

Safe Harbor Statement

The statements made by Applied DNA in this presentation may be "forward-looking" in nature within the meaning of Section 27A of the Securities Act of 1933, Section 21E of the Securities Exchange Act of 1934 and the Private Securities Litigation Reform Act of 1995. Forward-looking statements describe Applied DNA's future plans, projections, strategies, and expectations, and are based on assumptions and involve a number of risks and uncertainties, many of which are beyond the control of Applied DNA. Actual results could differ materially from those projected due to its history of net losses, limited financial resources, unknown future demand for its biotherapeutics products and services, the unknown amount of revenues and profits that will result from the Linea™ IVT platform, limited market acceptance for its supply chain security products and services, the declining demand for Applied DNA's COVID-19 testing services, the fact that there has never been a commercial drug product utilizing PCR-produced DNA technology and/or the Linea IVT platform approved for therapeutic use, and various other factors detailed from time to time in Applied DNA's SEC reports and filings, including its Annual Report on Form 10-K filed on December 14, 2022, as amended, its 10-Q filed on February 9, 2023, May 11, 2023, and August 10, 2023, and other reports it files with the SEC, which are available at www.sec.gov. Applied DNA undertakes no obligation to update publicly any forward-looking statements to reflect new information, events, or circumstances after the date hereof or to reflect the occurrence of unanticipated events, unless otherwise required by law.

		Three Months Ended June 30,			Nine Months Ended June 30,			
		2023		2022		2023		2022
Revenues								
Product revenues	\$	316,950	\$	219,765	\$	1,130,800	\$	1,454,42
Service revenues		425,694		182,796		826,813		570,759
Clinical laboratory service revenues		2,174,697		3,893,810		10,630,094		12,584,174
Total revenues		2,917,341		4,296,371		12,587,707		14,609,360
Cost of product revenues		368,902		307,049		1,103,843		1,211,959
Cost of clinical laboratory service revenues		1,279,121		2,950,064		6,029,428		8,760,520
Total cost of revenues		1,648,023		3,257,113		7,133,271		9,972,479
Gross profit		1,269,318		1,039,258		5,454,436		4,636,881
Operating expenses:								
Selling, general and administrative		3,292,304		3,032,877		9,440,734		11,341,176
Research and development		836,123		863,025		2,796,171		3,013,162
Total operating expenses	<u> </u>	4,128,427		3,895,902		12,236,905		14,354,338
LOSS FROM OPERATIONS		(2,859,109)		(2,856,644)		(6,782,469)		(9,717,457
Interest income		26,783		_		34,108		5,813
Transaction costs allocated to warrant liabilities		_		_		_		(391,335
Unrealized (loss) gain on change in fair value of warrants classified as a liability		(278,400)		1,758,200		334,700		2,540,700
Other (expense) income, net		(3,469)		(26,352)		6,396		(43,226
Loss before provision for income taxes		(3,114,195)		(1,124,796)		(6,407,265)		(7,605,505
Provision for income taxes								_
NET LOSS	<u>\$</u>	(3,114,195)	\$	(1,124,796)	\$	(6,407,265)	\$	(7,605,505
Less: Net loss attributable to noncontrolling interest		14,429		576		52,470		833
NET LOSS attributable to Applied DNA Sciences, Inc.	\$	(3,099,766)	\$	(1,124,220)	\$	(6,354,795)	\$	(7,604,672
Deemed dividend related to warrant modification				<u> </u>		<u> </u>		110,105
NET LOSS attributable to common stockholders	<u>\$</u>	(3,099,766)	\$	(1,124,220)	\$	(6,354,795)	\$	(7,714,777
Net loss per share attributable to common stockholders-basic and diluted	\$	(0.24)	\$	(0.13)	\$	(0.49)	\$	(0.94
Weighted average shares outstanding- basic and diluted		12,908,520		8,982,520		12,908,520		8,184,807

Comparative
Quarterly
Statement
of Operations
(unaudited)



Quarterly Consolidated EBITDA (unaudited)

Net Loss
Interest income
Depreciation and amortization
Change in provision for bad debt
Unrealized loss (gain) on change in fair value of Common Warrants
Stock based compensation expense
Total non-cash items
Consolidated Adjusted EBITDA (loss)

l Ended June 30,		
2022	2023	
(1,124,796)	\$ (3,114,195)	\$
-	(26,783)	
321,185	349,146	
-	34,169	
(1,758,200)	278,400	
272,914	340,042	
(1,164,101)	974,974	
(2,288,897)	\$ (2,139,221)	\$

	June 30,			September 30,	
		2023	2022		
ASSETS		(unaudited)			
Current assets:					
Cash and cash equivalents	\$	10,756,235	\$	15,215,28	
accounts receivable, net of allowance of \$75,000 and \$330,853 at June 30, 2023 and September 30, 2022, respectively		682,701		3,067,54	
eventories		276,422		602,24	
repaid expenses and other current assets		524,904		1,058,05	
otal current assets		12,240,262		19,943,12	
roperty and equipment, net		1,168,038		2,222,98	
Other assets:					
estricted cash		750,000		_	
apitalized transaction costs		275,726		_	
perating right of use asset		1,355,508		_	
pending right of the table.				98,99	
otal assets	\$	15,789,534	\$	22,265,11	
LIABILITIES AND EQUITY					
LIABILITIES AND EQUITI					
urrent liabilities:					
ccounts payable and accrued liabilities	\$	2,127,908	\$	3,621,73	
perating lease liability, current		487,425			
eferred revenue		275,885		563,53	
otal current liabilities		2,891,218		4,185,30	
ong term accrued liabilities		31,467		31,40	
perating lease liability, long term		868,081			
arrants classified as a liability		4,804,700		5,139,40	
otal liabilities		8,595,466		9,356,17	
		-,,,		. , ,	
ommitments and contingencies					
pplied DNA Sciences, Inc. stockholders' equity:					
referred stock, par value \$0.001 per share; 10,000,000 shares authorized; -0- shares issued and outstanding as of June 30 2023 and September 30, 2022, respectively		_		_	
eries A Preferred stock, par value \$0.001 per share; 10,000,000 shares authorized; -0- issued and outstanding as of June 30, 2023 and September 30, 2022, respectively		_		_	
eries B Preferred stock, par value \$0.001 per share; 10,000,000 shares authorized; -0- issued and outstanding as of June 30, 2023 and September 30, 2022, respectively					
		_			
ommon stock, par value \$0.001 per share; 200,000,000 shares authorized as of June 30, 2023 and September 30, 2022, 12,908,520 shares issued and outstanding as of ne 30, 2023 and September 30, 2022		12,909		12,9	
dditional paid in capital		306,091,402		305,399,00	
ccumulated deficit		(298,854,883)		(292,500,0	
pplied DNA Sciences, Inc. stockholders' equity		7,249,428		12,911,8	
oncontrolling interest		(55,360)		(2,8	
oncontrolling interest		7,194,068		12,908,9	
nui equity		7,134,000		12,900,9	
otal liabilities and equity	<u>\$</u>	15,789,534	<u>\$</u>	22,265,1	
22 Applied DNA Sciences Inc					

Quarterly Consolidated Balance Sheet



Einea IV/T

Better RNA...Faster



mRNA Manufacturing Pain Points

Bacterially derived plasmid DNA is currently the starting material for mRNA

Long lead times increase mRNA production timeline

Struggles with complex DNA sequences such as Poly(a) tails

Requires expensive enzymatic linearization and additional filtration steps

Increased regulatory scrutiny



Problematic inflammatory byproduct of conventional IVT

dsRNA removal is essential for safe and effective mRNA products

Currently removed via expensive and complex purification methods

Can cause quality control issues

Increased regulatory scrutiny





Next Generation mRNA Production

linearDNA[™] **IVT Templates**

- Cell free enzymatic production platform
- Rapid production timelines
- Eliminates the need for Plasmid linearization and purification
- Primer induced modifications enable simple 3' and 5' IVT template customization
- Stable homopolymer amplification

Linea[™] RNA Polymerase (RNAP)

- Reduces or eliminates dsRNA contamination
- mRNA yields that meet or exceed conventional IVT
- Proprietary and patent pending RNAP
- Includes DNA binding domain and high-fidelity T7
- High binding affinity for chemically modified linearDNA IVT templates



Significantly simplified mRNA production

Reduced or eliminated dsRNA contamination

Delivery of commercial scale IVT templates in as little as 14 days

Integrable into current mRNA workflows

better RNA...faster





Simplified mRNA Production



Linea[™] IVT mRNA Production

LinearDNA IVT Template Manufacture

In vitro transcription Purification step 1

Purification step 2

Encapsulate and sterile filtration

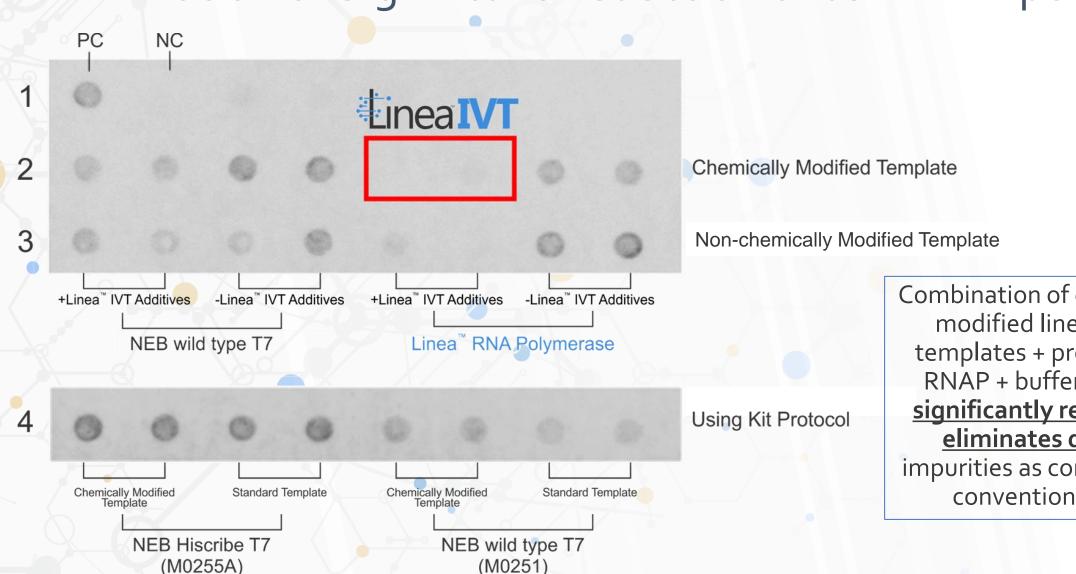
mRNA Product







Elimination or Significant Reduction of dsRNA Impurities



Combination of chemically modified linearDNA templates + proprietary RNAP + buffer system significantly reduces or eliminates dsRNA impurities as compared to conventional IVT

Revenue Growth Opportunities linear DNA in Diagnostics

Single Diagnostic Customer - Revenue Trend FY'23

