UNITED STATES SECURITIES AND EXCHANGE COMMISSION

Washington, D.C. 20549

FORM 8-K

CURRENT REPORT
Pursuant to Section 13 or 15(d)
of the Securities Exchange Act of 1934

Date of Report (Date of earliest event reported): December 11, 2020

DERMTECH, INC.

(Exact name of registrant as specified in its charter)

Delaware (State or other jurisdiction of incorporation) 001-38118 (Commission File Number) 84-2870849 (IRS Employer Identification No.)

	,	· · · · · · · · · · · · · · · · · · ·			
		11099 N. Torrey Pines Road, Suite 100 La Jolla, CA 92037 (Address of Principal Executive Offices and Zip Code)			
	Registr	rant's telephone number, including area code (858) 450-4222			
Chec	ck the appropriate box below if the Form 8-K filing is intended to simultaneously sati	isfy the filing obligation of the registrant under any of the following	ng provisions:		
	Written communications pursuant to Rule 425 under the Securities Act (17 CFR	230.425)			
	Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)				
	Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))				
	Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange	nge Act (17 CFR 240.13e-4(c))			
Secu	urities registered pursuant to Section 12(b) of the Act:				
	Title of each class	Trading Symbol(s)	Name of each exchange on which registered		
	Common Stock, par value \$0.0001 per share	DMTK	The Nasdaq Capital Market		
Indic chap	cate by check mark whether the registrant is an emerging growth company as defined oter).	in Rule 405 of the Securities Act of 1933 (§230.405 of this chapt	ter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§240.12b-2 of this		
Eme	erging growth company 🗵				
	n emerging growth company, indicate by check mark if the registrant has elected not to Exchange Act. $\;\Box$	o use the extended transition period for complying with any new o	or revised financial accounting standards provided pursuant to Section $13(a)$ of		

Item 7.01 Regulation FD Disclosure

The Company is furnishing with this Current Report on Form 8-K a copy of its current corporate presentation slides. The information in these slides shall not be deemed "filed" for purposes of Section 18 of the Securities Exchange Act of 1934, as amended, or the Exchange Act, or otherwise subject to the liabilities of that section, nor shall it be deemed incorporated by reference in any filing under the Exchange Act or the Securities Act of 1933, as amended, except as expressly set forth by specific reference in such a filing.

Item 9.01. Financial Statements and Exhibits.

(d) Exhibits

 Exhibit No.
 Description

 99.1
 Current Corporate Presentation

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

DERMTECH, INC.

By: /s/ Kevin Sun

Name: Kevin Sun Title: Chief Financial Officer

Date: December 11, 2020





Corporate Presentation

Q4 2020

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Disclaimer

This presentation includes forward-looking statements intended to qualify for the Safe Harbor from liability established by the Private Securities Litigation Reform Act of 1995. Forward-looking statements include all statements that are not historical facts and can be identified by terms such as "anticipates," "believes," "continues," "could," "seeks," "estimates," "targets," "guidance," "expects," "intends," "may," "ongoing," "plans," "potential," "predicts," "prospects," "projects," "should," "will," "would," or similar expressions intended to identify statements about the future and the negatives of those terms, although not all forward-looking statements contain these identifying words. These statements are based on management's current beliefs and expectations. These statements, including, but not limited to, statements regarding clinical utility tests, regulatory action, third-party payer reimbursement, and demand for our tests, are subject to substantial known and unknown risks, certainties, and other factors that could cause actual results to differ materially from those suggested or implied by these forward-looking statements. These factors include, but are not limited to, the following: regulatory action with respect to our existing and planned products (our "Products"); the commercial launch and future sales of our Products; our ability to achieve favorable pricing for our Products; third-party payor coverage and reimbursement for our Products and the impact of such coverage and reimbursement on our revenue growth; the accuracy of our estimates regarding anticipated capital requirements and our needs for additional financing; market size and market adoption of our Products by dermatologists and others; the timing, cost and other aspects of the commercial launch of our Products; the timing and cost of clinical utility tests for our Products, including whether such tests will be conducted at all; our ability to develop and commercialize additional tests and products; and our forward-looking statements.

The risks and uncertainties that may cause actual results to differ materially from our current expectations are more fully described in our reports filed with the Securities and Exchange Commission (the "SEC"). You may obtain these reports for free by visiting EDGAR on the SEC website at www.sec.gov. We assume no obligation to update any forward-looking statements after the date of this presentation or to conform any forward-looking statements to actual results, and has no intention of doing so except to the extent required by applicable law. You should, therefore, not rely on the forward-looking statements in this presentation as representing our views as of any date subsequent to the date of this presentation.

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Leadership Team



John Dobak, MD Chief Executive Officer

- Founder & Chairman, 10xBio (aesthetic medicine drugs)
- Chairman, Pantherics (anti-inflammatory drugs)
- MD, UCSD, Bachelors, UCLA



Todd Wood Chief Comm. Officer

- Allergan, VP US Sales, Dermatology, Ophthalmology, Aesthetics
 - - LabCorp



Daniel Visage, MBA Sr. VP, Payer Access

- Progenity, VP Payer Access
- OPKO/BRL, VP Managed Markets
- Florida Blue, Kaiser, Carecentrix



Claudia Ibarra Chief Operating Officer

- Exagen, Sr. VP Lab Operations
- Genoptix, Director Lab Operations
- Dexcom, VP Finance, Corporate Controller and Treasury, Interim CFO Biosite, FP&A, SEC Reporting, SOX Compliance
 - MBA, MS, Kelley School of Business at Indiana University

Chief Financial Officer



Kevin Sun, MS, MBA Zuxu Yao, PhD Chief Technology Officer



- · Post-doctoral, UCSD
- PhD, Memorial University of Newfoundland



Burkhard Jansen, MD Mike Howell, PhD Chief Medical Officer



· FDA Consultant Post-doctoral, University of Minnesota; MD, University of Graz



• Immunologist

Senior roles at Incyte, Medimmune

Inventor IL-36 receptor antibody, Spesolimab

Ph.D. West Virginia, thesis included use of tape stripping

MISSION: To transform dermatology with our non-invasive skin genomics platform, to democratize access to high quality dermatology care, and to improve the lives of millions

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DermTech, Inc. (NASDAQ: DMTK) is a leader in precision dermatology enabled by a non-invasive skin genomics platform



We address large market opportunities in skin disease with an initial focus in skin cancer



We are commercial stage and currently offer the Pigmented Lesion Assay (PLA) for early melanoma detection



We operate a **CLIA-certified and CAP-accredited commercial laboratory** in San Diego, CA. Our skin cancer product is available in all 50 U.S. states



We are commencing scale up around our recent Medicare coverage policy and new CPT Code (0089U) with favorable reimbursement

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Genomic Innovations Are Less Invasive, More Accurate, and Less Expensive

Focus	Company	Old Standard of Care	New-gen Diagnostic
Breast	Senomic Health	S. C.	CONSUMPLEX
Colorectal	exact sciences		Completed
Thyroid	veracyte.		1
Heart	CareDx*		■*AlloMap" ■*HeartCare
Lung	GUARDANT HEALTH	Banan and Banan	GUARDANT 366
Prenatal	% natera		panorama° natera prenatal screen
Skin	Derm Tech		Derm Tech

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PLA Brings Skin Cancer Diagnostics into the 21st Century

A Better Diagnostic Solution for Dermatology

Current Practice: More Art Than Science



Subjective	Low Accuracy
Invasive	High Cost

The Future: Non-invasive Skin Genomics

Adhesive Patch Sample Collection

Genomic Analysis CLIA/CAP Commercial Lab

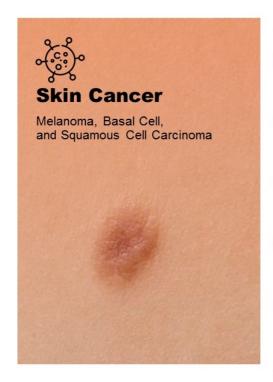
Physician Report (within 72 hrs.)

Objective	High Accuracy
Non-invasive	Low Cost

CAP: College of American Pathologists; CLIA: Clinical Laboratory Improvement Amendments.

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A Snapshot of the U.S. Skin Cancer Market



More people are diagnosed with skin cancer than all other cancers combined and 1 in 5 Americans will develop skin cancer by the age of 70. More than 50 MM patients have UV related skin damage that puts them at risk for skin cancer



Annual cost of treatment is estimated to be \$8.1 billion, \$4.8 billion for non-melanoma and \$3.3 billion for melanoma (2018)



~4.5 million cases of basal cell and squamous cell diagnosed per year in the US, ~11 MM diagnostic biopsies, with ~20,000 deaths



~180,000 new cases of melanoma were reported in 2018, ~4.0 MM diagnostic biopsies, with ~10,000 deaths



~3,000 new cases cutaneous lymphoma diagnosed, challenging diagnosis with average 3-6 years to definitive diagnosis, confusion with dermatitis leads to significant number of surgical biopsies (est. 420,000 annually)

Source: Cancer Facts & Figures 2018. American Cancer Society.

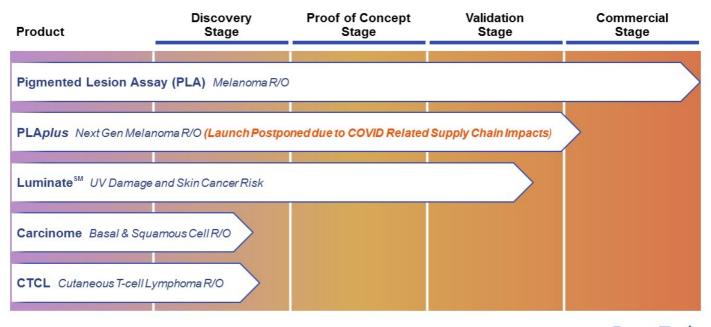
Source: AAD Skin Cancer Types Facts, aad.org

Source: Cutaneous Lymphoma Foundation, olfoundation.org

DermTech

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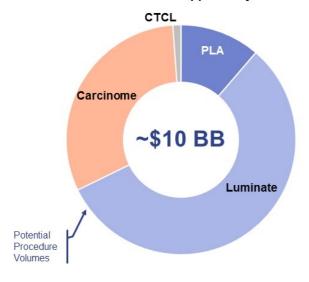
DERMTECH's Skin Cancer Product Pipeline



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Skin Cancer Market Opportunity

Estimated Total Potential Procedure Volumes and Market Opportunity



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Potential U.S. Patient and Procedure Volumes

- PLA/PLAp/usSM: melanoma¹ skin cancer
 - 4-4.5 MM diagnostic surgical biopsies
- Carcinome: non-melanoma skin cancer¹
 - 11-12 MM diagnostic surgical biopsies
- LuminateSM: UV damage and skin cancer risk²
 - 82 MM U.S. Adults aged 35-54, ~1/3 actinic sun damage
- CTCL: Cutaneous lymphoma³
 - Estimated biopsies for inflammatory conditions that my be confused with CTCL ~400K-700K

Source: IQVIA Skin Cancer Claims and Procedures Study
 Source: Kaiser Family Foundation, kkf. org and DermTech Estimate
 Source: DermTech KOL advisor estimates



Early Detection of Melanoma Is Critical to Survival

5-Year Relative Survival Rate1



When Diagnosed at the Local Stage

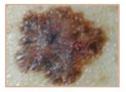


When Diagnosed at the Regional Stage



When Diagnosed at the **Distant** Stage

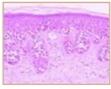
Early Detection Visually Is Extremely Difficult



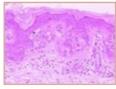




Skin Cancer







Skin Cancer

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Cancer Facts & Figures 2018. American Cancer Society.
 Hanke CW, Carcinoma and Keratoses, 1:2 2019

Unmet Need: Melanoma's Challenging Diagnostic Pathway

Diagnostic Inaccuracy and Unnecessary Surgery

Subjective Clinical Visual Assessment

Leads to ~25 surgical biopsies performed per melanoma found1

Subjective Pathology Assessment

Assess only 1% of the Lesion

58% of early-stage melanomas have diagnostic misinterpretation²

Leads to high proportion (>20%) unnecessary wide excisions³

Anderson A, et al., JAMA Dematology, 2018; doi:10.1001/jamadermatol.2018.0212
 Elmore JG, et al. BMJ. 2017;357;2813.
 Strazzula L, et al. Journal of the American Academy of Dematology, 2014; 71:1071-

s. ican Academy of Dermatology, 2014; 71:1071-8

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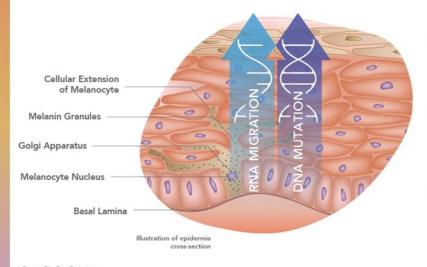
Most Biopsies Are NOT Melanoma

~4.0-4.5 million biopsies annually to identify 180k cases

Multiple biopsies and excisions of benign pigmented lesions. Courtesy of G. Peck, MD.



The First Non-invasive Gene Expression and Mutation Test to **Enhance Melanoma Detection**



Non-invasive Adhesive Patches

- Avoids unnecessary surgery^{1,2}
- · Improves patient care and comfort

Earliest Detection

- Detects genomic drivers of cancer^{1,3,4}
- Genomic changes can precede visual changes^{1,3,4}

Highly Accurate

- Reduces probability of missing melanoma (<1%)
- 100% of a lesion is sampled
- · Uses the precision of genomics

12 DermTech

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Better Care at a Lower Cost

Highly Validated Diagnostic Metrics



Enhance early detection of melanoma by identifying lesions with genomic atypia

- Elmore JG, et al. BMJ. 2017; 357;[2813 & Malvehy J, et al. BJD, 2014; 177:1099-1107
 Ferris L, et al. JAMA Dematol. 2018; 154(10):1229-1220 & Gerami P, et al., JAAD 2017, 76:114-120e
 Strazzula L, et al. JAAD 2017, 71:1071-8, Ferris L, et al. Melanoma Research, 2018 1:D0110.1097
 Homberger J and Siegel D. JAMA Dermatol, 2018 154(9):1-3
 Ferris L, et al. DOJ, 2019, 25(6):1-8
 Ferris LK, Moy RL, Gerami P et al. J Invest Dematol. 2019;139(5):1127-1134

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Superior Performance

Performance Metric	Biopsy & Histology	PLA Test	Improvement
Probability of Missed Melanoma ¹	17%	1%	17x
Biopsies per Melanoma ²	25	2.7	10x
Wide Excisions per Melanoma ^{2,3}	5.2	1.6	3x
Cost per Lesion Tested ⁴	~\$1,000	\$760	\$240

Strong Clinical Validation and Robust IP Portfolio

Milestone	Status	Sample Size
Analytical Validation	✓ Complete	125
Clinical Validation – pathology	✓ Complete	555
Clinical Validation – mutation	✓ Complete	626
Clinical Utility	✓ Complete	45 Derms
Real-World Utility	√ Complete	381
1-Year Follow-up	✓ Complete	734
Real-World Utility Registry	✓ Complete	3418
Adhesive Biopsy Validation	√ Complete	N/A
Health Economic	✓ Complete	326
CPTCodes	✓ Complete	N/A

IP Portfolio



Issued US patents provide broad protection for melanoma product through 2030

- Broad methods covering RNA analysis of adhesive patch collected skin: 7,183,057
- Method of detection of biological factors in epidermis: 6,720,145
- ullet Broad claims for melanoma gene classifier: 9,057,109
- · Broad claims for melanoma gene classifier: 10,407,729

CPT: Current Procedural Terminology; N/A: not applicable.
Publications available at www.dermtech.com

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Patents issued in select countries worldwide

· Multiple European countries, Canada, Japan, and Australia



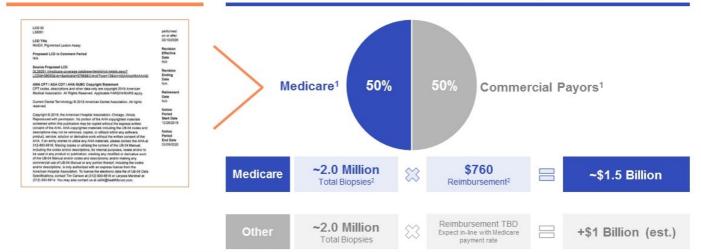
Trade secrets and technical know-how

- Low quantity, poor quality sample material requires special processes
- · Custom automation

CMS Pricing in Place Unlocks Significant Market Opportunity

Medicare Coverage Established in January 2020

Estimated Total Number U.S. Biopsies for Pigmented Lesions¹



Medicare Market Alone Has Potential for >\$1 Billion in Sales

Source: IQVIA Skin Cancer Claims and Procedures Study
 Clinical Laboratory Fee Schedule: https://www.cms.gov/medicaremedicare-fee-service-paymentdinicallabfeeschedclinical-laboratory-fee-schedule-files/20clabr

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DermTech's Goal: Eliminate Melanoma Deaths in the US

There are 5 Requirements to Eradicate a Disease

- Diagnostic test with 99% NPV & is agreeable with patients
- 2. Treatment protocol with 99% efficacy
- 3. Payer model to fund effective diagnosis & treatment
- 4. Widespread HCP adoption of diagnostic pathway & treatment protocol
- Activated consumers willing to be evaluated

Current Pathway Does Not Meet Requirements

- Invasive biopsy & pathology with 83% NPV & unpopular with patients
- √ 99% 5-year survival rate when detected early
- √ Biopsy & pathology widely covered
- · Diagnosis restricted to dermatology
- Very few patients visit dermatology



PLA Case Study – JAMA Dermatology Publication

Effectiveness Attributes of the PLA Test

Background

- 28-year-old female
- Family history of melanoma
- · Complained of bug bite
- · Refused surgical biopsy

DermTech Results

- · PLA test was positive
- Histopathology revealed 0.5-mm melanoma
- Curative wide excision performed
- Case study published in JAMA Dermatology¹



Childs MV. JAMA Dematol, 2018, 154(2):223
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Commercial Activities



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Commercial Channels

Dermatology Professional Channel



Direct sales reps



Inside sales reps



Regional managers



Medical science liaisons



Digital Consumer Channel



Social Media



Paid Search



Programmatic Display



Future Care Delivery Channel



Telemedicine



On- and near-site employer health clinics & retail clinics



Integrated Primary Care Networks





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Dermatology Professional Sales Force

- · 45-50 sales reps give coverage of 13,000 dermatology clinical professionals
 - ~80% sales manger hiring complete
- ~5,000 prioritized targets
 - Average ~66 biopsies/mo/clinician
 - Drive to average of 10 PLA tests/mo/clinician2
 - · Based on historical physician productivity metrics
 - · Minimal impact to biopsy practice
- ~10% served market penetration = ~446,000 tests per year
- Revenue potential @ 10% penetration = ~\$312 MM/year3

Source: IQVIA Skin Cancer Claims and Procedures Study
 DermTech estimate based on historical "good" account
 Assumes convergence of Medicare and commercial payer rates with average ASP of \$700/test

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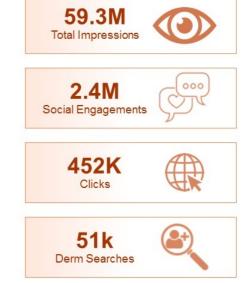
Each circle represents a prioritized dermatology target: high number melanoma biopsies, >50% proportion Medicare patients



Digital Consumer Channel is Driving Awareness and Patients to Take Action







Source: Socialbakers; Google Analytics; Acuity, May-Oct, 2020, select U.S. regions Data: Includes metrics across both published (organic and boosted) and unpublished content on owned social platform

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Future Cancer Care Dx Delivered via Telemedicine Channel

Telemedicine Technology Solution



- Store & Forward iPhone and Android App
- Secure email and text
- HCP network of users
- Remote interactive access

Patient Home Collection Kit



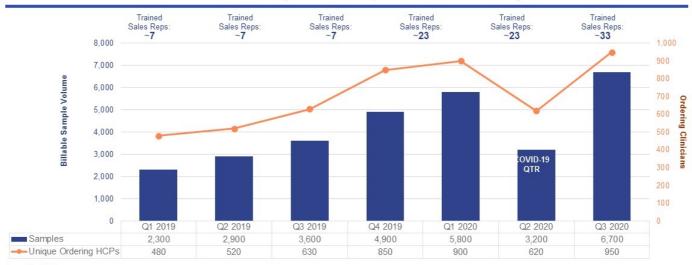
- · Sent to patient's home
- Supported by easy to use instructions
- QR Code activates instructional video

Potential Partnerships With Existing Real-Time Interactive Providers

- Employer service providers
- · Benefit providers
- · Integrated networks
- Other Telemedicine channels including skin disease

Ordering Clinician Installed Based and Sample Volume Resumed Growth Post Q2 COVID Shutdown

DermTech Historical Quarterly Billable Sample Volume and Ordering Clinicians



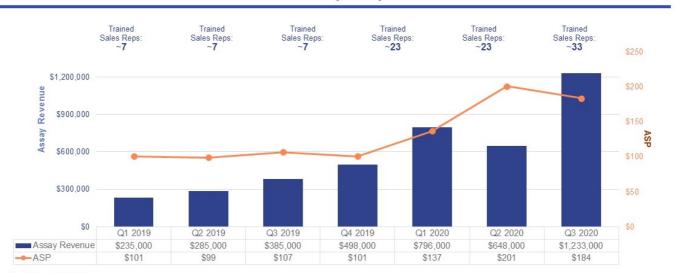
Source: DermTech, Inc.
Training program for new sales reps is 3 months

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Beginning to Monetize Sample Volume With Recent Medicare Coverage

Coverage Commercial Payer Coverage Expected To Further Accelerate Revenue Growth*

DermTech Historical Quarterly Assay Revenue and ASP



Source: DermTech, Inc., Capital IQ, Bloomberg *Recently contracted with BCBS of IL and BS of CA with early 2021 effective dates.

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Impact Discussion

- We are deemed an essential business and our commercial laboratory remains fully operational
- Dermatology offices have not yet re-opened to full capacity and future operations are dependent upon regional COVID-19 infection rates and regional restrictions
- We have been conducting virtual sales calls, providing client education, and introduced a telemedicine option for remote, clinician-guided sample collection
- We completed approximately 80% of our sales force expansion for the year by July 2020 and expect to finish 2020 with ~40 sales representatives
- We cannot predict which states will remain open, but we did recently see some encouraging data which underscores clinician acceptance of the DermTech PLA including:
 - Sample volume recovery in October 2020 to all time high levels
 - Sample volume for September and October 2020 increased 34%, compared to January and February 2020
 - In October 2020, we had an all-time high in the number of ordering clinicians who order 10 or more tests per month.

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Back Up



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THE PLAplus: Optimized Gene Targets

Increased Sensitivity to 97%

LINC00518^{1,2}

Long Intergenic Non-Coding RNA 518

- Novel marker discovered by DermTech
- · Member of a rapidly growing family of regulatory RNA molecules
- Important regulator of oncogenesis play role in melanoma proliferation and invasion
- · Overexpressed in melanoma

PRAME^{3,4}

Preferentially Expressed Antigen in Melanoma

- · Well described target involved in many tumors, including in melanoma progression and metastasis
- Independently validated- Haqq, Myriad, Castle
- Promotes tumor progression by interfering with retinoic acid receptor signaling
- · Overexpressed in melanoma

TERT^{5,6}

Telomerase Reverse Transcriptase Promoter

- Involved in many tumors, is found in 70-80% of melanoma
- Mutations lead to oncogenesis through functional increases in TERT protein, telomerase activity, telomere length, cell immortalization and proliferation
- · Associated with histopathologic features of aggressiveness and poor survival in melanoma

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^{1.} Lessard L, Liu M, Marzese et al. J Invest Dermatol. 2015;135(10):2464-74.

2. Sarkar D, Leung EY, Baguley BC, Finlay GJ, Askarian-Amiri ME. Epigenetics. 2015;10(2):103-21.

3. Ikeda H, Lethé B, Lehman F et al. Immunity. 1997;(2):193-203.

4. Hermes N, Kewitz S, Staege MS. Curr Cancer Drug Targets. 2016;16(5):400-414.

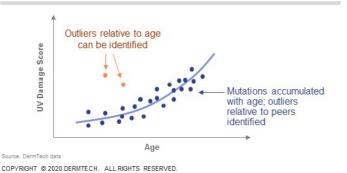
5. Ferris LK, Moy RL. Gerami P et al. J Invest Dematol 2019;139(5):1127-1134.

6. Nagore E, Rachakonda S, Kumar R. Oncottarget. 2019;10(16):1464-1549.

Consumer Health Offering

Mutation Burden Increases with Sun Exposure







- Quantifies UV damage to the genome in normal appearing skin by assessing driver mutations for basal cell and squamous cell cancer
 - Identifies risk for skin cancer and premature photoaging
- · Generates UV Damage score
 - Non-sun exposed areas of skin from the buttocks have zero score
 - Patients with a history of skin cancer have highest score
- Informs patients about actions and treatments that can be undertaken to repair damage and improve skin health

Research Partnerships and Collaborations

Indication*	Pilot	Phase 1	Phase 2	Phase 3
Pemphigus Vulgaris				
Hidradenitis Suppurative				
Psoriasis				
Mantle Cell Lymphoma				
Follicular Lymphoma				
Lupus				
Vitiligo				
Atopic Dermatitis				
Chronic Spontaneous Urticaria				

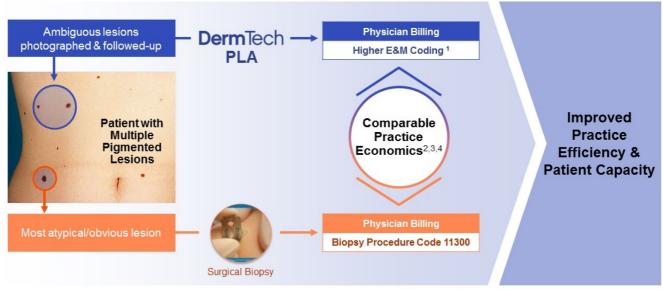
- ~\$12 million in research programs booked to date
- Robust pipeline of opportunities and pending contracts
- · Independent performance development
- Expansion to late-stage trials (phase II, III)



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Define PLA Positioning and Clarify Practice Economics

The PLA Provides Comparable Economics With Better Patient Care

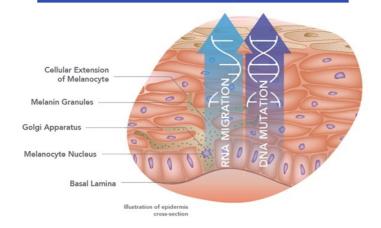


Based on AMA billing criteria for CPT E&M codes
 Net economics accounting for differences in cost for labor (physician work vs. nursing work) and supplies
 Sliopsy procedure costs based on AMA RVS Update Committee figures
 Reimbursement rates based on Medicare payments for established patient E&M CPT codes and biopsy procedure CPT codes, CodeMapTM

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Precision Dermatology Delivers More Accurate Melanoma Detection

RNA and DNA collected by the PLA/PLAplus spans across the entire lesion1,2



Significant improvement against current standard of care

PLA

Dermatopathology

Samples 100% of lesion

Samples less than 1%-2% of a potentially partial surgical biopsy

Provides additional objective genomic information

Relies on subjective visual criteria

Measures malignant changes that cannot be seen visually

Morphologic changes must be present and in the field of observation

Ferris LK, et al. JAMA Dermatol. 2017;153(7):675-680.
 Data on File. DermTech, Inc.

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