Pulse Biosciences

(Nasdaq: PLSE)

November 2021



Forward-looking statements

This presentation by Pulse Biosciences, Inc., contain estimates and forward-looking statements as of November 17, 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 conditions through a novel, proprietary, tissue sparing solution



Launching a New Category in Dermatology

Cell**FX**®

TIME TO GO

Say goodbye to unsightly cellular lesions with an innovative energy-based procedure







Raised, facial bumps sc

Dark, Rough, scaly spots persistent growths NPS Platform technology addresses unmet need aesthetically pleasing clearance of common spots, bumps and growths (benign skin lesions)

Built-in demand patients already visiting target aesthetic dermatologists and skin specialists

Positioned to be one of the strongest revenue and profitability categories for aesthetic dermatology clinics



Broadly Applicable Novel Bioelectric Medicine Platform





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

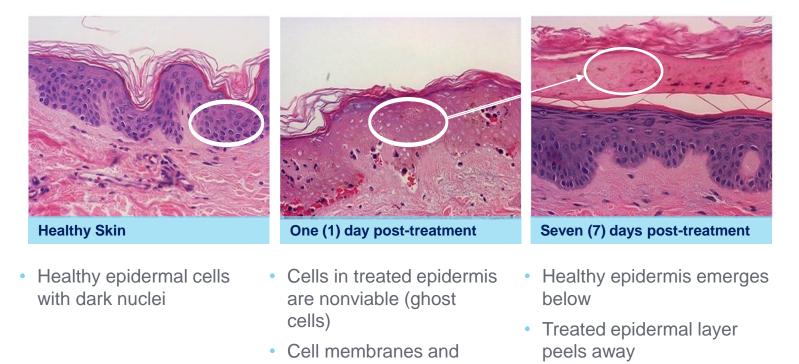
Unlike thermal (heat/cold) technologies, NPS directly impacts cellular structures while sparing non cellular tissue (primarily collagen)





Demonstrated Mechanism of NPS in Skin

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



surrounding non-cellular

tissue are intact



NPS Technology Platform

NPS delivered through the proprietary CellFX System



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







Initial applications in dermatology



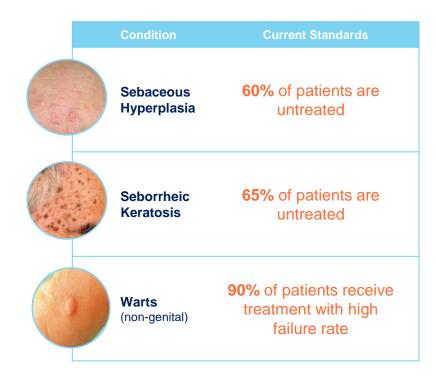
Unmet Need in Aesthetic Dermatology

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



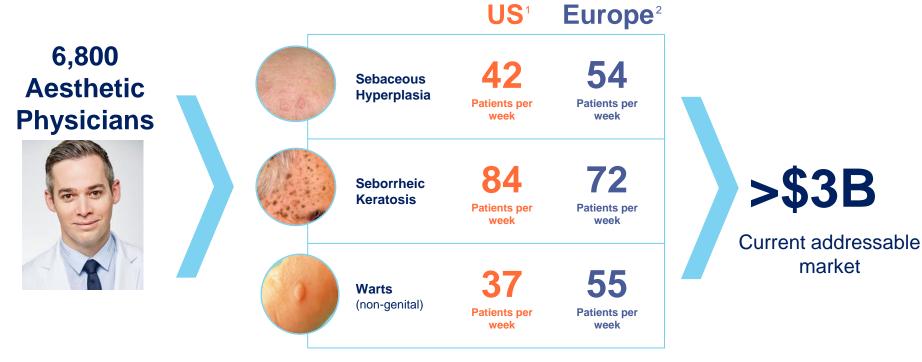
Aesthetic Patient Veterans

- 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





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

Published Peer-Reviewed Clinical Data

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

SH

Cleared SH at 60-day follow-up



Sebaceous Hyperplasia¹

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

Cleared Wart at 60-day follow-up



Warts (non-genital)²

- Up to 75% clearance efficacy, with majority of common warts cleared with a single procedure
- 48% of treated warts had failed previous modalities
- No evidence of recurrence within a 120-day observation period

SK

Cleared Raised SK at 106-day follow-up



Seborrheic Keratosis³

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



Munavalli G, Zelickson B, Selim M, Kilmer S, Rohrer T, Newman J, et al. Safety and Efficacy of Nanosecond Pulsed Electric Field Treatment of Sebaceous Gland Hyperplasia. Dermatologic Surgery 2020; 46: 803–809
 Nuccitelli R, LaTowsky BM, Lain E, Munavalli G, Loss L, Ross EV, Jauregui L and Knape WA (2021), Safety and Efficacy of Nano-Pulse Stimulation Treatment of Non-Genital, Cutaneous Warts (Verrucae). Lasers in Surgery and Medicine.
 <u>https://doi.org/10.1002/ism.23423</u>

3. Hruza G, Żelickson B, Selim M, Rohrer T, Newman J, et al. Safety and Efficacy of Nanosecond Pulsed Electric Field Treatment of Seborrheic Keratoses. Dermatologic Surgery 46(9):1183-1189, September 2020.

Early Commercial Experience



BEFORE

AFTER (42 days post-procedure)

Courtesy of Dr. Joe Overman, Arvada, CO



BEFORE



AFTER (42 days post-procedure)

Courtesy of Dr. Leslie Stewart, Greenwood Village, CO



Strong KOL Support and Advocacy

Deep network of leading aesthetic dermatologists

7 CLINICAL PUBLICATIONS ON NPS TECHNOLOGY





Scientific Advisors & Consultants



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 Commercial Launch



Transitioning from Controlled Launch program to broad commercialization

2021	Q1	Q2	Q3	Q4	
January 25 CE mark approval		June 25 Health Canada approval	August First clinic completes Controlled Launch and opts	November 15 68 clinics onboarded to date as part of the Controlled	
February 3		June 30	to acquire CellFX System for commercial use	Launch	
FDA clearance February 16 Start of Controlled Launch with first patient treated in the US		First patient treated in Canada June 30 49 clinics onboarded as part of the Controlled Launch	September 30 12 Controlled Launch Program participants opt to acquire their CellFX Systems and transition to commercial use	 Anticipated Complete onboarding of ~75 Controlled Launch participant c Continued Controlled Launch participant conversions 	
February 19 First patient t with the CellF	reated in EU			• Transition to full commercial lau	



Cycles Enable Treatment Delivery

Innovative utilization based business model aligns incentives



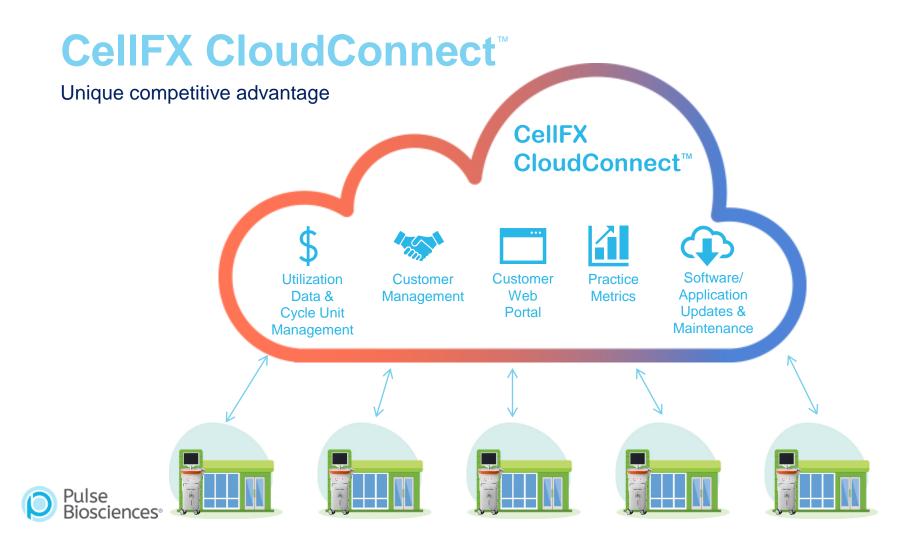
Patients want to pay based on number of lesions treated

 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







Practice Economics

MD Franchise Based on Expanding Utilization Over Time

Compelling Short-term Economics





Initial Investment \$45.000



Lesions Revenue per Patient per Patient

5 lesions / patient \$900-\$1,200*

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< Four Month **Return on**

Investment

Long-term Practice Franchise







Avg. Ten Patients / Week

- 10 x 15-minute treatment session (2.5 hrs)
- · Benign Lesion, SH, SK, & Warts
- Room for growth with additional apps

Avg. Five Lesions per Patient Net MD Rev

per lesion

\$150-\$200

Revenue to **Practice** ~\$500.000 / Year

Gross Profit to Practice ~\$400,000 / Year



*Average revenue per patient of \$1,050 used in ROI calculation yields \$63,000 revenue in < four months

Extensive CellFX Application Pipeline



EPIDERMAL

- Seborrheic keratosis
- (Raised & Macular)
- Warts (Pivotal Study)
- Common moles/nevi
- Cherry angioma
- Actinic keratosis
- Molluscum contagiosum
- .
- .

MID-DERMAL

- Sebaceous hyperplasia
- Acne
- Syringoma
- Intradermal nevi
- Oily skin
- Keloid scar
- Melasma
- White/blond hair
- Rhinophyma
- Xanthelasma
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Initial Applications

Early Feasibility Work Underway

Future Applications

DEEP DERMIS / SUBCUTANEOUS

- **Basal Cell Carcinoma**
- (BCC
- Sweat glands
- Congenital melanocytic nevi
- Subcutaneous fat
- Hidradenitis suppurativa
- Lipomas
- Sebaceous cysts

SUBCUTANEOUS

DERMIS

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 4Q21
 - **Completed 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 1H22
 - Completed treatments for FDA IDE approved treat and resect basal cell carcinoma feasibility study, tissue analysis expected to be completed 1Q22
- **CE mark approval** received January 22, 2021
- Health Canada approval received June 9, 2021
- Australian TGA approval received November 8, 2021

Upcoming Milestones

- Continue transition of Controlled Launch Program clinics to commercial use
- Onboarded 68 Controlled Launch Program clinics and expect to complete onboarding of 75 total clinics in 4Q21
- Clinical and regulatory activity to expand CellFX System clinical applications
 - Received Health Canada Medical Device License approval
 - Subsequent 510(k) submissions
 - SH planned for 4Q21
 - Non-genital warts planned for as early as end of 1H22
 - Tissue analysis as part of FDA IDE approved BCC feasibility study expected to be completed in 1Q22
- Progress Clinical and Scientific programs
 - Continued presence at scientific meetings



As of September 2021

120 issued patents globally owned & licensed

Additional 106 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

Financial Snapshot

Initial revenue recognized in 3Q21



- As of 9/30/2021 cash, cash equivalents and investments were \$42.0MM
- February 2021 launched At-the-Market Equity Offering of up to \$60MM

Pulse Biosciences®

	-	otember 30, 2021
		(In thousands)
Cash, cash equivalents & investments	\$	41,991
Accounts Receivable		72
Inventory		4,281
Prepaid expenses & other assets		3,231
Property, plant and equipment		2,446
Goodwill & intangibles		6,173
Right of use assets		8,954
Total assets	\$	67,148
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Accounts payable & accrued liabilities Deferred revenue	Ş	7,677
Lease liability		, 10,991
Note payable		1,086
Stockholders' equity		47,387
Total liabilities & stockholders' equity	\$	67,148

Shares outstanding (as of 9/30/2021)

29,631

Leadership Team

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

Management

Darrin Uecker*	President & Chief Executive Officer	gynesuncs' computermotion.
Edward Ebbers	EVP & GM, Dermatology	ZELTIQ thermage
Sandra Gardiner	EVP & Chief Financial Officer	CUTERA tria.
Mitch Levinson*	Chief Strategy Officer	Cerebrotech ZELTIQ thermage

Board of Directors

Bob Duggan, Chairman		Maky Zanganeh, D.D.S.	Mitch Levinson		
Spharmacyclics INTUITIVE		Spharmacyclics computermotion.	Cerebrotech ZELTIQ thermage		
Richard van den l	Broek	Manmeet Soni	Laureen DeBuono		
HSMR Advisors Pulse Biosciences°	©pharmacyclics [,]	<i>EREATA</i> 2 Alnylam <i>Spharmacyclics</i> ZELTIO	FLG PARTNERS R+F CIRCA		

Thank You