

Pulse Biosciences

(Nasdaq: PLSE)

May 2021

Forward-looking statements

This presentation by Pulse Biosciences, Inc., contain estimates and forward-looking statements as of May 12, 2021 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.

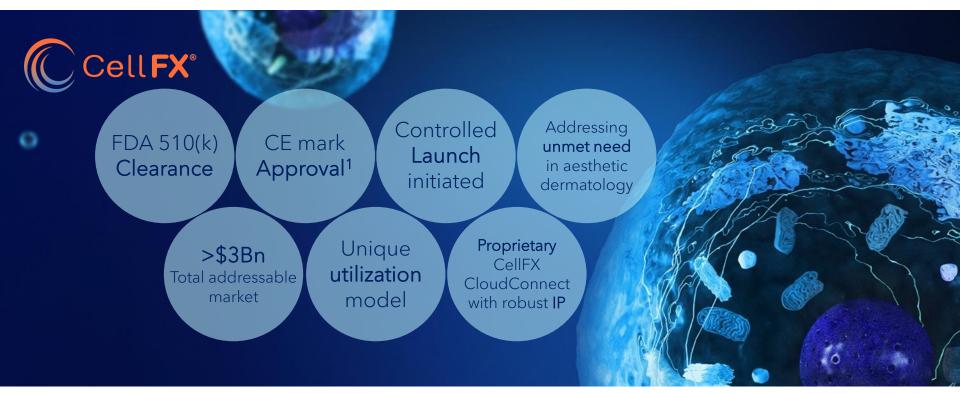


Provide superior treatment for a wide range of dermatological conditions through a novel, proprietary, tissue sparing solution





Novel Bioelectric Medicine Platform





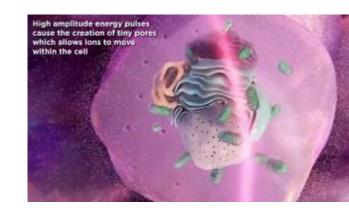
¹ Health Canada submission pending

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 (primarily collagen)





Demonstrated Mechanism of NPS in Skin

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



Healthy Skin



One (1) day post-treatment



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



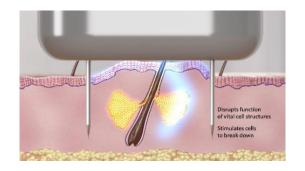
NPS Technology Delivery

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







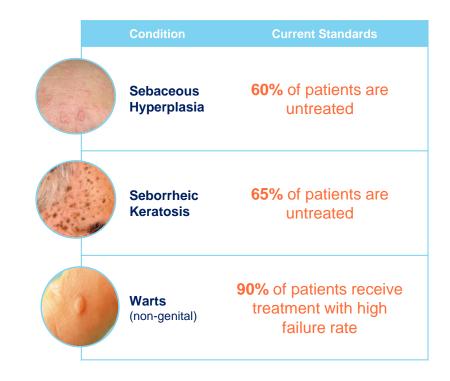
Unmet Need in Aesthetic Dermatology

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

6,800 **Aesthetic Physicians**









Surveys conducted by third-party market research firm on behalf of Pulse Biosciences, Inc., Full survey results available upon request 1. Based on 2019 Clinician (n=304) in the U.S.

- 2. Based on 2020 Clinician (n=46) in Europe

Excellent Clinical Data

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

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

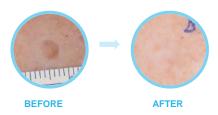


Warts (non-genital)

- Warts achieved complete clearance at a 79% rate, including recalcitrant warts
- 71% of completely cleared warts achieved with 1-2 treatment sessions
- Previous modalities failed for majority of patients

SK

Cleared Raised SK at 106-day follow-up



Seborrheic Keratosis

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

Strong KOL Support and Advocacy

Deep network of leading aesthetic dermatologists

Scientific Advisors







4

CLINICAL PUBLICATIONS ON NPS TECHNOLOGY







PODIUM PRESENTATIONS

4 2018

12

2019

16

2020

ACCEPTED ABSTRACTS

2021 (Apr/May)

Dr. Brian Biesman, Nashville, TN

Dr. Joel Cohen, Denver, CO

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, Boston, MA



CellFX Controlled Launch

A comprehensive and disciplined approach





KOL Controlled Launch Clinics across the U.S. and E.U.



KOLs experiences establish best practices for implementation of the CellFX System





Early adopters will be guided by KOL best practices

Track record for both clinical expertise and commercial acumen for introducing new technologies into a market

Shared experiences among peers at high visibility medical conferences and media outlets



Cycles Enable Treatment Delivery

Innovative utilization based business model aligns incentives



Patients want to pay based on number of lesions treated



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



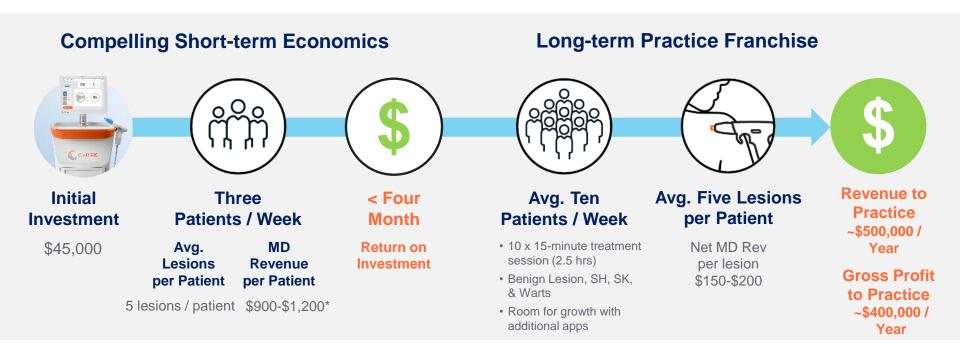


CellFX CloudConnect Unique competitive advantage Cell FX CloudConnect Software/ Utilization Customer **Practice** Customer Data & Management Web **Metrics** Application Updates & Cycle Unit Portal Maintenance Management



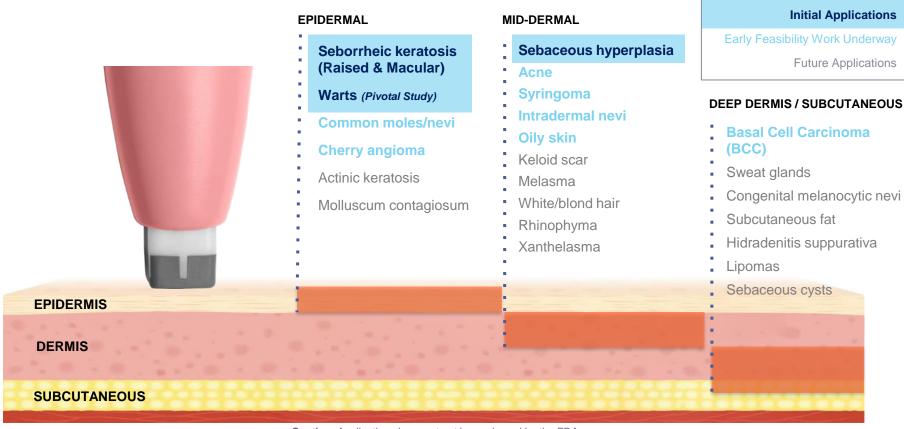
Practice Economics

MD Franchise Based on Expanding Utilization Over Time





Extensive CellFX Application Pipeline



CellFX System Regulatory Strategy

Expanding indications in US







- Received 510(k) Clearance for general dermatologic indication February 2, 2021
 - Progressed stepwise FDA strategy in pursuit of subsequent specific indications
 - Completed treatments in FDA IDE approved study to generate data for a 510(k) submission for the treatment of Sebaceous Hyperplasia, submission planned for 2Q21
 - **Completed 51 of 150 treatments** in FDA IDE approved study to generate data for a 510(k) submission for the treatment of common non-genital warts, submission planned for as early as 4Q21
 - Received FDA IDE approval for a treat and resect basal cell carcinoma feasibility study, enrollment of 30 patients expected to be completed by the end of 3Q21
- CE mark approval received January 22, 2021
- Health Canada Medical Device License application under review



Upcoming Milestones

- Onboarding of 75 total controlled launch participant clinics will continue throughout Q2 and conclude in Q3 of 2021
- Clinical and regulatory activity to expand of CellFX System clinical applications
 - Potential receipt of Health Canada Medical Device License as early as 2Q21
 - Subsequent 510(k) submissions
 - SH planned for 2Q21
 - Non-genital wart planned for as early as 4Q21
 - Enrollment of FDA IDE approved BCC feasibility study expected to begin in Q2 and end in Q3
- Progress Clinical and Scientific programs
 - Continued presence at scientific meetings





As of January 2021

112 issued patents globally owned & licensed

Additional 96 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



Cash Utilization

2020

\$34.6MM

1Q 2020

\$10.7MM

- As of 3/31/2021 cash, cash equivalents and investments were \$59.9MM
- February 2021 At-the-Market Equity Offering of up to \$60MM

Financial Snapshot

	March 31, 2021
	(In thousands)
Cash, cash equivalents & investments	\$ 59,883
Prepaids & other assets	2,458
Property, plant and equipment	2,430
Goodwill & intangibles	6,506
Right of use assets	 9,280
Total assets	\$ 80,557
Accounts payable & accrued liabilities	\$ 7,781
Lease liability	11,261
Debt	41,118
Stockholders' equity	 20,397
Total liabilities & stockholders' equity	\$ 80,557
Shares outstanding (as of 4/30/2021)	26,460



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





Thank You

