

#### **Safe Harbor**

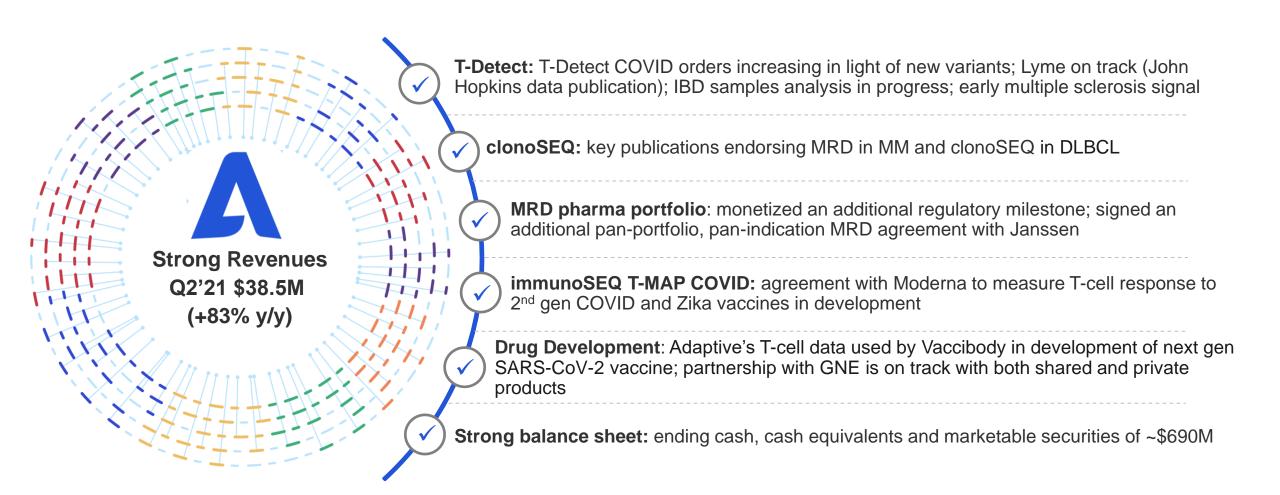
This presentation has been prepared by Adaptive Biotechnologies Corporation ("we," "us," "our," "Adaptive" or the "Company") and is made for informational purposes only. The information set forth herein does not purport to be complete or to contain all relevant information. Statements contained herein are made as of the date of this presentation unless stated otherwise. This presentation shall not constitute an offer to sell or the solicitation of an offer to buy securities, nor shall there be any sale of these securities in any jurisdiction in which such offer, solicitation or sale would be unlawful prior to registration or qualification under the securities laws of any such jurisdiction.

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," "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, including clonoSEQ, immunoSEQ and T-Detect products; discovery and development of neutralizing antibodies, 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, including the collaboration evidenced by our license agreement with Vaccibody; the estimated size of the market for our products and product candidates; the timing and success of our development and commercialization of our anticipated 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 24, 2021. 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.

Certain information contained in this presentation relates to or is based on studies, publications, surveys and other data obtained from third-party sources and the Company's own internal estimates and research. While the Company believes these third-party sources to be reliable as of the date of this presentation, it has not independently verified, and makes no representation as to the adequacy, fairness, accuracy or completeness of, any information obtained from third-party sources. In addition, all of the market data included in this presentation involves a number of assumptions and limitations, and there can be no guarantee as to the accuracy or reliability of such assumptions. Finally, while we believe our own internal research is reliable, such research has not been verified by any independent source.

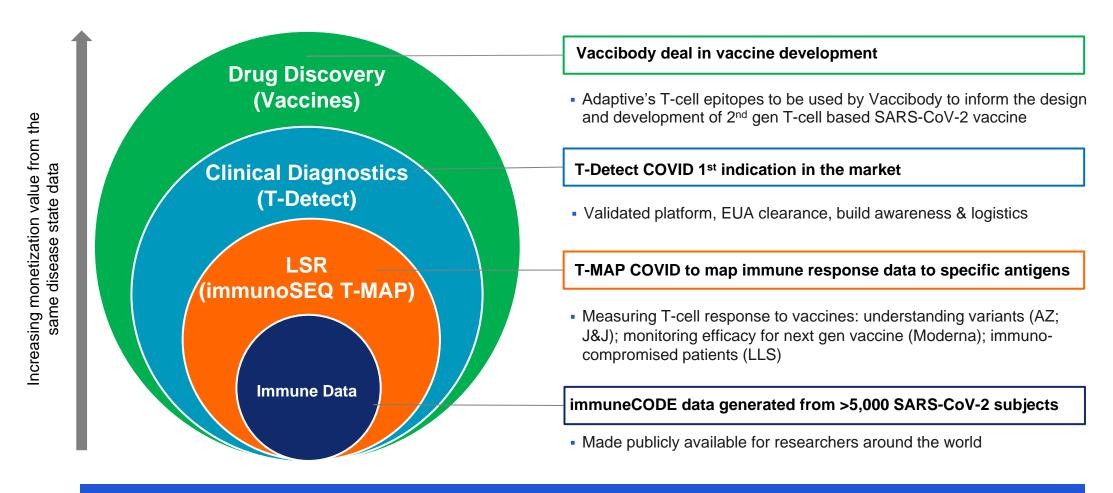


## **Q2 2021 Key Highlights**





## Value creation opportunities stemming from core data



Using disease-specific immune receptor data done for COVID, can be applied to ANY disease

## T-Detect: acceleration towards differential diagnosis & signal generation

1

2

3

## Improve SOC Dx Disease by disease

#### **T-Detect COVID**

- >10,000 tests ordered since launch
- Immune Response Website coming soon

#### **T-Detect Lyme**

- Publication of control data with John Hopkins
- ImmuneSense Lyme enrollment on track; completion by YE
- Commercial readiness underway – test online for CLIA around YE

## **Differential Diagnosis**Diseases with shared symptoms

#### **T-Detect IBD/Other**

- >5,000 IBD samples under current analysis. Data completion in 2021
- Significant market opportunity with high unmet need -- VOC (GIs, PCP) and payer research supports upstream value opportunity
- Early signal in Multiple Sclerosis



## Population Immunomics Any mapped disease

#### **Aspirational long-term vision**

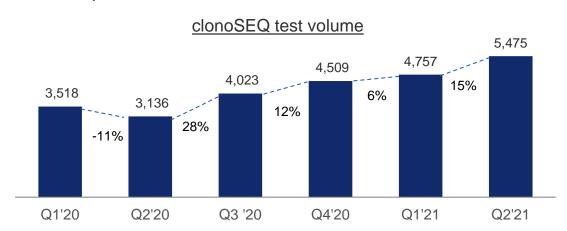
One blood test with many results



### clonoSEQ brand: Increase in clinical adoption and pharma partnerships

#### **clonoSEQ Clinical Testing**

- Q2'21 test delivered volume +75% vs P/Y; +15% vs P/Q
  - ~250 ordering accounts; ~950 ordering HCPs in Q2
- High-impact recent publications:
  - Use of MRD to guide clinical care for patients with MM
  - Use of MRD as a regulatory endpoint in clinical trials
  - clonoSEQ MRD assessment using ctDNA in DLBCL patients



#### **MRD Pharma Partners**

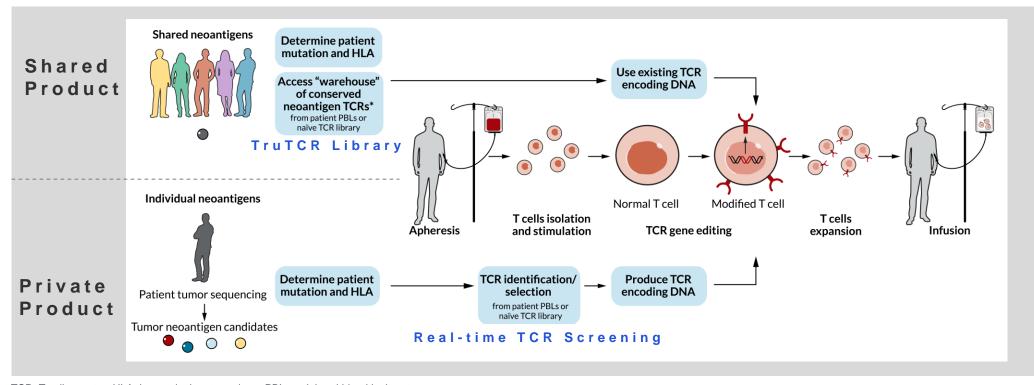
- Strong sequencing revenue growth Q/Q
- \$1.5M in development milestones in Q2
- Expanded MRD agreement with Janssen to a pan portfolio collaboration
  - Structured to contribute sequencing and development revenue





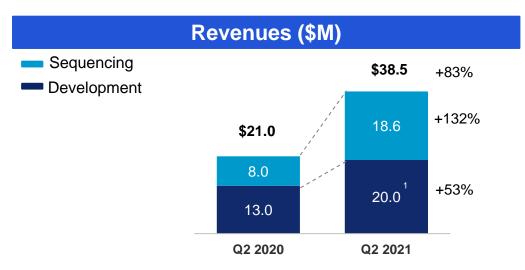
## Developing transformative T cell therapies in oncology

- Focus on advancing shared product and scaling private product development continues
  - Shared Product: on track to complete and deliver to GNE next shared product candidate data package by YE 2021
  - Private Product: on track to process blood from ~60 cancer patients by year end
- Advancing programs in parallel; clinical leveraging from the shared product(s) may also inform our private product process

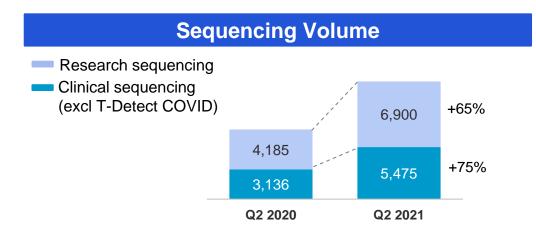


TCR: T cell receptor, HLA: human leukocyte antigen; PBL: peripheral blood leukocytes

### **Q2 2021 Key Financial Highlights**



<sup>&</sup>lt;sup>1</sup> Includes \$1.5M in MRD milestones



#### **Operating Expenses (\$M)**



#### **Balance Sheet & 2021 Guidance**

- \$689.5M in cash, cash equivalents and marketable securities as of 06/30/2021
- Revised 2021 full year revenue range \$148M \$155M



### 2021 2H upcoming catalysts

## Life Science Research

- Embed T-MAP COVID into SARS-CoV-2 vaccine trials
- Expand immunoSEQ T-MAP data offering into other disease categories
- Expand distribution of immunoSEQ RUO kit

# Clinical Diagnostics

- ImmuneSense study completion and T-Detect Lyme launch through CLIA
- T-Detect IBD: analysis of >5,000 IBD samples
- Confirm additional T-Detect signal(s)
- FDA clearance for clonoSEQ ALL in blood

#### **Drug Discovery**

- Complete 2nd shared product data package for GNE by YE
- Private product proof of concept data for ~60 cancer patients
- Vaccibody to initiate phase I/II study in Q4 2021 for T-cell SARS-CoV-2 vaccine
- Define neutralizing antibody pathway





## Clinical portfolio and pipeline

Diagnostic Product Plan	Signal Discovery	Clinical Validation	FDA Submission	FDA Clearance
Monitor MRD: clonoSEQ*	Multiple Myeloma (bone marrow)			✓
	Acute Lymphoblastic Leukemia (bone marrow)			✓
	Chronic Lymphocytic Leukemia (blood, bone marrow)			✓
	Non-Hodgkin's Lymphoma (Subtypes) <sup>1</sup>			
Accurate Detection:  T-Detect	COVID-19 <sup>2</sup>			(EUA)
	Lyme Disease			
	GI Diseases			
	<ul> <li>Ovarian cancer</li> </ul>			
Drug Discovery Product Plan	Early Development	IND Submission	Clinical Development	
TCR-Based Cell Therapies <sup>3</sup>	Shared			
	Personalized			
Neutralizing Antibodies <sup>4</sup>	— COVID-19			
Vaccines <sup>5</sup>	COVID-19			

clonoSEQ NHL subtypes: available to order as a CLIA-validated laboratory developed test (LDT) service. This use has not been cleared or approved by the FDA.
 T-Detect COVID: has received Emergency Use Authorization and is not FDA cleared or approved.
 TCR-Based Cell Therapies: product candidates in development as part of our worldwide collaboration and license agreement with Genentech. The product candidates refer to the lead clinical product candidates selected from our library of TCRs that target cancer antigens present in many cancer patients. Genentech will determine the timing of discussions with, and submissions to the FDA.
 Neutralizing Antibodies: Product candidates in development.
 Vaccines: Product candidates in development