

WELCOME TO OUR FIRST DIGITAL CAPITAL MARKETS DAY!



AGENDA

Welcome

Isabelle Ljunggren

Introduction to the future of medicine

Dr. Robert Langer

Bioconvergence agenda and strategic focus areas 2021

Erik Gatenholm

Financial update Q1, 2021 and M&A agenda

Gusten Danielsson

Introduction to Biosciences

Dr. Jonas Schöndube

Break (5 min)

Introduction to Bioprinting

Artur Aira & Cecilia Edebo

The BIO MDX Series

Dr. Héctor Martínez & Cecilia Edebo

How we work and interact with our engaged customers and partners

Dr. Itedale Namro Redwan & Mariana Andrade

Break (5 min)

Introduction to Industrial Solutions

Dr. Holger Eickhoff

Q&A session

Moderator Ulrik Trattner

Thank you

Isabelle Ljunggren

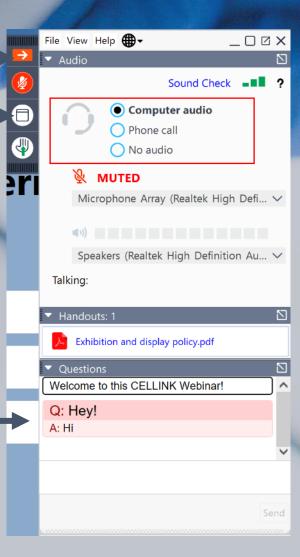


Expand control options •

Maximize window size



Ask questions and chat with the presenter





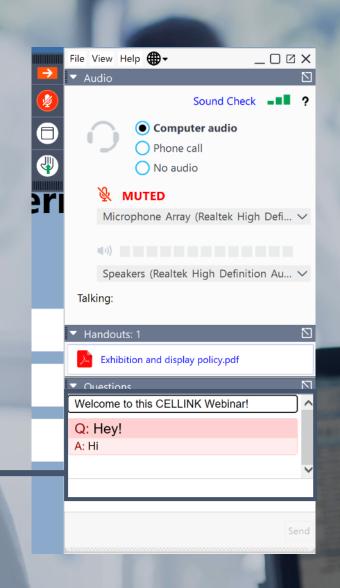
Q&A SESSION

PARTICIPATE BY SENDING YOUR

UQUESTIONS TO THE QUESTION BOX

FUTURE

Write your questions here!





INTRODUCTION TO THE FUTURE OF MEDICINE



Dr. Robert LangerMIT and co-founder of Moderna,
Scientific Advisor CELLINK



INTRODUCTION TO THE FUTURE OF MEDICINE

Dr. Robert Langer MIT and co-founder of Moderna, Scientific Advisor CELLINK





Erik Gatenholm CEO, CELLINK



BIOCONVERGENCE AGENDA AND STRATEGIC FOCUS AREAS

Erik Gatenholm, CEO



PASSION - INSPIRATION - PERSISTENCE

2016

CELLINK = bioprinting

Bioprinting

7 employees

TAM, \$1.4Bn+

20 laboratories

5 publications

Market Cap MSEK 645



CELLINK = CELLINK Group,
Business Areas and Group Companies

Bioconvergence

700 employees

TAM, \$150Bn+

2,000+ laboratories

1,700+ publications

Market Cap BSEK 22.63

















MODERN HEALTHCARE CHALLENGES THAT SHAPE THE FUTURE OF BIOCONVERGENCE.





9 out of 10 drugs in development fail in clinical trials. It can take 10 to 12 years to develop a new drug at a cost of more than \$2Bn.



While 108,000 chronically ill patients wait on organ donation lists, only 39,000 transplants are performed in the U.S. each year. For the lucky few who do find a donated organ, however, the rejection failure rate is an astounding 50%.

Sources: organdonor.gov (U.S. government on organ donation and transplantation), U.S. department of health and service, The American Society of Transplantation and Organ procurement and transplantation during the COVID-19 pandemic (The Lancet).

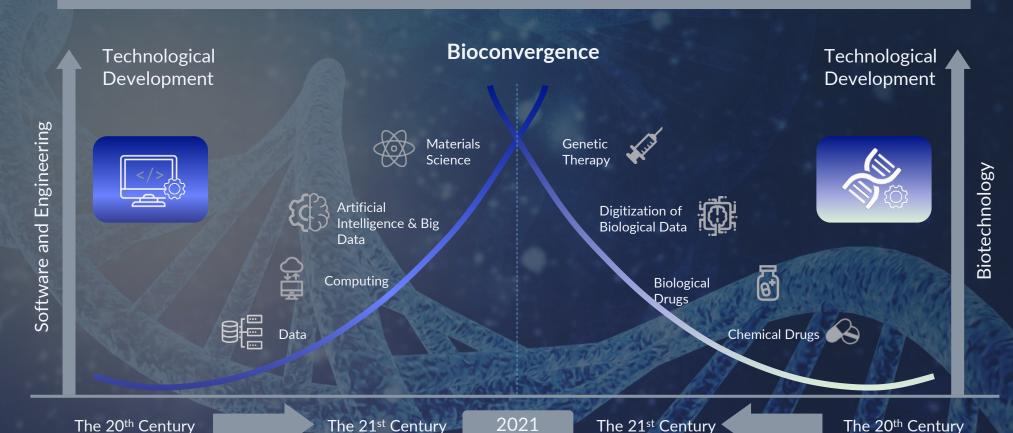
CELLINK 2021, All rights reser





BIOCONVERGENCE TO FORM THE FUTURE BASE OF MEDICINE AND TO RESHAPE THE GLOBAL HEALTH INDUSTRY

Bioconvergence connects various technologies from the fields of biology and engineering to identify and develop precise, personalized and effective medical solutions



© CELLINK 2021, All rights reserved.



THE BIO REVOLUTION*

Human health is one of the most significant domains where biological advances are being applied. Biology is already helping save lives through innovative treatments tailored to our genomes and microbiomes. In the future, we estimate that almost half of the global disease burden could be addressed through applications that are scientifically conceivable today.



Four arenas of bio innovations



Biomolecules

Mapping and engineering
intracellular molecules



Biosystems Mapping and engineering cells, tissues, and organs



Biomachine interfaces
Connecting nervous
systems of living
organisms to machines



Biocomputing
Using cells and cellular
components for
computation

\$2T-\$4T

of annual direct economic potential globally in 2030–40 (significantly higher with downstream and secondary effects)



THE SOLUTION IS FOUND IN BIOCONVERGENCE





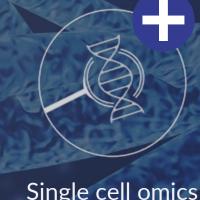






Diagnostics and

biological sensors





- tissue printing



Single cell omics

Optics and Imaging

Biopharmaceuticals

© CELLINK 2021, All rights reserved.



THE BIOCONVERGENCE COMPANY





THE BIOCONVERGENCE COMPANY

Bioprocessing

Reagents

Microscopy

Live cell imaging

Cell culture

Genomics

Biosensors

Bioinformatics



POC

OMICS

Proteomics

Bioreactors

Diagnostics

Single cell sorting

Liquid handling

Bioprinters & Bioinks

Robotics

Cell line development

Consumables

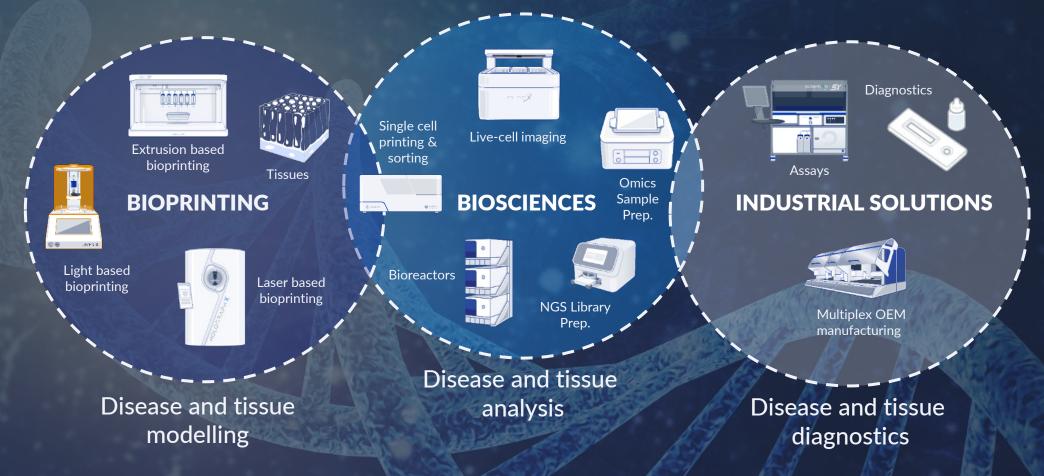
© CELLINK 2021, All rights reserved.

NGS



IT IS THE ERA OF BIOCONVERGENCE

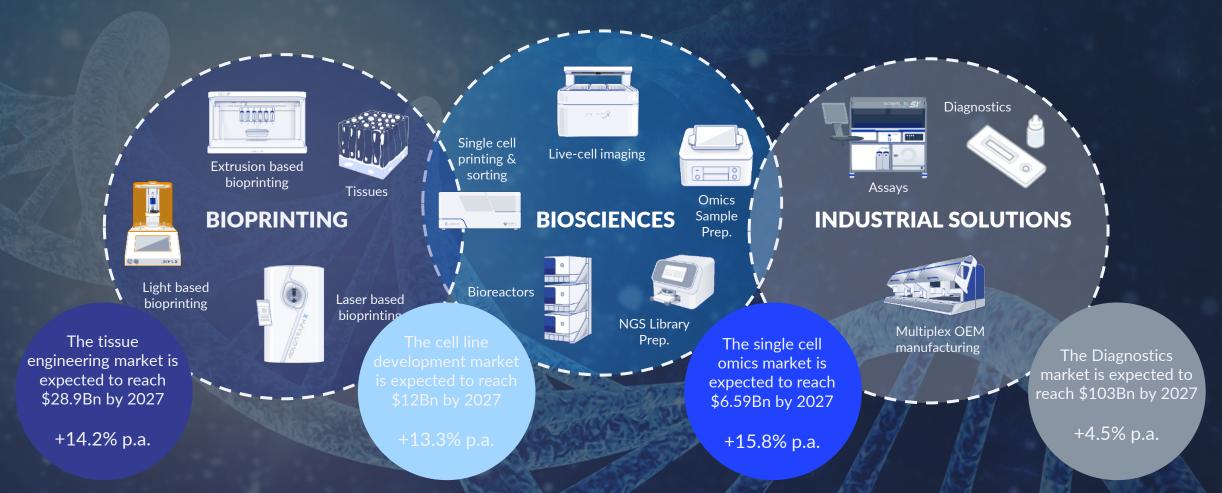
Serving our customers in the best possible way by offering market leading workflows



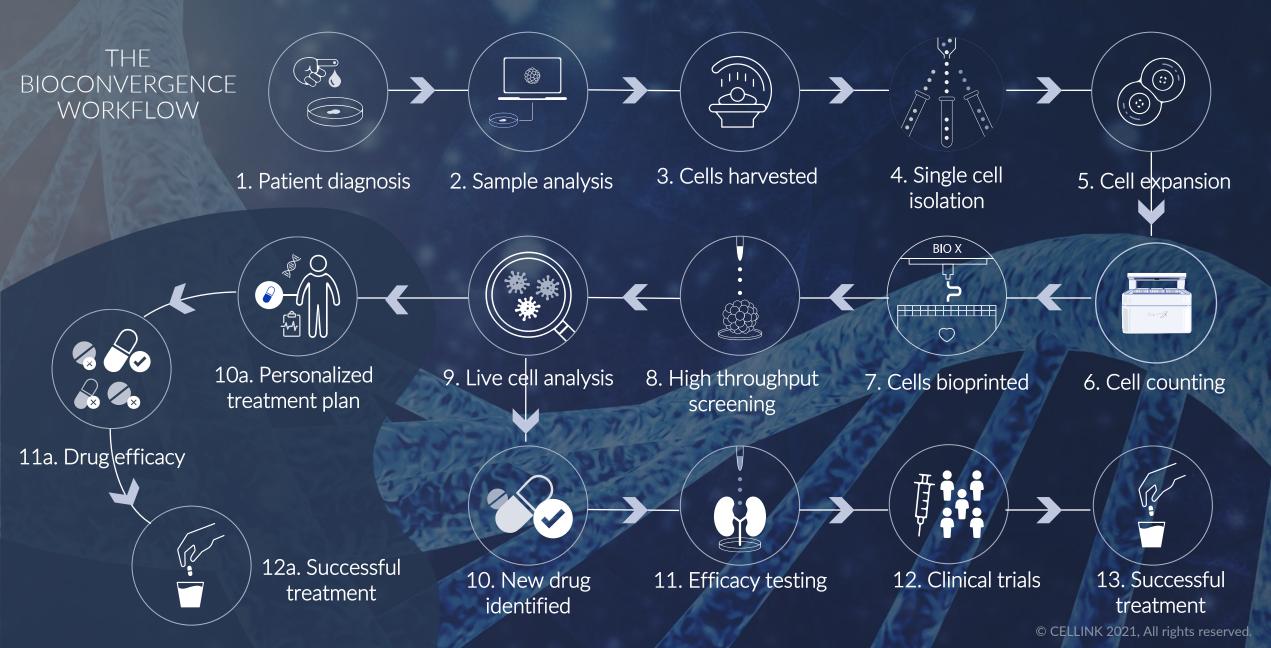


BIOCONVERGENCE MARKET

APPLICATION AREAS \$150Bn+



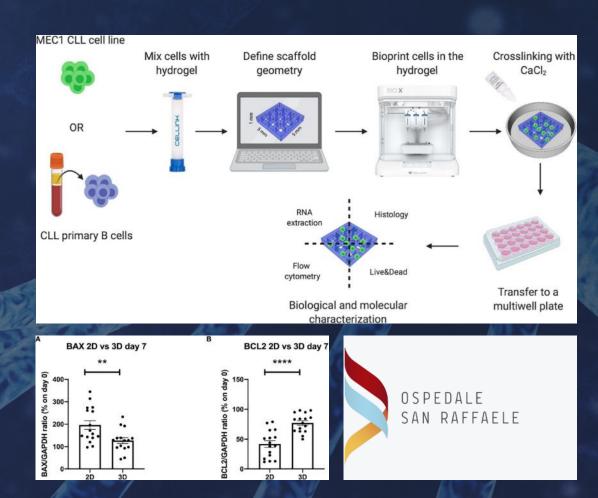






3D Bioprinting allows the establishment of long-term 3D culture model for chronic lymphocytic leukemia cells

- Chronic Lymphocytic Leukemia (CLL) represents the most common leukemia in the western world and remains incurable. Leukemic cells organize and interact in the lymphoid tissues, however what actually occurs in these sites has not been fully elucidated yet.
- Studying primary CLL cells in vitro is very challenging due to their short survival in culture and also to the fact that traditional two-dimensional in vitro models lack cellular and spatial complexity present in vivo. Based on these considerations, we exploited for the first time three-dimensional (3D) bioprinting to advance in vitro models for CLL. This technology allowed us to print CLL cells (both primary cells and cell lines) mixed with the appropriate, deeply characterized, hydrogel to generate a scaffold containing the cells, thus avoiding the direct cell seeding onto a precast 3D scaffold and paving the way to more complex models.
- Using this system, we were able to efficiently 3D bioprint leukemic cells and improve their
 viability in vitro that could be maintained up to 28 days. We monitored over time CLL cells
 viability, phenotype and gene expression, thus establishing a reproducible long-term 3D
 culture model for leukemia.
- Through RNA sequencing (RNAseq) analysis, we observed a consistent difference in gene expression profile between 2D and 3D samples, indicating a different behavior of the cells in the two different culture settings. In particular, we identified pathways upregulated in 3D, at both day 7 and 14, associated with immunoglobulins production, pro-inflammatory molecules expression, activation of cytokines/chemokines and cell-cell adhesion pathways, paralleled by a decreased production of proteins involved in DNA replication and cell division, suggesting a strong adaptation of the cells in the 3D culture.
- Thanks to this innovative approach, we developed a new tool that may help to better mimic
 the physiological 3D in vivo settings of leukemic cells as well as of immune cells in broader
 terms. This will allow for a more reliable study of the molecular and cellular interactions
 occurring in normal and neoplastic conditions in vivo and could also be exploited for clinical
 purposes to test individual responses to different drugs.





STRATEGIC FOCUS AREAS 2021

Strengthen our position as *the* bioconvergence company in the world through strong business areas and dedicated group companies



M&A and tech development

Strengthen bioconvergence position



Financials

Deliver on financial targets



Customers

Best customer care, design, quality and supply chain in the industry



People

Happy and motivated team



Sustainability

Develop a sustainability agenda



STRATEGIC FOCUS AREAS 2021

Area

Focus 2021

M&A and tech development agenda

Strengthen bioconvergence position

- Active customer centric M&A agenda
- Develop and capitalize on our strong R&D and tech development agenda (Group synergies)
- Continue collaboration through partnerships with Academia, research organizations and customers

Financials

Deliver on financial targets

- Deliver on financial targets (35% organic growth and show positive EBITDA)
- Focus on sales and value drivers for growth at Group company level

Customers

Best customer care, design, quality and supply chain in the industry

- Continue to build direct sales organization on main/growth markets
- Unified global service capabilities at Business Area level
- Focus on product design based on user experience by global design team
- Implement lean and efficient supply chain
- Implementation of ERP- and CRM system



WORLD-LEADING CUSTOMERS IN FOUR SEGMENTS

Universities & research organizations

- biomaterials research, cell biology, teaching etc.
- Exploratory research that, if successful, will convert to clinical applications.
- ~600 institutions in 65 countries (>11.000 relevant academic institutions on a global basis).

Pharmaceutical companies

- Cancer research (in vitro. clinical and preclinical studies), biopharmaceuticals, cell line development for biologics, gene therapy, tissue culture & engineering.
- Over 50 customers with top 20 pharma's as customers.

Diagnostic companies

- Multiplex assays, lateral flow tests, microfluidic devices, IVD assays and array printing.
- Drug development, cancer research, biopharmaceuticals, gene therapy and tissue culture &
- Long relationships and contracts with many of the larger actors in the field.

Cosmetic companies/others

- Toxicity and cosmetic tests on human tissues, working with major players in cosmetics and injectables. Large potential in cosmetics in 3-5 years.
- Wide range of application areas (e.g., packing solutions, car materials, skin tissues and transplants). Large potential in the next 5-10 years.





Imperial College

KOCHINSTITUTE



HARVARD UNIVERSITY

Tutts































AstraZeneca 22























PARTNERSHIPS AND COLLABORATIONS EXPLORING THE BIOCONVERGENCE AGENDA

PARTNERSHIPS 2020-2021



May 2020: collaboration with AstraZeneca to utilize CELLINK's 3D-bioprinting technology for liver organoid culture.



June 2020: CELLINK and Lonza joined forces to Offer Complete 3D Cell Culture Workflows.



August 2020: strategic partnership with Carcinotech to advance 3D-bioprinting technology for cancer research.



August 2020: collaboration with AstraZeneca to develop plate based micro-bioreactors for cell line development workflows.



Nov 2020: CELLINK will develop 3D bioprinted personalised scaffolds for tissue regeneration of ankle joints as part of the TRIANKLE project.



Dec 2020: CELLINK and Atelerix team up to enable the shipping at room temperature of fragile 3D bioprinted constructs.



Jan 2021: Third year extension of extended partnership to utilize CELLINK's 3D bioprinting technology for drug discovery.



March 2021: collaboration agreement for cellulose-based bioink.

SIGNIFICANT COLLABORATIONS



Several research projects, for instance, creating a 3D-Bioprinted biomimetic heart value with correct mechanical properties.



Study the influence of microgravity and hypergravity on living systems.

CELLINK sent neural crest cells to space.



ETH summer school program focused on 3D printing.



Development of bioprinted liver and skin tissue models.



Bioprinting of patient-specific cancer tumours.

SELECTED DISTRIBUTORS

















STRATEGIC FOCUS AREAS 2021

Area

Focus 2021

People

Happy and motivated team

- Integration strategy for acquisitions, initial 100-day plan
- Create a shared digital workplace for the Group and encourage knowledge sharing
- Continuous monitoring employee satisfaction
- Training and development on Group level thru CELLINK academy

Sustainability

Develop sustainability agenda

- Sustainability agenda for the Group with sustainability targets (to be launched 2021/2022)
- Development of products and services through technologies for minimizing animal trials
- Mapping towards UN:s Sustainable Development Goals













Gusten Danielsson CFO, CELLINK





LONG TERM FINANCIAL TARGETS 2019-2022



FinancialsDeliver on financial targets

Organic growth

CELLINK's objective is to achieve an annual organic sales growth of >35%, supplemented by strategic acquisitions.

Outcome 2020

Organic growth was 48% (77%)

EBITDA margin

CELLINK's objective is to have a positive EBITDA margin.

Outcome 2020 EBITDA margin was 0.2% (-7.8%)

Capital structure

CELLINK aims to maintain a ratio of Net Debt to EBITDA of 3.0x, and may temporarily exceed this level (e.g., as a result of acquisitions).

Outcome 2020 cash excl. leasing debt ounted to MSEK 756,

ncluding net debt MSEK 676

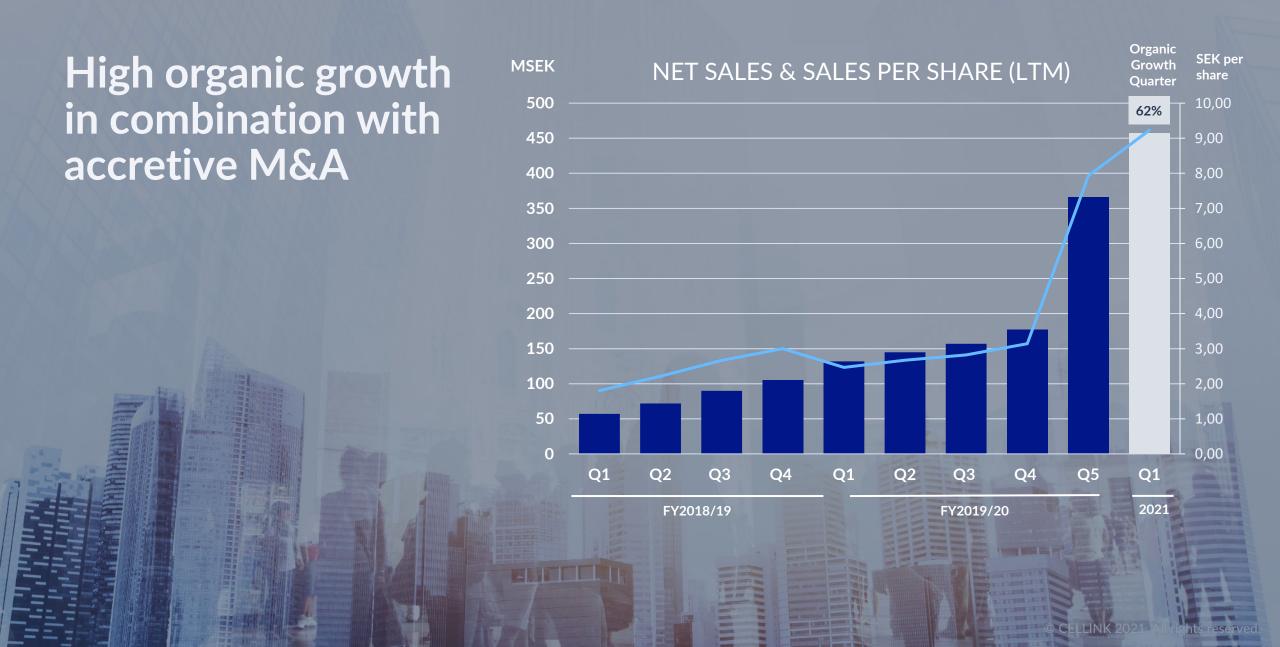


HIGH ORGANIC GROWTH, STRENGTHENED GROSS MARGIN AND STRATEGIC ACQUISITIONS

Q1: January-March 2021

- Net sales amounted to MSEK 129.5 (38.0), which corresponds to an increase of 241% (55) compared to the corresponding period previous year, of which 62% (25) was organic growth.
- EBITDA amounted to MSEK -34.9 (-5.7), corresponding to a margin of -26.9% (-14.9). The operating profit was affected by acquisition costs totalling MSEK 20.5.
- Profit/loss for the period amounted to MSEK -47.8 (-33.5), which generates earnings per share after dilution of SEK -0.90 (-0.80). The result was positively affected by the market valuation of the company's short-term investments of MSEK 4.4 (-22.7).
- The gross margin amounted to 77.3% (74.8), mainly due to higher revenue per product, improved product mix and increased share of sales in services and consumables.
- Rolling twelve-month net sales from consumables amounted to MSEK 46.2 (19.2), an increase of 140%. The share of total product sales was 12.9% a decrease of 0.2 percentage points (13.1% in the comparison period).





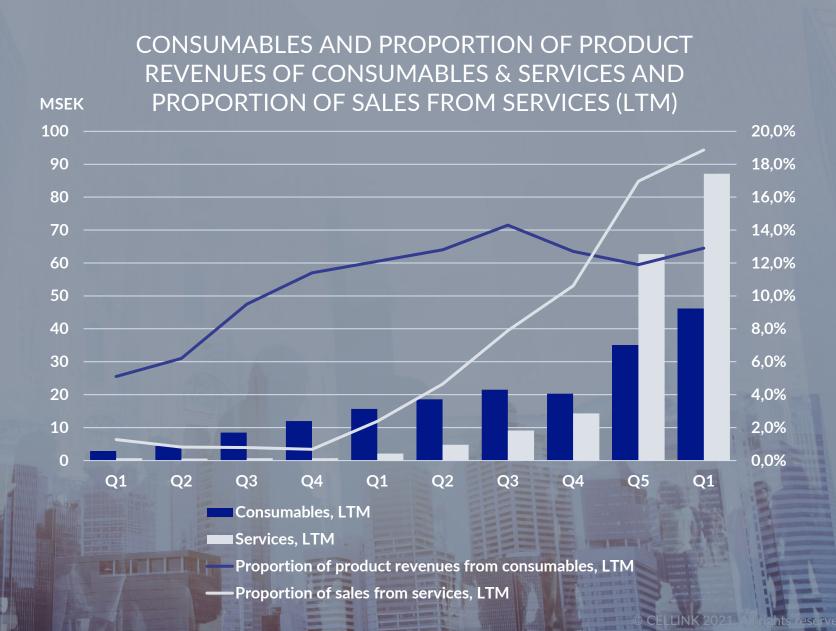








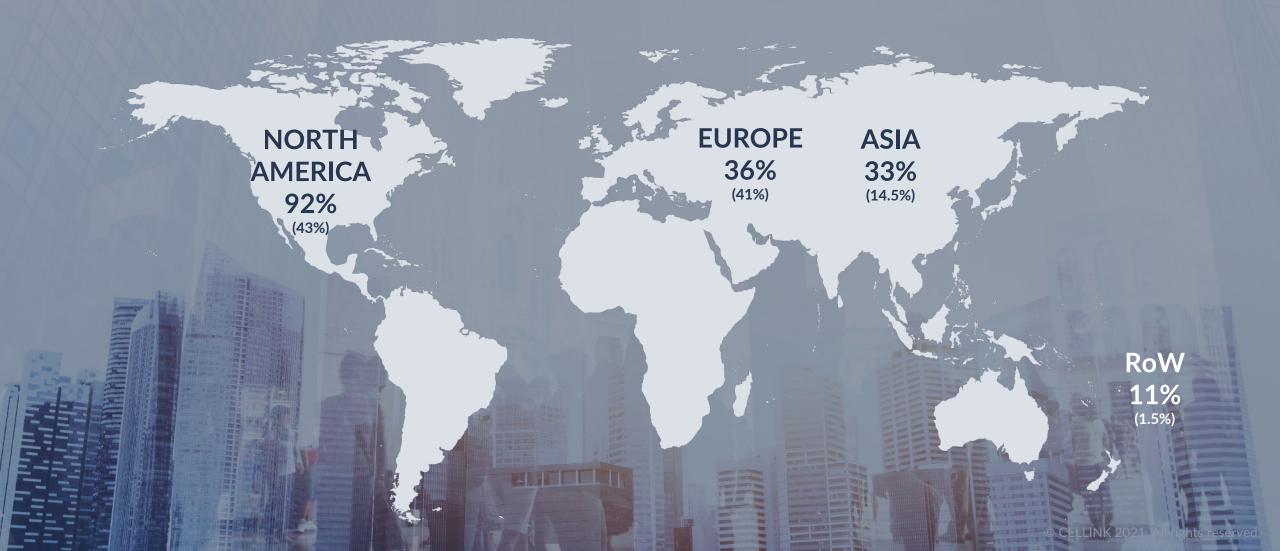
Increasing revenue from consumables and services





ORGANIC GROWTH PER REGION Q1, 2021 vs. Q1, 2020

(Share of sales, Q1 2021)







M&A and tech development Strengthen bioconvergence position

CELLINK issued senior unsecured convertible bonds convertible into Class B shares and a Class B share issue in an overall amount of SEK 3.0 billion on March 12, 2021

- Assuming full conversion, the Bonds will entail a dilution of 4.8% of the total number of outstanding shares and 3.8% of the votes in CELLINK.
- The Bonds will bear a coupon of 2.875% per annum, payable semi-annually.
- The conversion price was set at SEK 598.50, representing a premium of 42.5% above the reference share price.
- Placement of approximately 1.04 million existing B Shares of CELLINK at a placement price
 of SEK 420 per share on behalf of certain subscribers of the bonds who wished to sell these
 in short sales to purchasers to hedge the market risk to which they are exposed with respect
 to the Bonds that they acquire in the Offering.
- The share issue consisted of 3,571,429 New B Shares, equal to approximately 7.0% of the current outstanding Class B share capital of CELLINK and 6.8% of the current outstanding total share capital of CELLINK.



CUSTOMER CENTRIC M&A AGENDA

CELLINK invests in entrepreneurs' enthusiasm and passion for what they do. We aim to find and acquire companies that are built on determined people with a strong desire to create the future of medicine.

FINANCIAL TARGETS

- Revenue growth potential in line with CELLINK's financial targets
- Potential for EBITDA margins above industry average
- Proven historical track record of products and customers

STRATEGIC TARGETS

- Increase share of ownership of the value chain and improved value proposition
- Increased market power and know-how
- Potential to branch out into new additional verticals or strengthen regions
- Cross-selling potential and improved customer offering



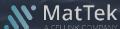
M&A and tech development Strengthen bioconvergence



SUCCESSFUL INTEGRATION OF **ACQUIRED COMPANIES**



M&A and tech development Strengthen bioconvergence





EV/sales multiples







dispendix



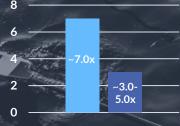
- €5 MM in transaction value - 60% in shares / 40% in cash
- Part of the bioprinting and drug development screening process
- Cross-sales opportunities and increased customer value with CELLINK

At acquisition

Post acquisition (per May 2021)







- ~€30 MM in transaction value - 60% in shares / 40% in cash
- ~40% EBITDA margin
- Strengthen position in research workflow and with big pharma (~90% of customer base)
- Cross-sales opportunities with CELLINK and Dispendix

scienion





- ~€80 MM in transaction value - 50% / 50% in shares/cash
- ~25% EBITDA margin
- Strengthen position in single-cell handling
- Creating a low volume dispensing powerhouse with Dispendix and a dominant player in singlecell dispensing with **CYTENA**

 ~€70 MM in transaction value - 40% / 60% in shares/cash

GINOLIS

EV/sales multiples

- ~12% EBITDA margin
- Strengthen position in automation and diagnostics
- Creating an automation and low-volume dispensing powerhouse with SCIENION and **Business area Bioprinting**



- ~MUSD 72 in transaction value - 20% / 80% in shares/cash
- ~22% EBITDA margin
- Strengthen position in tissue focusing on advanced in-vitro technology for clinical and pre-clinical studies offering in-vitro human tissue model innovation, cell isolation and cell culture
- Creating synergies in **Business area Bioprinting**



M&A and tech development Strengthen bioconvergence position



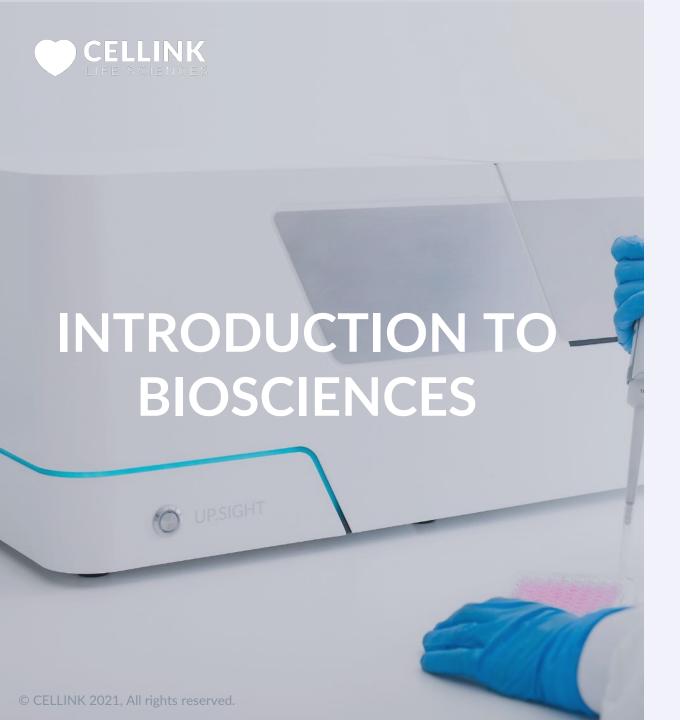
FinancialsDeliver on financial targets

FOCUS 2021

Deliver on our financial targets.

Continue our active and customer centric M&A agenda.

Identify cost efficiency synergies in the Group.





Dr. Jonas SchöndubeBusiness Area Manager
Biosciences



INTRODUCTION TO BIOSCIENCES

Dr. Jonas Schöndube Business Area Manager



BUSINESS AREA BIOSCIENCES







Market segments

Cell Line Development, Live-cell imaging, Single-Cell Omics, Liquid Handling

Customer base

Biopharma & Biotech companies,

Main competitors

Berkeley Lights; BeckmanCoulter, Sartorius: Becton, Dickinson and Company (BD)

TAM

\$11.11Bn with expected growth of 13% p.a.

Product offering

UP.SIGHT, C.SIGHT, F.SIGHT, C.BIRD, CELLCYTE X, I.DOT, C.WASH

Sales model

Instruments, consumables and services

Market position

>15 of the top 25 pharma companies are using CELLINK products to develop cell lines. In the other segments we are fast growing challengers



MAJOR TRENDS - MARKET DRIVERS

1

Biosimilars/bio-betters

More and more
companies develop
biopharmaceutical
drugs, because many
plockbusters fell of the
patent cliff.
Cell line development is

2

Cell and Gene Therapy

Revolutionary
therapeutic approache
are looking for the
right manufacturing
technologies.

3

Software/Automation

More value by software on existing hardware.
The pandemic has broken the barrier of hesitation for many customers to adopt automation.

4

Single-cell multi omics

Most cell analysis is or will be single-cell analysis in the near future. We provide the tools.



M&A and tech development

Strengthen bioconvergence position

CELL LINE DEVELOPMENT





Transfection

First round of cloning

Second round of cloning

Proof of clonality & colony growth monitoring

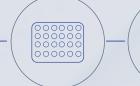
Plate Imaging





24-well plate 24 deep well plate 350-800 µl

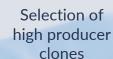
2-3 ml



Complicated upscaling process



Shaking flask 10-20 ml



up to 13 weeks time saving

99.99% assured clonality and increased throughput

UP.SIGHT workflow



Transfection

UP.SIGHT™



Single-cell cloning & multiple proof of clonality

C.BIRD™



Early suspension culture & upscaling



Bioreactor 15-50 ml

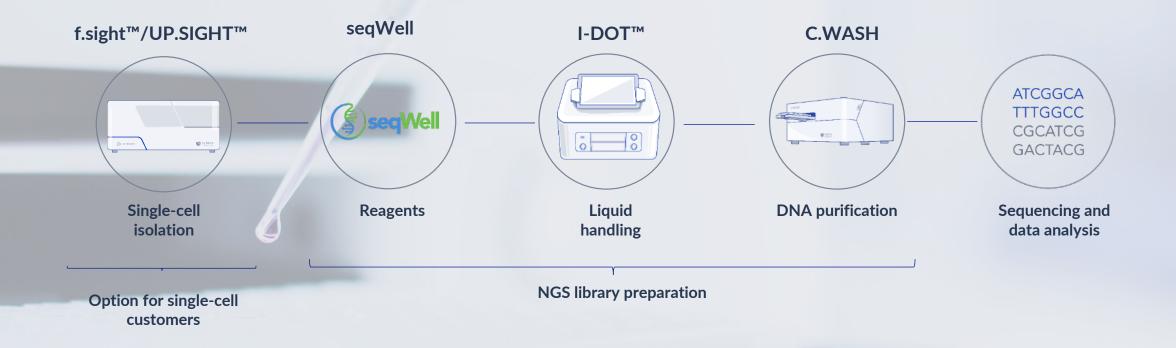


Selection of high producer clones



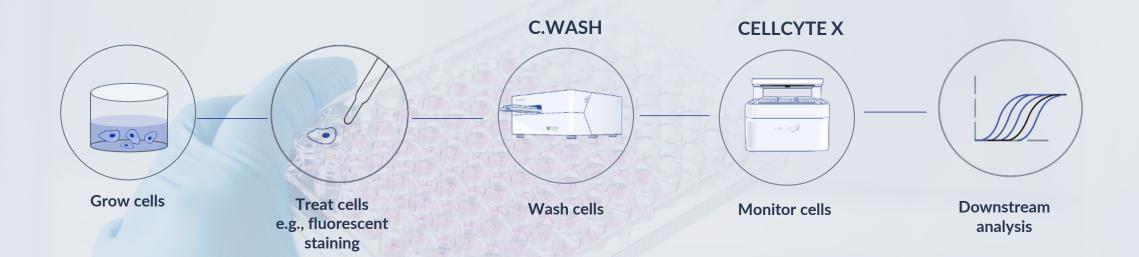


SINGLE-CELL OR COVID-19 SEQUENCING WORKFLOWS





HIGH-CONTENT SCREENING WITH CELL-BASED ASSAYS





FOCUS 2021

1

Accelerate revenue growth, especially for consumables and reagents.

Whilst keeping profitability positive 2

Applications and Workflows

- Cell line development
- Single-cell multi-omics
- High content screening

3

Software development

- The CELLINK
 Biosciences SW
 team is growing
 heavily.
- Software products will be a main value driver in the future.







= BREAK -

NEXT SESSION: BIOPRINTING

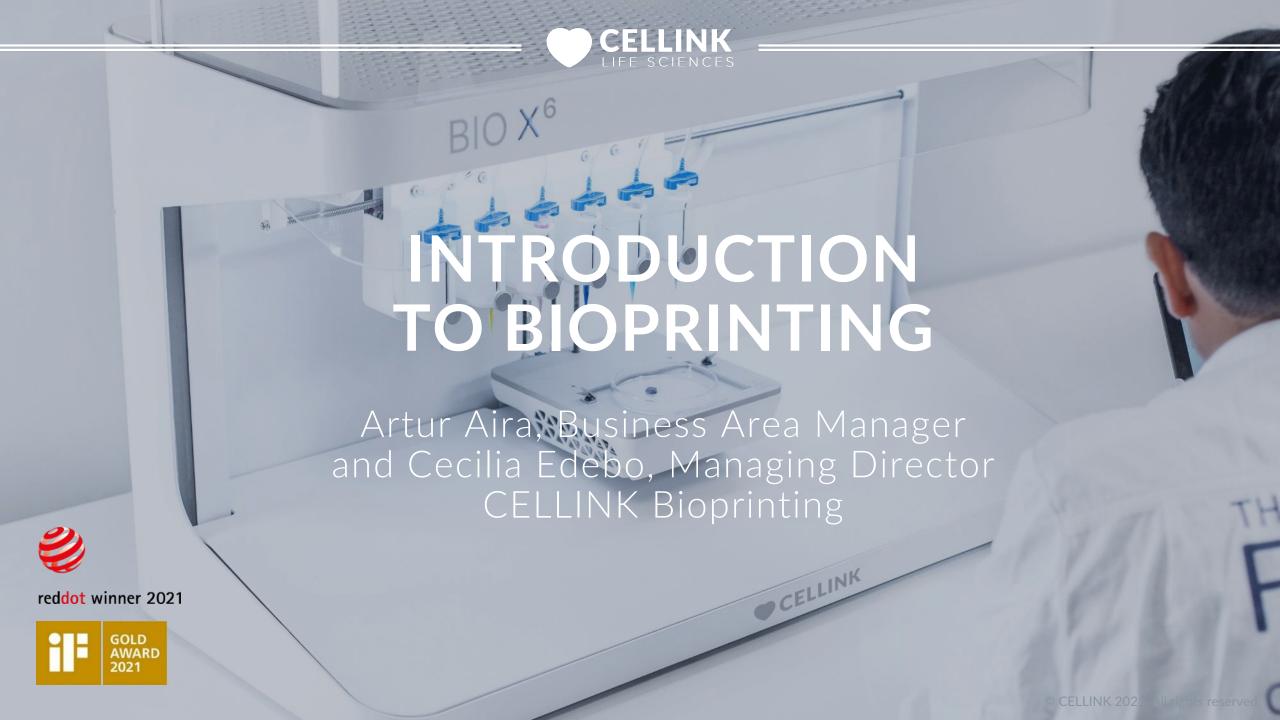




Artur AiraBusiness Area Manager Bioprinting



Cecilia EdeboManaging Director
CELLINK Bioprinting





BUSINESS AREA BIOPRINTING



Market segments

Research: regenerative medicine, 3D cell culture, drug research

Clinical/medical: blood vessels, bone, cartilage, skin, pills, tissue, implants/prosthetics, Skin applications, microfluidics

Consumer product testing

Market position

Market leading position

Customer base

Pharmaceutical-, cosmetics- and industrial food companies, med tech industry, academia and research organizations

Product offering

Light based bioprinting
Laser based bioprinting
Extrusion based bioprinting
Tissue models
Bioinks

Main competitors

Corning Thermo Fisher Scientific Sigma-Aldrich Episkin Greiner Bio-One

TAM

The 3D printing market is expected to reach between \$1.4-4Bn by 2024 with a CAGR of 20-35% annually

Sales model

Instruments Consumables Services



MAJOR TRENDS - MARKET DRIVERS

1

3D Bioprinting

2

Tissue Models

3

Microfluidics

4

Personalized Medicine





Long-term 3D bioprinting market potential

The cell culture market is expected to reach \$33Bn by 2025 with a CAGR +11.8% annually. The 3D printing market is expected to reach between \$1.4-4bn by 2024 with a CAGR of 20-35% annually.

Scaffold-free

Scaffold-based

Traditional 2D Cell Culture

3D Cell Culture

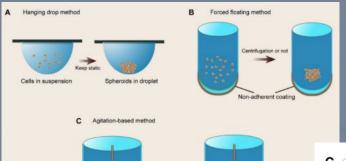
Cell culture

Bioprinting today Including extrusion, ink jet, light based, etc.

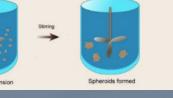


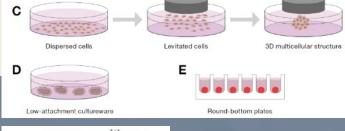
3D Cell Culture

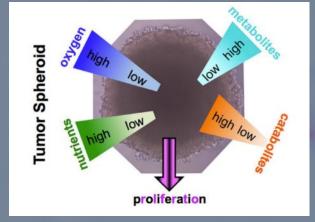
Scaffold-free Methods



Primarily **spheroids** and **organoids** cultured in media.

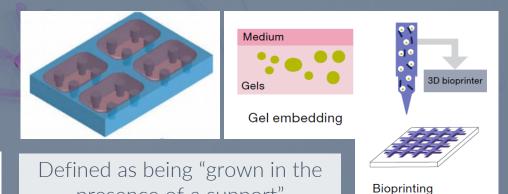






Mainstream Biology 3D Cell Culture primarily uses these methods today.

Scaffold-based methods



Could be simple like **ECM** or **TME** scaffolding (gel) around a simple spheroid or organoid.

presence of a support"

Or scaffolding could also be used to support complicated builds like mini-organs, tissue, etc. replicating the architecture, function, or topography of the in vivo environment.

M&A and tech

developmentStrengthen bioconvergence position





Organs for implant



Our Customers

Mini-organs

"The Biologist"



Scaffold-based methods



Scaffold-free 3D Technique



To scale organs









Sustainability
Develop a sustainability
agenda

THE INCREASE IN TISSUE MODEL TESTING

REDUCTION IN ANIMAL TESTING

FASTER DRUG DEVELOPMENT PROCESS

SUPERIOR MODELS OF HUMAN TISSUE

OPPORTUNITIES WITH IN-VITRO TOXICOLOGY MARKET TISSUE MODELS IN 3D BIOPRINTING

DEVELOPING THE RIGHT THERAPY FOR THE RIGHT DISEASE FOR THE RIGHT INDIVIDUAL

Sustainability



U.S. representatives have introduced bipartisan Develop a sustainability legislation to modernize testing standards, end mandatory animal testing, and lower drug prices

"This reform would allow the use of the nonclinical test methods most likely to predict how a drug will react in humans, including state of the art nonclinical models based on human biology," said Gerry R. Boss, M.D., board member of the Center for a Humane Economy and a long-time researcher in drug development. "It will ultimately streamline drug development, spur innovation, and move drug development forward, benefiting both patients and industry."

Buchanan, Luria, Mace, Sherrill, and **Boyle Introduce FDA Modernization Act** to End Mandatory Animal Testing, **Lower Drug Prices**

April 20, 2021

Legislation Aims to Get Safer Drugs to Patients More Quickly, Embracing Innovation and Shedding Costly, Cruel, Non-Predictive Animal Tests to Revamp FDA New Drug Testing Protocols That Will Reduce Animal Testing and Enable Use of Most Scientifically Advanced Methods



and enable FDA to require the most effective testing methods, regardless of whether animals are used

The FDA Modernization Act would lift requirements for animal testing for any new drug development and enable FDA to require the most effective testing methods, regardless of whether animals are used.



MatTek - MelanoDerm - Skin Lightening



Transfer tissues from agarose to assay medium

Harvest tissues for endpoint analysis

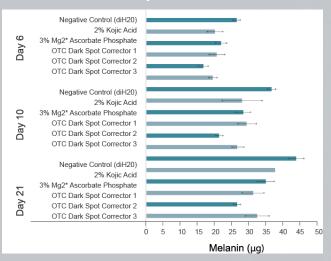
Incubate 1 hour at 37°C, 5% CO₂

Stop exposure by rinsing with DPBS

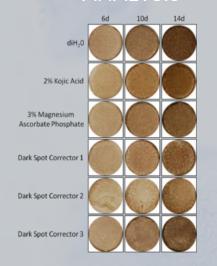
Apply test materials to apical surface of tissues

Repeat applications for 14-21 days

MELANIN QUANTITATION



MACROSCOPIC ANALYSIS



MICROSCOPIC MELANOCYTE ANALYSIS



HISTOLOGICAL ANALYSIS





THE BIOPRINTING ASSORTMENT

















Sustainability
Develop a sustainability



M&A and tech development Strengthen bioconvergence position

CELLINK LIFE SCIENCES

FOCUS 2021

1

Continue with strategic
acquisitions and
capitalizing on the
synergies in the group.
Focus on technologies that
fill gaps in our product
offering as well as broaden
our portfolio for reagents
and consumables.

2

Continue our mission to contribute to decrease of animal testing by providing market leading solutions.

3

Explore the commercial opportunities in; personalized medicine, microfluidics, vascularization and the regeneration of organs.







Dr. Héctor Martínez CTO, CELLINK



Cecilia EdeboManaging Director

CELLINK Bioprinting



THE BIO MDX SERIES

Dr. Héctor Martínez, CTO, CELLINK Cecilia Edebo, Managing Director CELLINK Bioprinting





Ushering in a new era of bioprinting

THE BIO MDX SERIES



High throughput Bioprinting



Precise control at multi-scale resolution from single-cell to spheroid to bulk tissue



Unique modular technology capabilities



Strength across

KEY APPLICATION AREAS



M&A and tech development Strengthen bioconvergence

3D Cell-based assays for high throughput drug screening



Tissue engineering



Cosmetic & wound healing





Superior insights for drug discovery



Personalized implants



Reduced dependency on animal models



Total climate control for utmost sterility

Onboard intuitive & powerful software

High throughput with capacity for up to 27 microplates



Multi-channel bioprinting

Max build volume: 82.5x36x15 cm







FEATURED WORKFLOW

The end-to-end solution for batch biofabrication of personalized medical devices

Load cells in cartridges and initiate human tissue bioprinting protocol





Automated bioprinting at high-throughput



Stimulate tissue development with growth factors



Tissue stabilization & Quality Assurance



Tissue maturation & non-destructive analysis



Personalized implant ready



















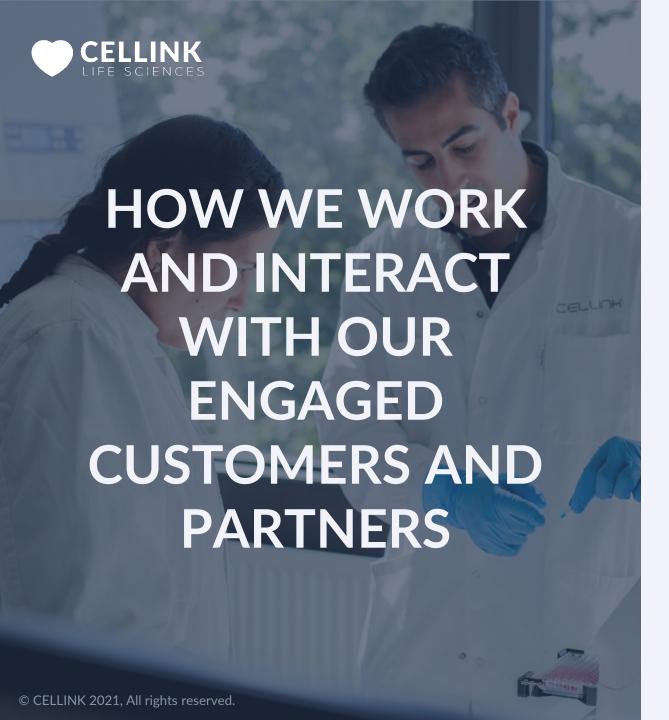












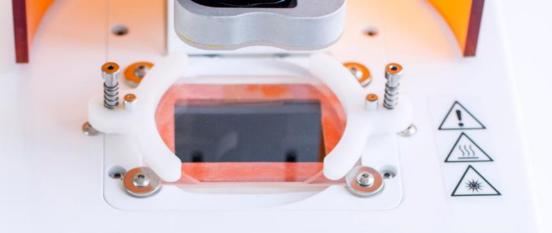


Dr. Itedale Namro RedwanCSO Bioprinting

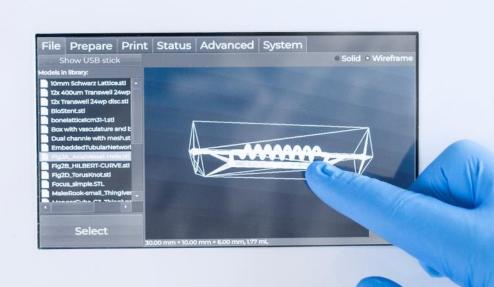


Mariana Andrade
Head of Customer success





















RODUCTS A

APPLICATIONS

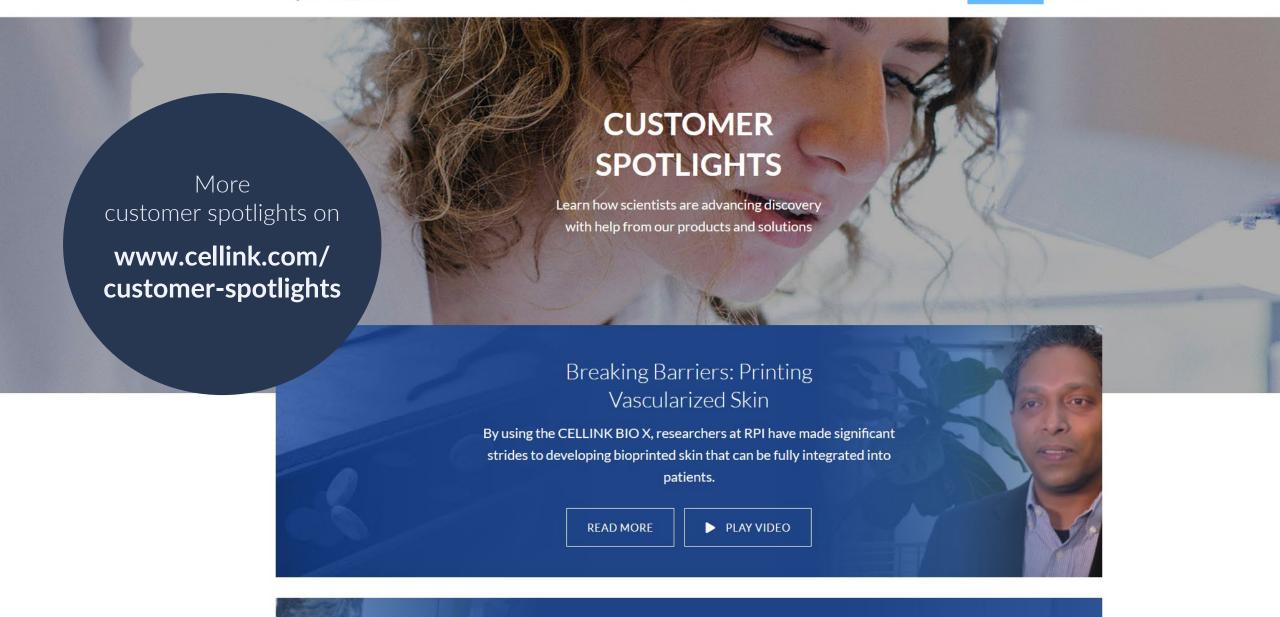
ESOURCES

MPANY

S











Dr. Holger EickhoffBusiness Area Manager
Industrial Solutions





BUSINESS AREA INDUSTRIAL SOLUTIONS

scienion cellenion



Market segments

Diagnostics, Biosensors, Single-Cell Omics and Liquid Handling

TAM

Diagnostics & Biosensors: \$40Bn Single Cell Analysis \$2.1Bn CAGR 10% / 17.2% p.a

Sales model

Instruments, consumables, services and contract manufacturing

Product offering

sciFLEXARRAYER S3-S100. LFDA 1-8. Pixie. Cecilia-L Dispenser, Ginger Software, cellenONE, cellenONE FL, cellenCHIP, proteoCHIP, sciREADERs, Contract Manufacturing,

Main competitors

Tecan, ATS, HP, Sanmina, Nordson, 10xGenomics, Nanocellect

Customer base

Diagnostics companies, Bioprocessing companies, Pharma companies and Research organizations and Academic Institutions

Market position

>15 of the top 25 diagnostics companies are using CELLINK products to manufacture diagnostics. Precision dispensing market leader. We are a very fast mover in single cell genomics and proteomics.



MAJOR TRENDS - MARKET DRIVERS

1

Molecular Diagnostics & Wearables

Disease management with personalized diagnostics is growing. Same is true for precision farming in agbio. The number of tests and platforms is growing, which we support with instrumentation, services and contract manufacturing.

2

Precision medicine

Most cell analysis is or will be single-cell analysis in the near future. We provide instruments, consumables and contract services for our customers.

3

COVID Testing

More and more companies
launch COVID tests. Our
offering is compatible with
testing formats for Antigen
and NAT / PCR Testing. We
are manufacturing partner for
several technologies.

4

Software/Automation

The pandemic has broken the barrier of hesitation for many customers to adopt automation, so that new work regimes will increase the demand for automated workflows in the lab.





Next-Generation Technology for Low Volume Precision Dispensing

SCIENION is the only company capable of dispensing both biological reagents and viable single cells at an industrial scale

Precision Dispensing in pL to μL Range

Single Cell Isolation and Handling

sciDROP NANO

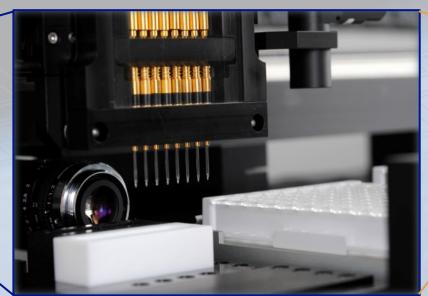
Nanoliter to microliter dispensing in bulk or aspirate/dispense mode

Core Technology

sciDROP PICO

High precision droplet dispensing in the pico- to nanoliter range









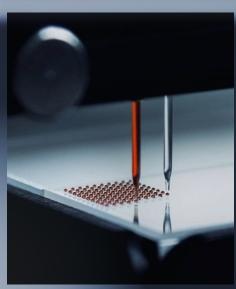
A SCALEABLE APPROACH TO NEXT GENERATION OF DIAGNOSTICS

ADVANCED AUTOMATION

Cutting-edge robotics and automated solutions for manufacturing of diagnostic devices including lateral flow and microfluidics









LATERAL FLOW DEVICE MANUFACTURING

We provide production automation solutions for lateral flow device manufacturing using only very limited amounts of precious bioreagents. Our standard modules give you the flexibility to make a wide range of products with a single automation line, without compromising quality and taking a minimal footprint in your production facilities.

Ginolis Lateral flow device assembly (LFDA) is a standard system for the assembly and packaging of rapid tests

Flexible

Assemble different test variants and products on same line

Intelligent

Vision guided robot operation and quality control provide high quality assurance

Modular

Integrate additional cells for multiple strips, RFID tags, printing, labeling, ultrasonic welding and cap assembly





MEDICAL DEVICE MANUFACTURING THROUGHHIGH QUALITY AUTOMATION

Specialized in modular desktop solutions for the automated production and processing of micro components and medical device products.



Surface Treatment

(Plasma, spray coating, UV curing)

Testing

(Leak, occlusion, functional)

Cutting

(Die, rotary)

Bonding

(Laser & ultrasonic welding, gluing, heat sealing)

Packaging

(Pouching, kitting)

Dispensing

(Array printing, membrane striping)

Marking

(Printing, labelling, laser)

Drying

(Climate chamber, vacuum chamber)

Assembly

(Feeders, bulk feeders, cutters, grippers)

Quality & Process Control

(Machine vision, SPC, testing)

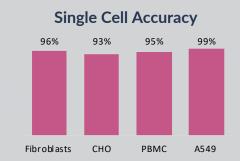


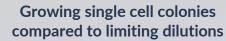
Core Technology: cellenONE

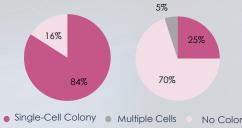
cellenONE allows high-throughput, automated dispensing of individual cells from cell suspensions onto any substrate

Benefits

- Real-time, high accuracy single cell isolation and dispensing
- Gentle acoustic wave for droplet generation
 - Outstanding cell viability for cloning
 - Maintains protein expression for sequencing
- High recovery processing
 - Wide range of samples: rare cells from minute cell suspensions containing just a couple of microliters and a few dozen cells to much larger samples containing thousands of cells
- High resolution optical isolation
 - Wide range of particles (including smaller than < 5μm), from bacteria to nuclei to microbeads to large mammalian cells







Rapid Cell Line Development



Technical Specifications

· · · · · · · · · · · · · · · · · · ·	
Dispensing technology	Acoustic dispensing
Droplet volume	Typically 350pL - 480pL
Single Cell Accuracy	Up to 100%
Frequency	One plate (96 single cells) in 4 min
Capillary material	Borosilicate glass
Dimensions	1300x700x1590 mm
Weight	205kg
Additional commentary	Dead volume: Down to 0 μL Cell imaging: HD camera

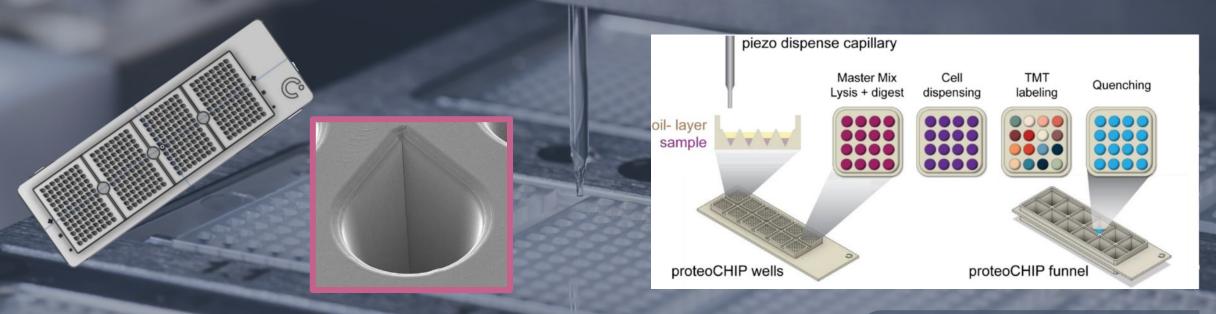






cellenCHIP

A consumable to miniaturize and automate our customers' single cell sample preparation



Offer full solutions to our customers (system + consumables + reagents) for single cell genomics AND single cell proteomics

Fast market access: Allows miniaturization of existing microplate-based protocols

Enable development of tomorrow's leading single cell (multi)omics protocols including a picture of each cell deposited



BIOCONVERGENCE IN INDUSTRIAL SOLUTIONS

1. POC/LATERAL FLOWS

Increased life expectancy, Early and efficient increased frequency of chronic disease e.g., cancer, heart disease and diagnosis can and diabetes will rise Global health expenditure expected to

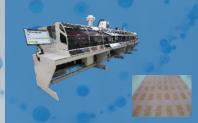
Around 50% of the population in the US are considered chronically ill and these patients account for about 85% of the total expenditure on healthcare services.

\$10T by 2022.



medical intervention prevent or delay most chronic diseases.

The health system pays greater attention to early, efficient intervention and preventative medicine.



Technological breakthroughs, innovative genetic and digital technologies can assist in identifying and contending with the complexities inherent in chronic diseases.



Help identify the "dormant" stage of diseases to preempt outbreak of symptoms. Health systems make the transition from a Volume Based Model to a Value Based Model.



M&A and tech development Strengthen bioconvergence

Technological innovation that can meet the new challenges and needs of the health system are required.





BIOCONVERGENCE IN INDUSTRIAL SOLUTIONS

M&A and tech development Strengthen bioconvergence

2. SINGLE CELL/HIGH PRECISION GENOMICS MARKET

The genomic revolution, the dramatic decline in the cost and increased speed of DNA sequencing alongside Artificial Intelligence and Big Data are today leading to the development of advanced diagnostic technologies that are based on genomics, proteomics and clinical data.

Two of the other fields developing alongside biotechnology are that of gene therapy and synthetic biology, based on the combination of innovative technologies such as DNA sequencing, creation and writing of new genes, among others by using CRISPR technology, behavioral modeling of specific genes, and precise measurement of gene behavior.



scWG-Sea







cellen ?NE

0 0 0 . .









FOCUS 2021

M&A and tech development Strengthen bioconvergence

1

Combined
SCIENION/Ginolis/
CELLENION offering for
early detection and
monitoring of diseases with
affordable diagnostics.
Offer complete solutions
for medical device
manufacturing and
drug delivery.

2

Launch the cellenCHIP and proteoCHIP.
Enter the single cell genomics and proteomics markets.

3

Focus on development of a unique cloud and bioinformatics offering for diagnostics and single cell analysis.

ISO 13485 certification

Asia expansion



RECORDED PRESENTATIONS

www.cellink.com/investors

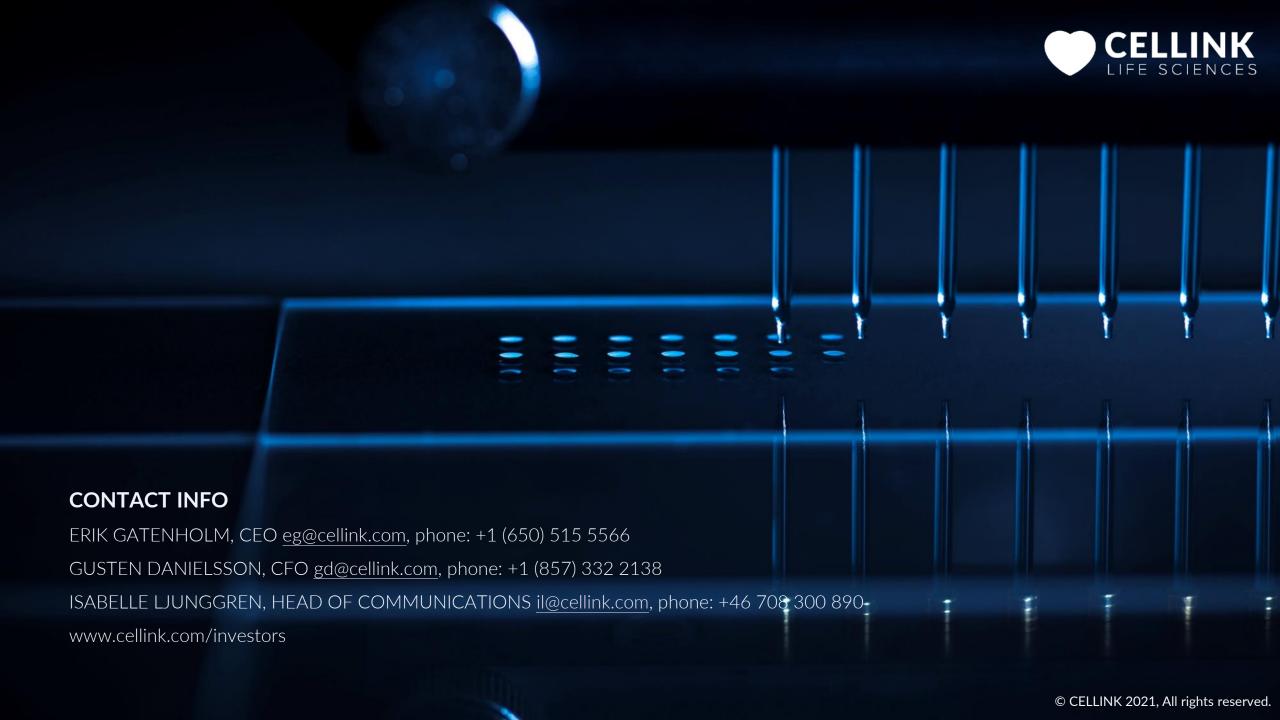














DISCLAIMER

You must read the following before continuing. The following applies to this document and the information provided in this presentation by CELLINK AB (publ) (the "Company") or any person on behalf of the Company and any other material distributed or statements made in connection with such presentation (the "Information"), and you are therefore advised to carefully read the statements below before reading, accessing or making any other use of the Information. In accessing the Information, you agree to be bound by the following terms and conditions.

The Information does not constitute or form part of, and should not be construed as, an offer of invitation to subscribe for, underwrite or otherwise acquire, any securities of the Company or a successor entity or any existing or future subsidiary or affiliate of the Company, nor should it or any part of it form the basis of, or be relied on in connection with, any contract to purchase or subscribe for any securities of the Company or any of such subsidiaries or affiliates nor shall it or any part of it form the basis of or be relied on in connection with any contract or commitment whatsoever. Specifically, this presentation does not constitute a "prospectus" within the meaning of the U.S. Securities Act of 1933, as amended.

The Information may not be reproduced, redistributed, published or passed on to any other person, directly or indirectly, in whole or in part, for any purpose. The Information is not directed to, or intended for distribution to or use by, any person or entity that is a citizen or resident of, or located in, any locality, state, country or other jurisdiction where such distribution or use would be contrary to law or regulation or which would require any registration or licensing within such jurisdiction. The Information is not for publication, release or distribution in the United States, the United Kingdom, Australia, Canada or Japan, or any other jurisdiction in which the distribution or release would be unlawful.

All of the Information herein has been prepared by the Company solely for use in this presentation. The Information contained in this presentation has not been independently verified. No representation, warranty or undertaking, expressed or implied, is made as to, and no reliance should be placed on, the fairness, accuracy, completeness or correctness of the Information or the opinions contained herein. The Information contained in this presentation should be considered in the context of the circumstances prevailing at that time and has not been, and will not be, updated to reflect material developments which may occur after the date of the presentation. The Company may alter, modify or otherwise change in any manner the content of this presentation, without obligation to notify any person of such revision or changes.

This presentation may contain certain forward-looking statements and forecasts which relate to events and depend on circumstances that will occur in the future and which, by their nature, will have an impact on the Company's operations, financial position and earnings. The terms "anticipates", "assumes", "believes", "can", "could", "estimates", "estimates", "forecasts", "forecasts", "intends", "may", "might", "plans", "should", "projects", "will", "would" or, in each case, their negative, or other variations or comparable terminology are used to identify forward-looking statements. There are a number of factors that could cause actual results and developments to differ materially from those expressed or implied in a factor that could cause these differences include, but are not limited to, implementation of the Company's strategy and its ability to further grow, risks associated with the development and of the Company's products, ongoing research and development, the ability to commercialize the Company's products, technology changes and new products in the Company's products, the impact of competition, changes in general economy and industry conditions and legislative, regulatory and political factors. While the Company always intends to express its best judgment when making statements about what it believes will occur in the future, and although the Company bases these statements on assumptions that it believe to be reasonable when made, these forward-looking statements are not a guarantee of its performance, and you should not place undue reliance on such statements. Forward-looking statements are subject to many risks, uncertainties and other variable circumstances. Such risks and uncertainties may cause the statements to be inaccurate and readers are cautioned not to place undue reliance on such statements. Many of these risks are outside of the Company's control and could cause its actual results to differ materially from those it thought would occur. The forward-looking statements to reflect future event