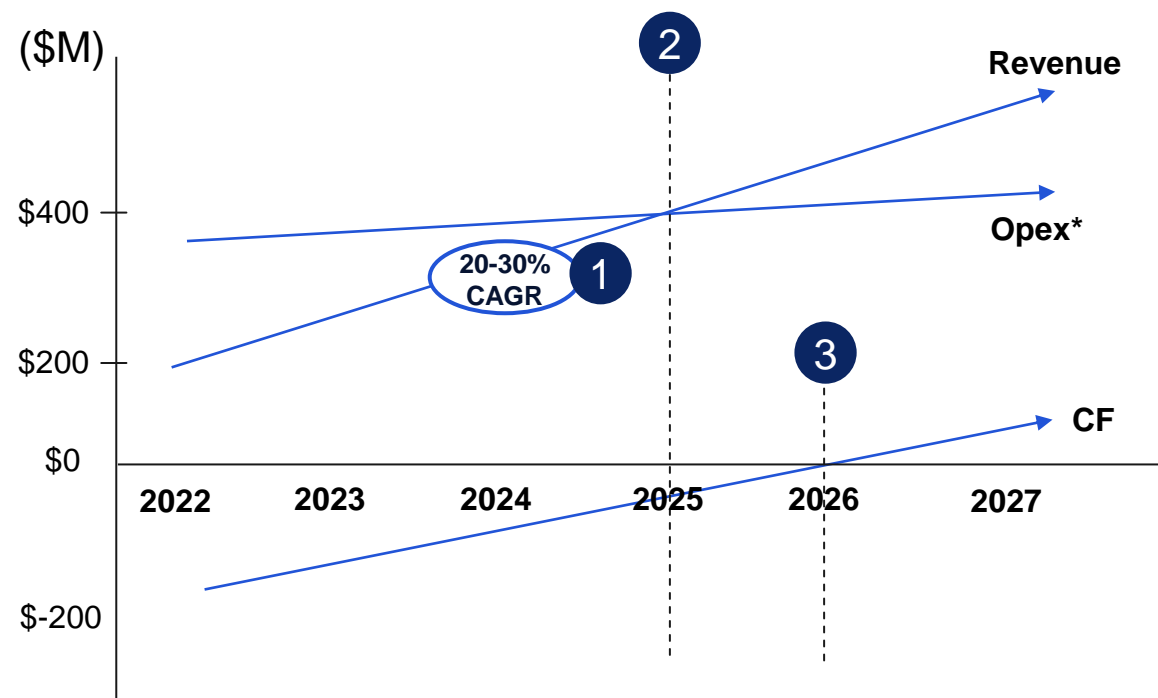
Long-term expectations

Path to Profitability / Cash Flow breakeven

- 1 Revenue CAGR from 2022-2027 to be 20-30%
 - MRD contribution higher in the near-term
- 2 Adj EBITDA¹ positive 2025
 - Prudent spend management: maintain operating expenses levels at low growth
- 3 Cash Flow Breakeven 2026
 - Cash on hand >3 years

¹ Adjusted EBITDA excludes stock comp

Estimated 5 yrs P&L progression



* Opex in this chart excludes stock comp, depreciation and amortization
Chart not at scale

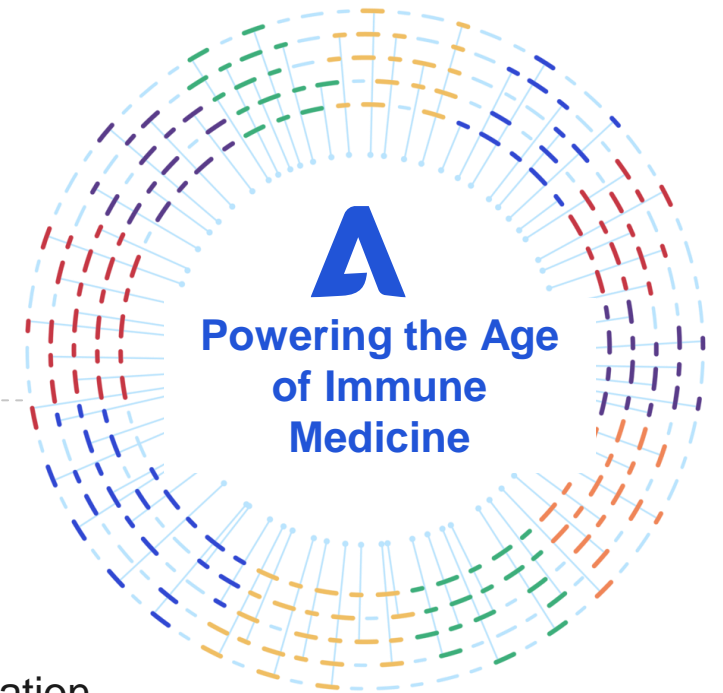
Updated Key Catalysts 2022

MRD

- ✓ Medicare **coverage of DLBCL**
- ✓ CLIA **launch DLBCL**
- ✓ Expand adoption of MRD status as a co-/primary clinical endpoint
- **Read-out data** for use in blood in MM/DLBCL

Immune Medicine

- Genentech collaboration:
 - ✓ Selected TCR candidate to progress as a potential therapeutic product candidate
 - Deliver 2 additional TCR data packages for consideration
 - Establish private product specifications
- Scale drug discovery opportunities with pharma





Thank You.

Reconciliation between Adjusted EBITDA and net loss attributable to Adaptive Biotechnologies Corporation

	Three Months Ended September 30,		Nine Months Ended September 30,	
	2022	2021	2022	2021
Net loss attributable to Adaptive Biotechnologies Corporation	\$ (45,281)	\$ (55,903)	\$ (160,063)	\$ (145,846)
Interest and other income, net	(765)	(327)	(1,454)	(1,429)
Interest expense	653	—	653	—
Depreciation and amortization expense	5,383	3,528	15,634	9,104
Restructuring expense	—	—	2,023	—
Share-based compensation expense	14,142	11,643	41,183	31,376
Adjusted EBITDA	\$ (25,868)	\$ (41,059)	\$ (102,024)	\$ (106,795)