

Safe Harbor

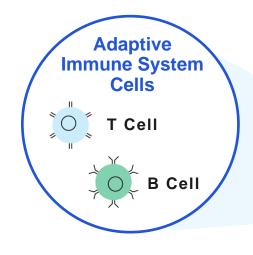
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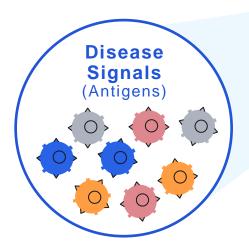
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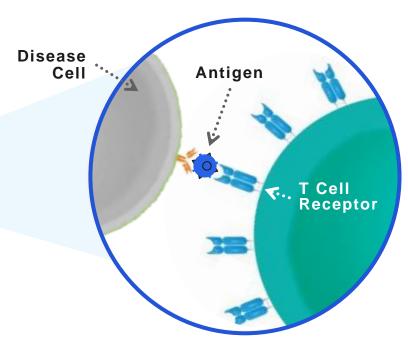
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The Immune System Detects & Treats Most Diseases in the Same Way







Receptors bind to specific signals of disease



Using the immune system as the source-code for immune medicine





Business areas of focus

MRD

TAM ~\$6B¹

Highly sensitive NGS-based assessment of minimal residual disease for use in clinical practice and drug trials.

Clinical Testing



MRD Pharma Partnerships

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

Immune Medicine

TAM ~\$48B²

Clinical diagnostics, drug discovery, and research informed by our TCR-Antigen Map.

Clinical Testing

T-Detect"

Drug Discovery

- T-Cell therapeutics
- Antibodies
- Vaccines

Immune Medicine Partnerships

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



The Management Team



Chad Robins
Chief Executive Officer,
Co-founder,
Chairman of the Board



Harlan Robins, PhD
Chief Scientific Officer
& Co-founder



Julie Rubinstein
President



Tycho Peterson
Chief Financial Officer



Sharon Benzeno, PhD
Chief Commercial Officer, Immune
Medicine



Nitin Sood
Chief Commercial Officer, MRD



Mark Adams, PhD
Chief Operating Officer



Francis LoChief People Officer

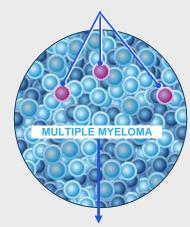


MRD

Monitoring minimal residual disease (MRD) in select blood cancers

- Patients are living longer on new therapies
- Clinicians need to monitor disease burden
- Pharma needs earlier response measures
- Guidelines include MRD in multiple diseases





Patient-specific clonal sequence

- Universal applicability to patients with lymphoid malignancies
- Sensitivity of 1 in a million cells*
- Minimally invasive



^{*} Given sufficient sample input

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



- FDA cleared in MM, ALL (BM*) and CLL (BM*, blood)
- Broad coverage: >240M covered lives in MM & ALL; >150M in CLL
- Widely adopted: Clinical use in all 31 NCCN centers



 Pharma supports lifecycle expansion which drives clinical use

 Clinical usage drives inclusion as an endpoint in pharma trials MRD Pharma Trials

Broad use: most major pharmaceutical companies developing lymphoid cancer drugs use clonoSEQ as the test of choice in their trials.





MRD Business: clonoSEQ clinical testing

Growth experienced across the board...

- FY '21 and Q1'22 test delivered grew +48% and +45% vs P/Y respectively
 - ~320 ordering accounts in Q1 (+36% vs P/Y)
 - ~1,200 ordering HCPs in Q1 (+53% vs P/Y)
- ~30% of MRD tests delivered by blood

clonoSEQ test volume



clonoSEQ test volume under new reporting to include tech transfer volume from international sites

Strategy to cement leadership in lymphoid cancers...

Three priority areas for investment

- HCP education & adoption: field force expansion & training
- Product development: expanding into NHL using cfDNA
- Customer experience: integrating into customer ordering systems

400K patients

Expand into

Multiple Myeloma in blood



~\$950-\$1000



Penetrate deeper in institutional accounts



NHL

ASP ~\$800

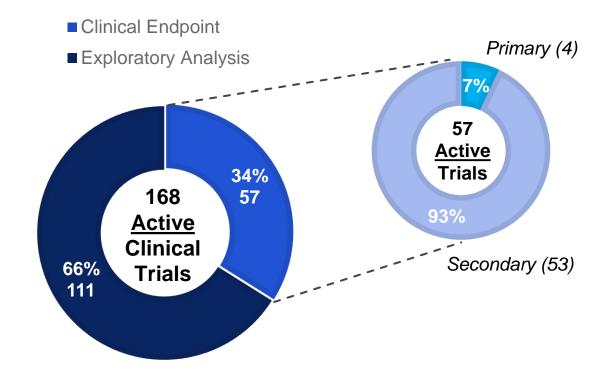
MRD Business: pharma portfolio

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

Portfolio Overview

- >60 companies, 168 active clinical trials
- 5 Recent FDA drug approvals containing clonoSEQ data (Blincyto, Darzalex, Abecma)
- Regulatory milestones: >\$330M in regulatory milestones available from active and future trials







MRD has evolved into an essential clinical decision-making tool

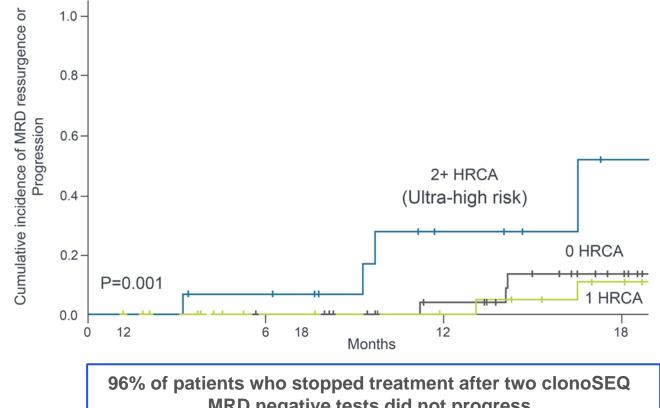
MASTER trial

Cumulative incidence of MRD resurgence or progression

Assess response to therapy

Escalate/de-escalate therapy

Detect relapse early



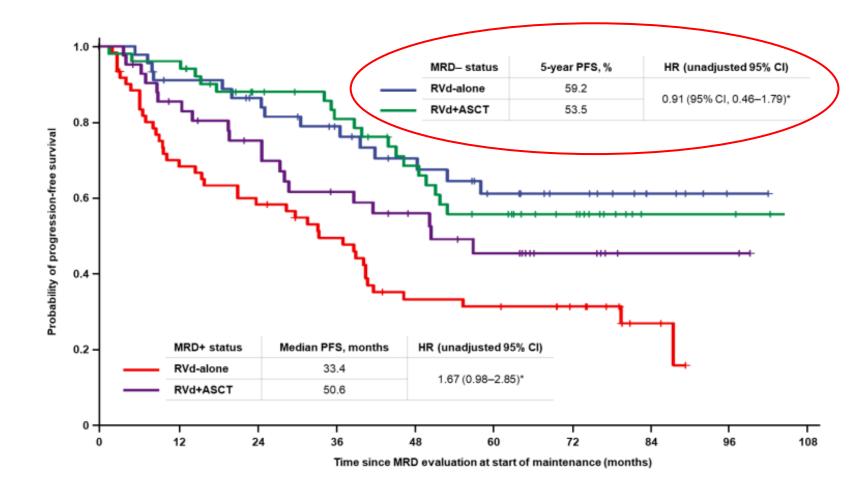
MRD negative tests did not progress



Patients who reached MRD negativity had the best outcomes, even without transplant

DETERMINATION trial

"The elimination of minimal residual disease is of increasing importance in tailoring treatment, in informing clinical care, and as a treatment goal given its prognostic value for better outcomes."



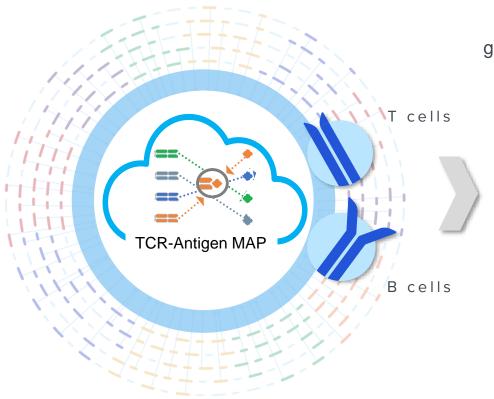




Immune Medicine

Platform synergies will drive growth opportunities and generate revenue

Immune Medicine Platform



3 Growth Areas

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

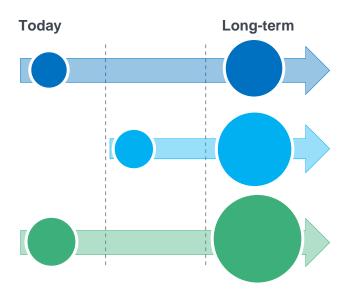
Pharma

Clinical Testing

Drug Discovery

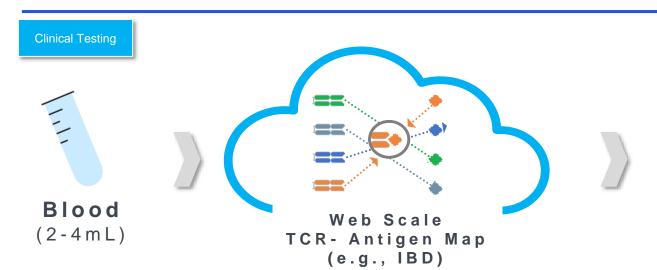
Revenue Contribution

(Illustrative)



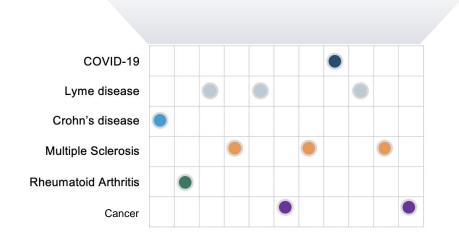


T-Detect[™] — A new class of **T-cell** based molecular test











Validate TCR-based disease signatures

Scale mapping of TCRs to disease antigens

Execute toward early disease interception



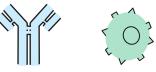
T-Detect platform — near term strategic priorities and status

<u>Infectious diseases (COVID, Lyme)</u>



To be pursued opportunistically

Autoimmune / inflammatory



Focus of T-Detect, given alignment with technology, high unmet need, spend, and biopharma interest

COVID

- Continue offering to consumer
- COP opportunities ongoing
- Launched T-Detect brand

Lyme

- CV study confirmed double sensitivity vs SOC/serology
- Build CLIA infrastructure
- Brand building

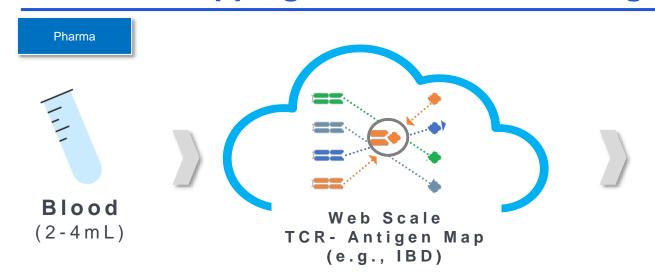
<u>GI</u>

Neuro

- On track to initiate clinical validation in IBD; deliver MVP* target
- Launch one autoimmune disease test by end of 2023
- Brand expansion



T-MAP — Mapping TCRs to disease antigens



Antigen-specific T-Cell Signature

immunoSEQ® T-MAP™

(detection + monitoring)



2022 YTD T-MAP deals

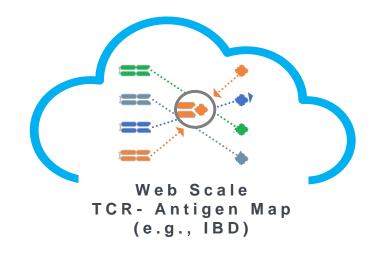
COVID T-MAP deals (Moderna, AZ, Janssen)

RSV T-MAP deal with Janssen



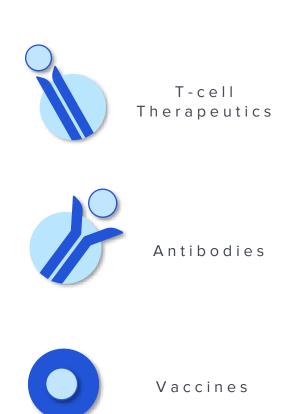
Deploy the platform to discover novel targets and develop differentiated therapeutics across three drug modalities

Drug Discovery





3 THERAPEUTIC MODALITIES





Cell Therapy in Oncology; Partnership with Genentech

Characterize TCRs against cancer antigens for cellular therapy



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

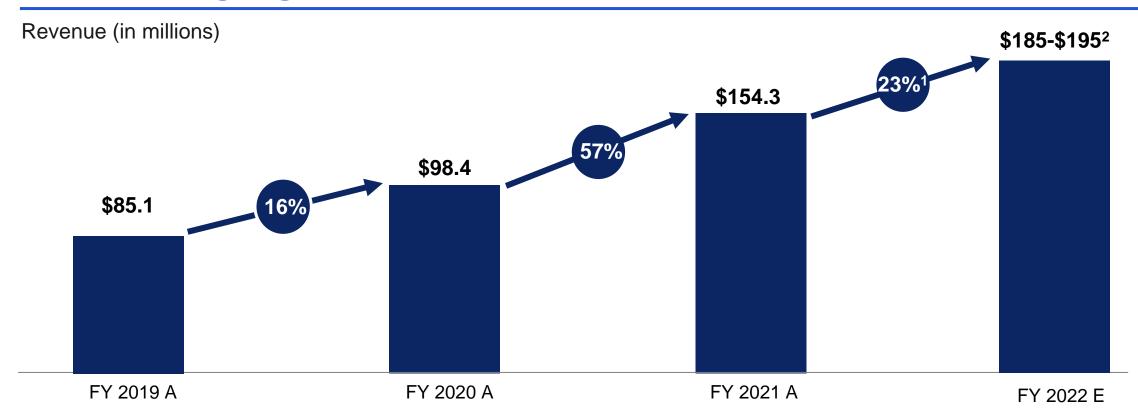
Build private product process



- 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



Financial highlights



\$500 million in cash, cash equivalents and marketable securities as of 03/31/2022. No Debt

Note: bar charts not at scale

2. FY 2022 revenue guidance range



^{1.} Implied growth rate at mid-point of 2022 revenue guidance range

Why Adaptive is well positioned to Thrive?

- ✓ Differentiated vision
- ✓ Unparalleled science & data
- ✓ Significant growth opportunity
- ✓ Diverse & experienced team
- ✓ Well capitalized



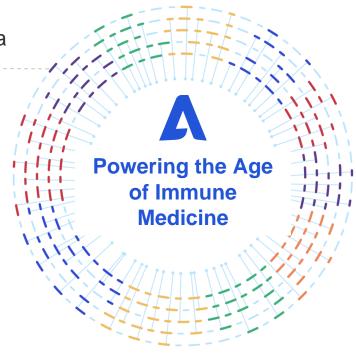
Key Catalysts 2022 – Multiple levers to drive value

Immune Medicine

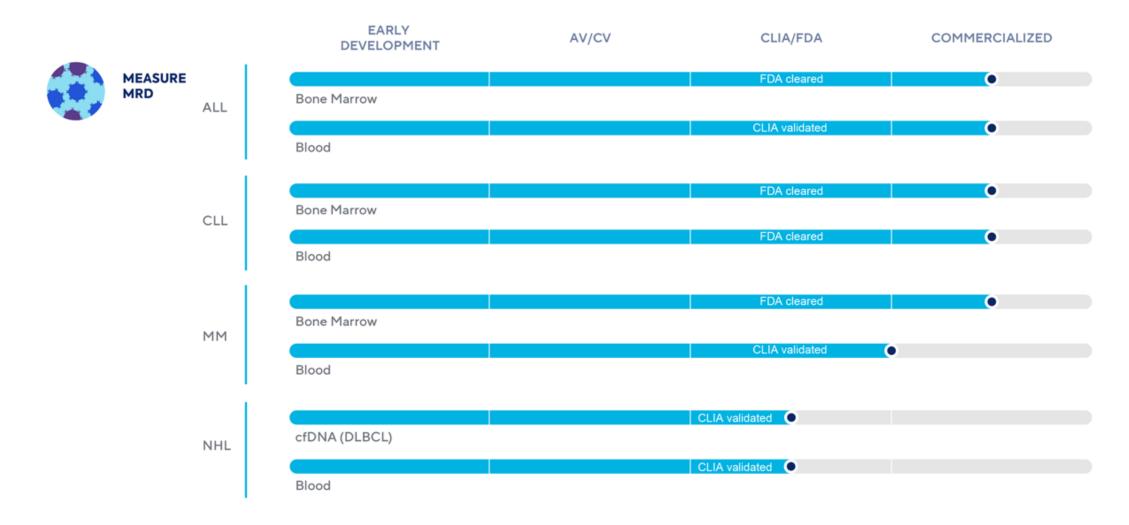
- **T-Detect COVID**: Enhance product profile (correlate of protection)
- **T-Detect Lyme**: T-Detect Lyme available through CLIA in 2H
- **T-Detect AI**: Increase sensitivity/specificity in IBD, MS for market readiness
- Genentech collaboration:
 - ✓ Selected TCR candidate to progress as a potential therapeutic product candidate
 - Deliver 2 additional TCR data packages for consideration
 - Establish private product specifications
- Nykode collaboration: Phase 1/2 clinical trial 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

