

Pulse Biosciences

(Nasdaq: PLSE)

November 2020

Forward-looking statements

This presentation by Pulse Biosciences, Inc., contain estimates and forward-looking statements as of November 9, 2020 including, among others, statements regarding Pulse Biosciences' future business plans, products, commercial applications, intellectual property strategy, clinical trials, regulatory processes and pathways, and markets for its technologies, as well as patient and physician behavior and preferences and other future events.

You should not place undue reliance on forward-looking statements, as they involve known and unknown risks and uncertainties that are, in some cases, beyond the Company's control and could cause actual results to differ materially from the information expressed or implied. Factors that could materially affect actual results are described in detail in the Company's recent Securities and Exchange Commission filings.

Pulse Biosciences undertakes no obligation to revise or update forward-looking statements to reflect future events or circumstances.



Provide superior treatment for a wide range of dermatological conditions through a novel, proprietary, tissue sparing solution





Nano-Pulse Stimulation Technology

Non-thermal modality that delivers nanosecond duration pulses of electrical energy

High speed nanosecond energy pulses penetrate the cell membrane and disrupt internal cellular function, leading to regulated cell death

Unlike thermal (heat/cold) technologies, NPS directly impacts cellular structures while sparing non cellular tissue





Demonstrated Mechanism of NPS in Skin

Safe, precise targeting and elimination of cells while sparing the dermis



Healthy Skin





Seven (7) days post-treatment

- Healthy epidermal cells with dark nuclei
- Cells in treated epidermis are nonviable (ghost cells)
- Cell membranes and surrounding non-cellular tissue are intact
- Healthy epidermis emerges below
- Treated epidermal layer peels away



As of October 2020

106 issued patents globally owned & licensed

Additional 102 patent applications pending worldwide

Robust IP Portfolio

Multipronged Patent Strategy

- Pioneering IP for the use of nanosecond pulses in biology
- Covering methods and tools for the application of nanosecond pulses in biology
- Pioneering IP and continued development of IP with focus on skin-based applications
- Continued development and patent filings covering systems, applications, and methods of combining nanosecond pulsing with other biological technologies and agents



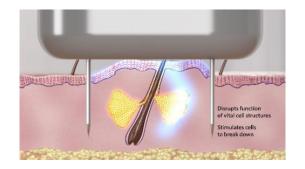
NPS Technology Delivery

Through the CellFX System proprietary platform



Broadly applicable platform with the ability to address multiple treatment targets in commercially significant areas of unmet clinical need







Excellent Clinical Data

Safety and efficacy across multiple difficult-to-treat dermatologic conditions

SK

Cleared Raised SK at 106-day follow-up



Seborrheic Keratosis

- 82% efficacy rate
- Clear or mostly clear with single treatment
- 78% satisfaction rate



SH

Cleared SH at 60-day follow-up



Sebaceous Hyperplasia

- 99.5% efficacy rate
- Clear or mostly clear 90% with single treatment
- 77% satisfaction rate

W

Cleared Wart at 60-day follow-up



Warts (non-genital)

- 79% rate of complete clearance in Recalcitrant Warts
- 71% of completely cleared warts achieved with 1-2 treatment sessions
- Previous modalities failed for majority of patients

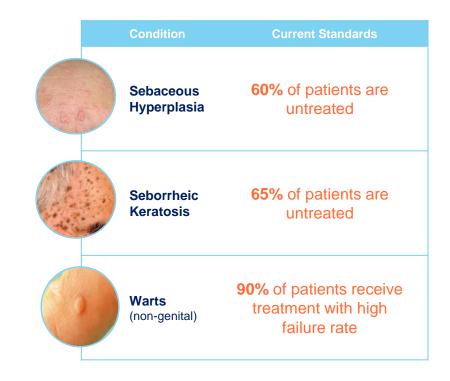
Unmet Need in Aesthetic Dermatology

Current treatments lack acceptable cosmetic outcomes or no standard of care exists



Aesthetic Patient Veterans

- Visit skin specialists for cosmetic procedures several times per year
- Seek latest treatment options
- Willingness to invest in procedures to look their best
- Aesthetic procedures are generally elective and cash-pay



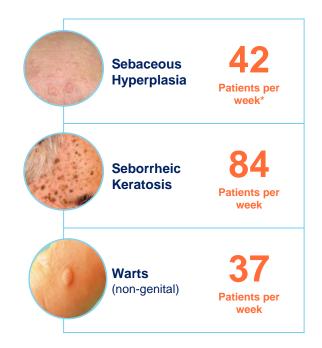


Market Opportunity

Widespread prevalence among patients visiting aesthetic dermatologists today











Commercialization Strategy

Innovative utilization based business model aligns incentives



Dermatologists want to treat all patients regardless of condition size/type/amount and want their costs aligned with what they treat



CellFX System is designed to support a business model based on each lesion treated, aligning the patient, physician, and Pulse



Practice Economics

MD Franchise Based on Expanding Utilization Over Time

Compelling Short-term Economics Long-term Practice Franchise Coultex Revenue to Initial **Three** < Four Avg. Ten **Avg. Five Lesions Practice** Patients / Week per Patient Investment **Month** Patients / Week ~\$500.000 / MD Return on • 10 x 15-minute treatment Net MD Rev \$45,000 Avg. Year session (2.5 hrs) Lesions Revenue Investment per lesion **Gross Profit** · Benign Lesion, SH, SK, \$150-\$200 per Patient per Patient & Warts to Practice · Room for growth with ~\$400,000 / 5 / patient \$900-\$1,200* additional apps Year



CellFX CloudConnect Unique competitive advantage Cell FX CloudConnect Software/ Utilization Customer **Practice** Customer Data & Management Web **Metrics** Application Updates & Cycle Unit Portal Maintenance Management



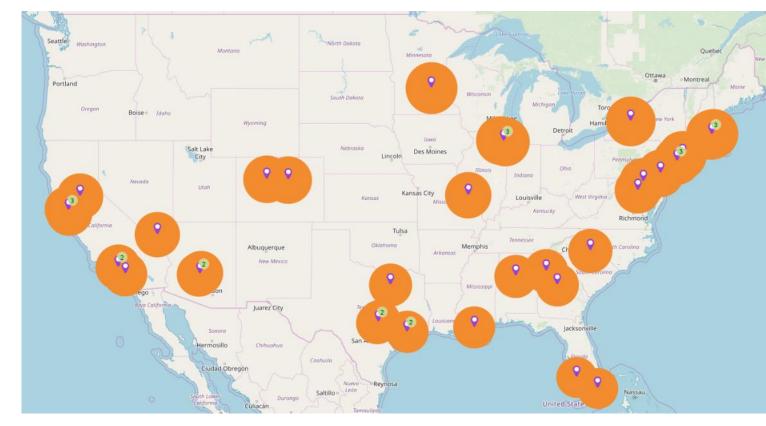
U.S. Target Market

80% of target early adopter physicians

located within 100-mile radius of the 40 controlled launch participants selected for controlled launch

4.6 M average population located within 100-mile radius of the 40 controlled launch participants selected

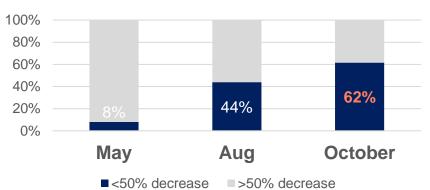
for controlled launch





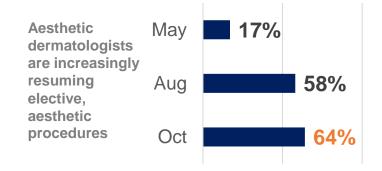
October survey of 39 aesthetic dermatologists on COVID-19 Impact shows positive signs of recovery in dermatology clinic activity

Aesthetic procedure volumes trending towards normal

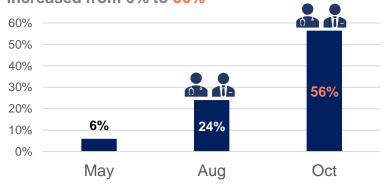


Majority believe procedure volumes will take another 6 months to return to normal volumes

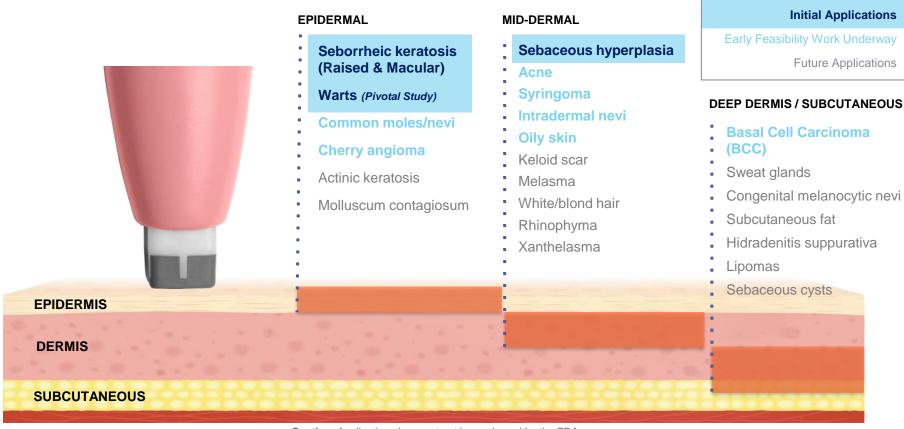




In-person interaction with industry reps increased from 6% to 56%



Extensive CellFX Application Pipeline



CellFX System Regulatory Strategy

Potential 510(k) clearance, CE mark and Health Canada approval by end of 1Q21







- Submitted 510(k) application for an initial FDA clearance with a general dermatologic indication in 4Q20.
- Received FDA IDE Approval and fully enrolled 60 patient comparative study to generate data for a subsequent 510(k) submission for a specific indication for the treatment of Sebaceous Hyperplasia, submission planned for 1Q21.
- Planned FDA IDE submission for common non-genital warts in 4Q20.
- Submitted Technical File in pursuit of the CE mark in 3Q20
 the regulatory approval required for commercialization in the EU.
- Submitted a Medical Device License application to Health Canada in 4Q20.



Strong KOL Support and Advocacy

Deep network of leading aesthetic dermatologists

PUBLICATIONS ON NPS TECHNOLOGY









PODIUM PRESENTATIONS

4 2018

12

2019

16

2020

(as of October)

2021 ABSTRACTS SUBMITTED

3

ASLMS



Scientific Advisors







- Dr. Brian Biesman, Nashville, TN
- Dr. Eric Bernstein, Ardmore, PA
- Dr. Chris Harmon, Birmingham, AL
- Dr. Suzanne Kilmer, Sacramento, CA
- Dr. Darius Mehregan, Ann Arbor, MI
- Dr. Gilly Munavalli, Charlotte, NC
- Dr. Tom Rohrer, Chestnut Hill, MA
- Dr. Vic Ross, San Diego, CA
- Dr. Elizabeth Tanzi, Chevy Chase, MD
- Dr. Brian Zelickson, Edina, MN
- Dr. Brian Berman, Aventura, FL
- Dr. Mark Nestor, Aventura, FL
- Dr. James Newman, San Mateo, CA
- Dr. Robert Pierce, Seattle, WA

Leadership Team

Proven and experienced in growing medical technology from clinical to commercial stage

Management

Darrin Uecker	President & Chief Executive Officer	gynesUnics computermotion.	
Edward Ebbers	EVP & GM, Dermatology	ZELTIQ thermage	
Sandra Gardiner	EVP & Chief Financial Officer	cutera tria.	

Board of Directors





Upcoming Milestones

- Regulatory Steps for the CellFX System
 - Potential receipt of initial FDA 510(k) clearance for general dermatologic indication as early as 1Q21
 - Subsequent 510(k) submission for SH indication planned for 1Q21
 - Potential receipt of Health Canada Medical Device License as early as 1Q21
- European controlled launch potentially as early as 1Q21
- Progress Clinical and Scientific programs
 - Continued presence at scientific meetings
- Continue to Expand Pipeline of Clinical Applications





Cash Utilization

2019 \$34.2MM 1Q 2020 \$9.5MM 2Q 2020 \$7.9MM 3Q 2020 \$8.2MM

- As of 09/30/20 cash, cash equivalents and investments were \$29.6MM
- Closed over-oversubscribed Rights
 Offering in June 2020 with \$29.5MM in
 net proceeds
- Additional gross proceeds of \$4.5MM through the exercise of issued warrants
- No debt



Financial Snapshot

	September 30, 2020		
	(In th	(In thousands)	
Cash, cash equivalents & investments	\$	29,586	
Prepaids & other assets		1,113	
Property, plant and equipment		2,562	
Goodwill & intangibles		6,839	
Right of use assets		9,595	
Total assets	\$	49,695	
Accounts payable & accrued liabilities Lease liability	\$	5,543 11,448	
Debt			
Stockholders' equity		32,704	
Total liabilities & stockholders' equity	\$	49,695	
Shares outstanding		25,342	

Thank You

