

## **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 various disease states, including T cell responses to COVID-19 or other infectious diseases, as well as autoimmune disorders and cancer, the ability to leverage any such findings to advance solutions to diagnose, treat and prevent infectious diseases or other 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 T-MAP COVID and T-Detect products, planned non-IDE clinical studies, clinical trials and preclinical activities, research and development costs, current and prospective collaborations, including our collaboration with Genentech, the estimated size of the market for our products and product candidates; the timing and success of our development and commercialization of our current and anticipated product candidates and the expansion of existing product lines such as T-Detect into additional indications; the availability and extent of reimbursement coverage by government and private payors; 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 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.

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.

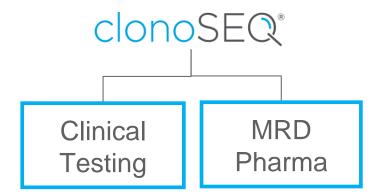


# **Business areas of focus going forward**

## **MRD**

Driven by clonoSEQ/MRD assay applications

TAM ~\$6B<sup>1</sup>

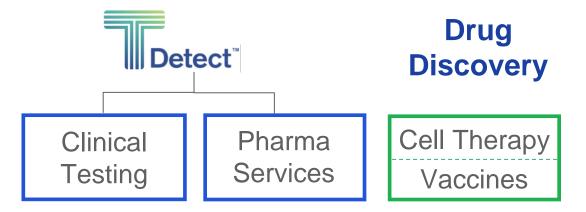


1. Global TAM: \$4.5B clonoSEQ clinical testing; \$1.5B Pharma partnerships (including regulatory milestones)

## **Immune Medicine**

Driven by immune receptor data opportunities

TAM ~\$48B<sup>2</sup>



2. Global TAM: illustrative TAM for 3 indications for T-Detect (one infectious disease, one autoimmune disease, one oncology) and drug discovery in cell therapy oncology



# **Key Achievements through 2021**



#### **MRD Business**

- Key data read outs demonstrating clinical utility
- Expanded commercial team, added product line enhancements for CLL patients
- Signed 2 significant pan-portfolio MRD partnerships across all heme indications
- Recognized \$10M in milestones from pharma partners

#### **Immune Medicine Business**

- Genentech selected TCR candidate targeting a shared cancer neoantigen
- Successfully completed initial POC screens for personalized product
- Extended platform to vaccines with Nykode collaboration; phase 1/2 enrolling
- T-Detect COVID: granted EUA by the FDA with >30k tests ordered
- Completed first T-Detect clinical validation study with ImmuneSense Lyme
- Identified multiple T-Detect signals in autoimmune disorders
- T-MAP COVID agreements with partners including J&J, AZ, Moderna



# Our MRD Heme business: synergistic value of pharma and clinical diagnostic

## **Tests Delivered '21**

•Q4: 6,356 (+7% vs Q3)

•FY: 22,516 (+48% Y/Y)

## **Ordering HCPs**

•1,735 (+57% Y/Y)

## **Ordering Accounts**

• 372 (+32% Y/Y)



> 60 Companies

155 Active trials

> \$330M Milestones



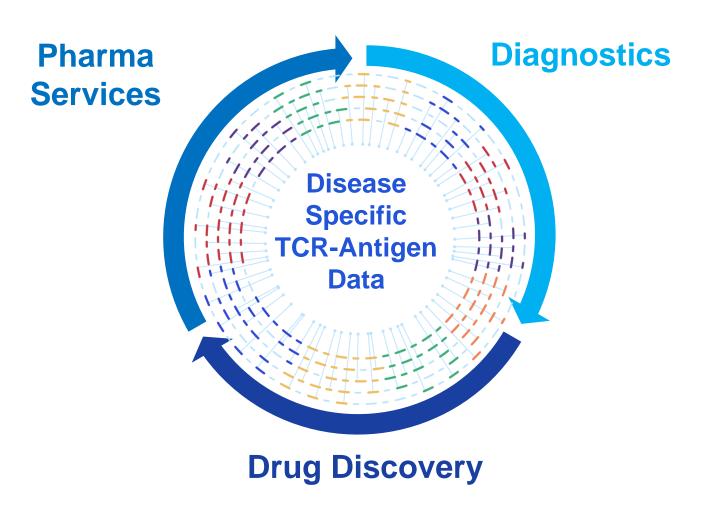
Embedded as clinical endpoint across multiple drug modalities and diseases



# Unique ability to map & identify disease specific TCR sequences

# Immune Medicine Strategy

Create multiple value opportunities stemming from the same core disease data





# T-Detect, advancing towards "one test, many diseases"

#### COVID



**Only** FDA authorized test, available to consumers



- Focus on studying correlate of protection for T cells
- Enable vaccine manufacturers to incorporate T cell response

#### **AUTOIMMUNE**



Clinical signals in **five** diseases Crohn's, UC, celiac, MS, RA



- Maintain high specificity (rule-in) while improving sensitivity levels needed for CLIA launch
- Initiate clinical validation in IBD

## LYME



ImmuneSense Study Completed



- Enable CLIA testing during Lyme season
- Leverage consumer-pay experience



# Genentech selects TCR candidate against a shared cancer neoantigen



#### Cell Therapy - Progress on Shared and Private Programs

#### Shared

TCRs targeting shared cancer antigens



- TCR candidate selected to progress as a potential therapeutic product candidate
- Efficacy and safety data reviewed by both Adaptive and Genentech
- Deliver 2 additional TCR data packages for consideration by YE

#### Private

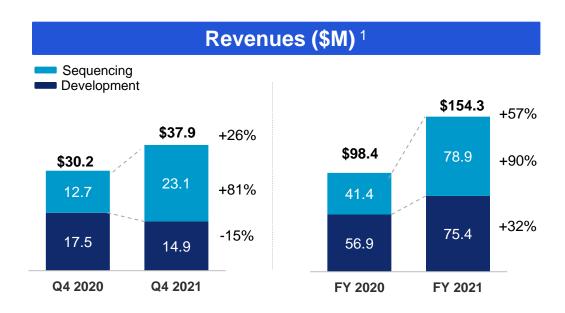
Build private product process

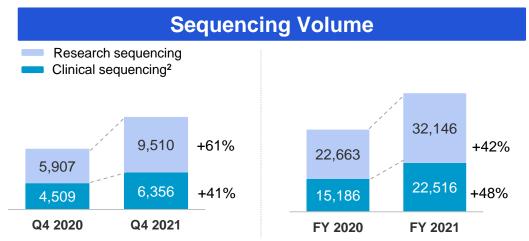


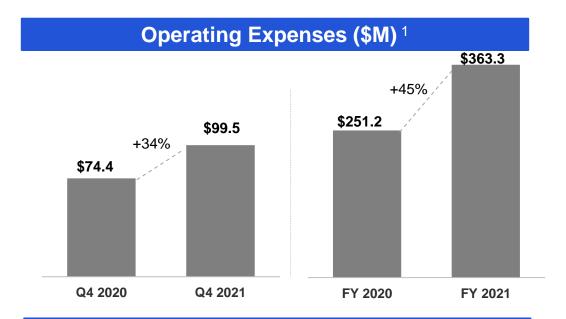
- Completed initial POC screens using samples from ~ 60 cancer patients
- Establish private product specifications and build data package
- Start to define steps toward early product development



# **Q4 and FY 2021 Financial Highlights**







#### **Strong Balance Sheet**

- \$570.2 M in cash, cash equivalents and marketable securities as of 12/31/2021
- No debt

<sup>1\$</sup> and % figures are rounded

<sup>&</sup>lt;sup>2</sup> Excludes T-Detect COVID volume

## FY 2022 Guidance

#### ■ Revenue: 2022 full year revenue range \$185M - \$195M

- □ Sequencing revenue represents ~60% of total revenue at mid-point of range
- Include low to mid double digit in potential milestones (risk-adjusted)
- Q1 expected to be the lowest quarter of the year with no milestone anticipated during the quarter

## **■ FY 2022 Operating Expenses:**

- Decelerating expense growth rate versus 2021 reflecting leverage from significant investments made in 2021
- Targeting operating expenses to grow at lower rates than revenue



# **Key Catalysts 2022 -- Multiple levers to drive value**

# **Immune Medicine**

- **T-Detect COVID**: Enhance product profile (correlate of protection)
- **T-Detect AI**: Increase sensitivity/specificity in MS, IBD, RA for market readiness
- Genentech collaboration:
  - ✓ Selected TCR candidate to progress as a potential therapeutic product candidate
  - On track to deliver 2 additional TCR data packages for consideration
  - Establish private product specifications
- Nykode collaboration: phase 1/2 data

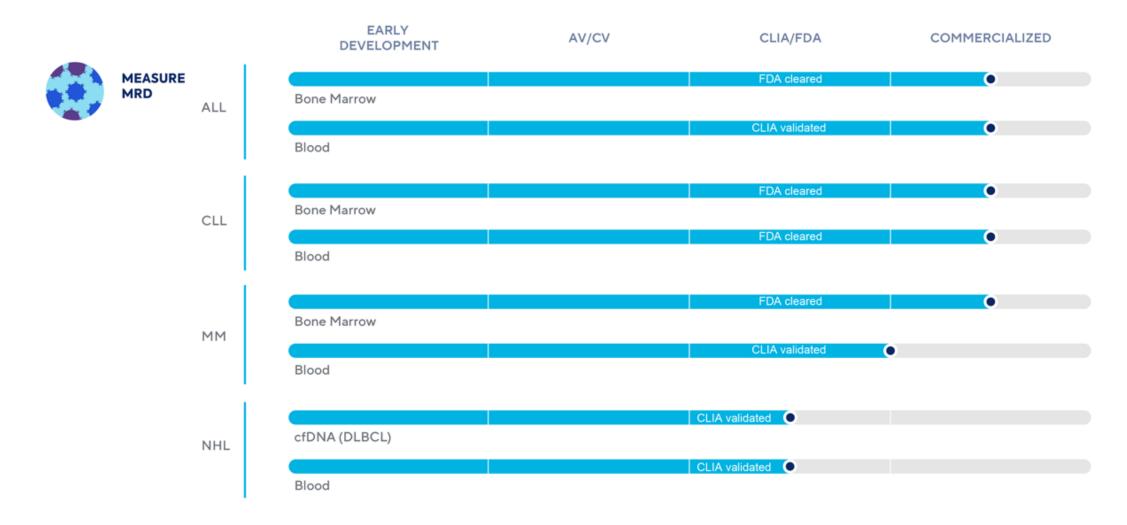
# **MRD**

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





# **Appendix: clonoSEQ Pipeline**





# **Appendix: Drug Discovery Pipeline**





# **Appendix: T-Detect Pipeline**



