

Safe Harbor

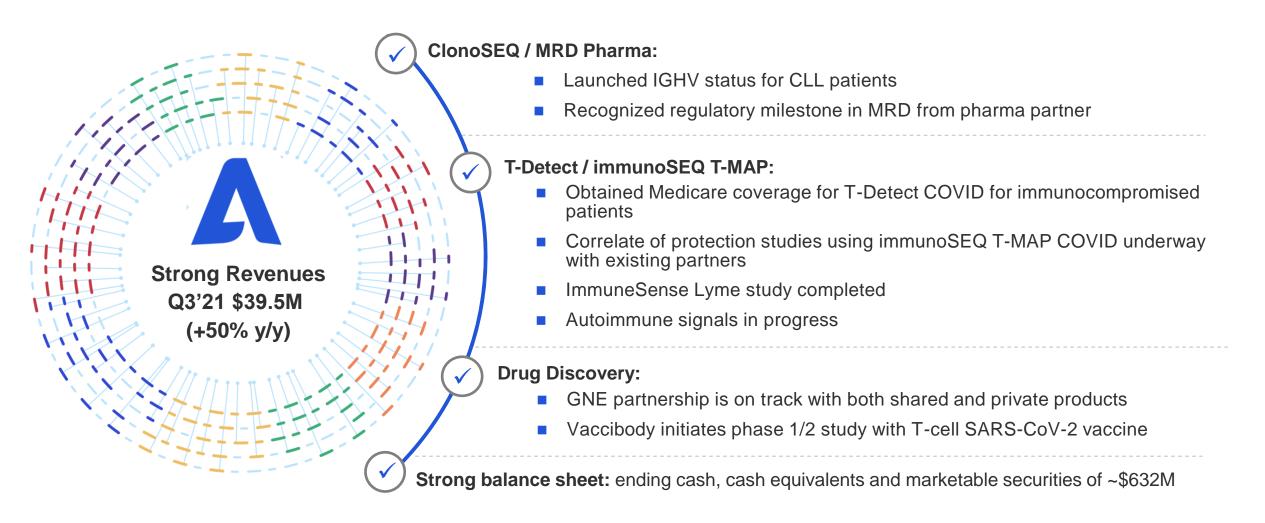
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Q3 2021 Key Highlights





clonoSEQ brand: Increase in clinical adoption and pharma partnerships

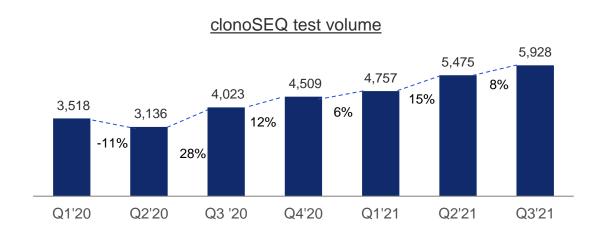
clonoSEQ Clinical Testing



MRD Pharma Portfolio

- Q3'21 test delivered volume +47% vs P/Y; +8% vs P/Q
 - ~260 ordering accounts; ~1,020 ordering HCPs in Q3
- Launched IGHV status for CLL patients
 - Enable HCPs to assess IGHV mutation status and obtain trackable sequence for MRD monitoring from one ID blood sample.

- Significant value driver to the overall MRD franchise
- Structured to contribute sequencing and development revenue
 - □ >\$330M in future regulatory milestones pipeline
 - Recognized \$1.5M in a development milestone in Q3







T-Detect COVID focus; leveraging Coverage and T-MAP pharma partnerships

T-Detect update: Infectious Disease



Leveraging Partnerships

COVID with Pharma, Biotech and Research

Widening adoption of immunoSEQ T-MAP COVID / T-Detect

COVID

Medicare coverage for immuno-compromised patients

- Prioritization of timing and resource allocation to market T-Detect COVID for immunocompromised patients and prepare formal reimbursement launch in 2022
- immuneSense Lyme study completed: nearly 2x more sensitive than STTT at detecting early signs of disease
- Make T-Detect Lyme available by Lyme season



Published data showing key role of T cell in the wake of variants; ongoing discussions to expand studies to assess correlate of protection



Correlate of protection in immunocompromised patients



Samples in house: effectiveness of vaccine in eliciting COVID specific T-cell response



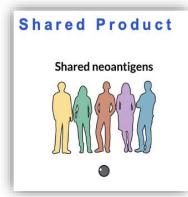
Understand T-cell response in individuals with AZ vaccine + booster developed by Clover

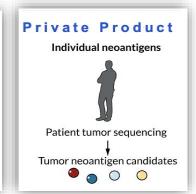
Progress on our open-ended growth story in Drug Discovery

Cell Therapy: identify TCRs as Therapeutics



- Shared Product: on track to assess efficacy and safety data with GNE to enable a decision by YE to move our lead product into early development
- Private Product: on track to process blood from ~60 cancer patients by YE





Vaccines: Identify Antigens for vaccines

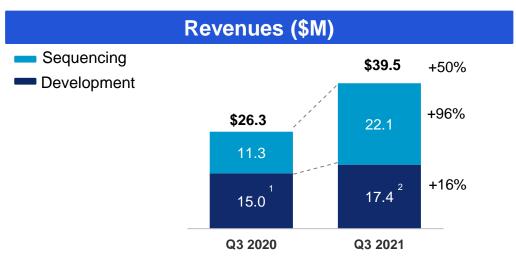


 Vaccibody announced start of 2-arm Phase 1/2 trial in previously vaccinated individuals with two vaccine candidates

- RBD Candidate: Vaccibody program designed to generate RBD specific antibody and T-cell immunity
- <u>T Cell Candidate</u>: Uses Adaptive identified and validated T cell epitopes; expected to result in a broader immune response against variants



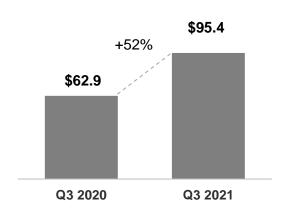
Q3 2021 Key Financial Highlights



¹ Includes \$2.5M in MRD milestones

Sequencing Volume Research sequencing Clinical sequencing (excl T-Detect COVID) 6,541 4,023 Research sequencing 8,710 5,928 +47%

Operating Expenses (\$M)³



³ Excludes amortization of intangible assets

Balance Sheet & 2021 Guidance

- ~\$632M in cash, cash equivalents and marketable securities as of 09/30/2021
- Reiterate 2021 full year revenue range \$148M \$155M



² Includes \$1.5M in MRD milestones

2021 remaining milestones

- Embed T-MAP COVID into SARS-CoV-2 vaccine studies
- T-Detect IBD: analysis of >5,000 IBD samples
- Confirm additional T-Detect signal(s)
- FDA decision for clonoSEQ ALL in blood
- Shared product: decision to move lead candidate into early development
- Private product: proof of concept data for ~60 cancer patients





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 (Subty	/pes) ¹		
Accurate Detection: T-Detect	COVID-19 ²			(EUA)
	Lyme Disease			
	GI Diseases			
	— Ovarian cancer			
Drug Discovery Product Plan	Discovery	IND Submission	Clinical Development	
TCR-Based Cell Therapies ³	Shared			
	Personalized			
Neutralizing Antibodies ⁴	— COVID-19			
Vaccines ⁵			—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