

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 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, 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 Report 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 29, 2024. 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 forwardlooking statements. Except as required by applicable law, we do not plan to publicly update or revise any forward-looking statements contained herein.

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



Q3 highlights: continued to fuel growth with stronger financial profile



Fueling growth

- Q3 MRD revenue \$37.5M
 - +52% Y/Y; +32% Y/Y excluding MRD milestone
- Strengthened clinical profile with increased gapfill rate/test
- Fueling growth: MCL coverage,+ODAC impact



Reducing spend

- Q3 Operating spend
 - -11% Y/Y
- Gross Margin 64%
 - 56% sequencing GM; 13 ppts increase Y/Y¹
 - Significant steps taken to reduce cost/sample in the lab



Managing cash

- ~\$267M in cash²
 - 38% reduction in YTD³ cash burn (~\$79M in '24 vs ~\$127M in '23)
- Focused on targeted IM investments with highest upside value



¹ Excludes one-time lab move costs in Q3 2023

² Cash, cash equivalents and marketable securities as of 9/30/2024

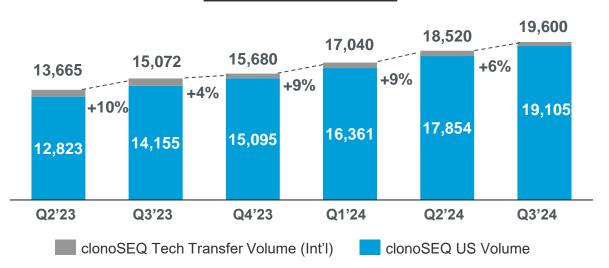
³ YTD = as of Q3 for the respective periods

MRD clinical testing: strong performance with milestones fueling future growth

Q3 clonoSEQ® clinical revenue growth of 39% Y/Y

- Volume: clonoSEQ test volume +30% Y/Y; +6% Q/Q
- Blood based: 41% of clonoSEQ MRD test in blood
- Ordering HCPs: 2,974; 39% growth Y/Y
- Ordering accounts: 610; 16% growth Y/Y

clonoSEQ test volumes



Key milestones and future drivers of growth

- Gapfill rate + ASP growth initiatives
 - New Medicare gapfill rate at \$2,007/test
- MCL Medicare coverage
 - Obtained Medicare coverage for MCL at new gapfill rate
- EPIC integrations
 - 11 accounts completed to date
 - Largest account live



MRD pharma: strong growth; continued positive momentum from ODAC

Q3 2024 revenue growth of 73% Y/Y, including recognition of \$5.0M in regulatory milestone revenue

ODAC vote on MRD as endpoint in MM: palpable impact on our pharma and clinical businesses



BioPharma increasing MM investments

- 16 new studies in MM closed YTD
- 11 closed post ODAC vote

Increasing use of MRD as primary endpoint

- 3 studies converted secondary to primary
- 2 under consideration for upgrade

Halo effect in clinical use

- Growing recognition of clinical utility will drive MRD adoption in clinics
- New protocols depend on MRD testing availability



Immune Medicine (IM) programs in cancer and autoimmunity

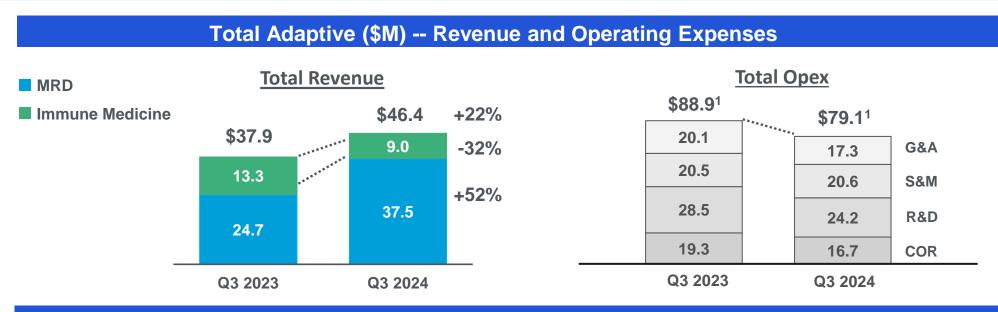
Program		Overview	Progress
Cancer	Cell Therapy	TCR-based cell therapy products targeting tumor neoantigens	 Establishing personalized product profile with highest POS and a focus on reducing cost and TAT (days vs. weeks)
Autoimmunity	Antibody Dev	Develop antibodies that target and eliminate or block disease-causing T cells	 Successfully completed several mouse immunization campaigns in MS, T1D and several other indications On track to select differentiated antibodies and start functional characterization by YE
	Target Discovery	Target discovery in select, prioritized autoimmune indications	 Making good progress in T1D to confirm disease-causing biology and find the protein to which 'pathogenic' TCRs bind

POS = probability of success
TAT = turnaround time

Gate R&D investments ● Make quick go/no-go decisions ● Reduce cash burn



Q3 2024 financial highlights



Segment Reporting

(\$M)	MRD				IM		Unallocated Corporate				
	Q1'24	Q2'24	Q3'24	Q1'24	Q2'24	Q3'24	Q1'24	Q2'24	Q3'24		
Revenue	32.6	35.3	37.5	9.2	7.9	9.0		N/A			
Opex ²	59.9	55.5	52.5	23.8	21.7	20.7	6.9	6.0	5.8		
Adj. EBITDA ³	(17.3)	(11.3)	(6.1)	(6.9)	(7.0)	(5.2)	(4.0)	(3.1)	(3.0)		

¹ Includes ~\$0.4M in amortization of intangible assets; ² Excludes one-time long-lived asset impairment charges of \$7.2M in Q2'24



³ Adj. EBITDA is a non-GAAP financial measure.

FY 2024 updated guidance

Narrowed FY 2024 revenue guidance range:

MRD revenue between \$143M and \$145M vs previous guidance of \$140M-\$145M

Reduced FY 2024 operating expenses:

OPEX between \$335M and \$340M¹ vs previous guidance of \$340M-\$350M¹

Reduced FY 2024 annual cash burn:

Cash burn of \$105M¹ vs previous estimate of \$115M¹



¹ Excluding one-time costs from strategic review pertaining to resources elimination

Appendix: Reconciliations between Adjusted EBITDA and net loss attributable to Adaptive Biotechnologies Corporation & Segment Information

■ The following table sets forth a reconciliation between our Adjusted EBITDA and net loss attributable to Adaptive Biotechnologies Corporation, the most directly comparable GAAP financial measure, for each of the periods presented (in thousands):

	Three Months Ended September 30,					Nine Months Ended September 30,				
		2024		2023		2024		2023		
Net loss attributable to Adaptive Biotechnologies Corporation	\$	(32,071)	\$	(50,300)	\$	(125,800)	\$	(155,809)		
Interest and other income, net		(3,474)		(4,282)		(11,462)		(10,918)		
Interest expense		2,939		3,652		8,628		10,788		
Depreciation and amortization expense		4,591		5,763		14,808		16,839		
Impairment of long-lived assets		_				7,205				
Restructuring expense		193				1,917				
Share-based compensation expense		13,522		15,336		40,778		47,352		
Adjusted EBITDA	\$	(14,300)	\$	(29,831)	\$	(63,926)	\$	(91,748)		



Appendix: Reconciliations between Adjusted EBITDA and net loss attributable to Adaptive Biotechnologies Corporation & Segment Information

The following tables set forth our segment information for the three months ended September 30, 2024 and 2023 (in thousands):

	Three Months Ended September 30, 2024							
		MRD		Immune Medicine		Unallocated Corporate		Total
Revenue	\$	37,470	\$	8,965	\$	_	\$	46,435
Operating expenses		52,538		20,689		5,840		79,067
Adjusted EBITDA		(6,120)		(5,212)		(2,968)		(14,300)
Reconciliation of Net Loss to Adjusted EBITDA:								
Net loss	\$	(15,068)	\$	(11,724)	\$	(5,305)	\$	(32,097)
Net loss attributable to noncontrolling interest		_				26		26
Net loss attributable to Adaptive Biotechnologies Corporation		(15,068)		(11,724)		(5,279)		(32,071)
Interest and other income, net		_		_		(3,474)		(3,474)
Interest expense		_		_		2,939		2,939
Depreciation and amortization expense		2,428		1,728		435		4,591
Restructuring expense		167		26		_		193
Share-based compensation expense		6,353		4,758		2,411		13,522
Adjusted EBITDA	\$	(6,120)	\$	(5,212)	\$	(2,968)	\$	(14,300)

	Three Months Ended September 30, 2023								
		MRD		Immune Medicine		Unallocated Corporate		Total	
Revenue	\$	24,668	\$	13,251	\$	_	\$	37,919	
Operating expenses		55,977		26,400		6,498		88,875	
Adjusted EBITDA		(21,616)		(4,986)		(3,229)		(29,831)	
Reconciliation of Net Loss to Adjusted EBITDA:									
Net loss	\$	(31,309)	\$	(13,148)	\$	(5,869)	\$	(50,326)	
Net loss attributable to noncontrolling interest		_		_		26		26	
Net loss attributable to Adaptive Biotechnologies Corporation		(31,309)		(13,148)		(5,843)		(50,300)	
Interest and other income, net		_		_		(4,282)		(4,282)	
Interest expense		_		_		3,652		3,652	
Depreciation and amortization expense		2,489		2,546		728		5,763	
Share-based compensation expense		7,204		5,616		2,516		15,336	
Adjusted EBITDA	\$	(21,616)	\$	(4,986)	\$	(3,229)	\$	(29,831)	



Appendix: Reconciliations between Adjusted EBITDA and net loss attributable to Adaptive Biotechnologies Corporation & Segment Information

■ The following tables set forth our segment information for the nine months ended September 30, 2024 and 2023 (in thousands):

	Nine Months Ended September 30, 2024								
		MRD		Immune Medicine		Unallocated Corporate		Total	
Revenue	\$	105,380	\$	26,118	\$	_	\$	131,498	
Operating expenses		170,785		70,663		18,762		260,210	
Adjusted EBITDA		(34,668)		(19,172)		(10,086)		(63,926)	
Reconciliation of Net Loss to Adjusted EBITDA:									
Net loss	\$	(65,405)	\$	(44,545)	\$	(15,928)	\$	(125,878)	
Net loss attributable to noncontrolling interest		_		_		78		78	
Net loss attributable to Adaptive Biotechnologies Corporation		(65,405)		(44,545)		(15,850)		(125,800)	
Interest and other income, net		_		_		(11,462)		(11,462)	
Interest expense		_		_		8,628		8,628	
Depreciation and amortization expense		7,733		5,777		1,298		14,808	
Impairment of long-lived assets		2,819		4,386		-		7,205	
Restructuring expense		1,195		722		_		1,917	
Share-based compensation expense		18,990		14,488		7,300		40,778	
Adjusted EBITDA	\$	(34,668)	\$	(19,172)	\$	(10,086)	\$	(63,926)	

	Nine Months Ended September 30, 2023							
		MRD		Immune Medicine		Unallocated Corporate		Total
Revenue	\$	71,977	\$	52,515	\$		\$	124,492
Operating expenses		170,946		88,753		20,760		280,459
Adjusted EBITDA		(71,081)		(11,149)		(9,518)		(91,748)
Reconciliation of Net Loss to Adjusted EBITDA:								
Net loss	\$	(98,969)	\$	(36,236)	\$	(20,632)	\$	(155,837)
Net loss attributable to noncontrolling interest		_				28		28
Net loss attributable to Adaptive Biotechnologies Corporation		(98,969)		(36,236)		(20,604)		(155,809)
Interest and other income, net		_		_		(10,918)		(10,918)
Interest expense		_		_		10,788		10,788
Depreciation and amortization expense		6,812		7,907		2,120		16,839
Share-based compensation expense		21,076		17,180		9,096		47,352
Adjusted EBITDA	\$	(71,081)	\$	(11,149)	\$	(9,518)	\$	(91,748)

