

Investor presentation

March 10-11, 2021

Erik Gatenholm, CEO, CELLINK Gusten Danielsson, CFO, CELLINK



BIOCONVERGENCE



The modern healthcare challenges



High costs and lengthy processes for drugs to reach patients. <u>9 out of 10</u> <u>fail in clinical stages!</u>



A life is lost <u>every hour</u> of the day due to lack of organ transplants



Animal studies are <u>poor</u> <u>indicators</u> of success for human drug development



Biology + Technology

BIOCONVERGENCE IS THE FUTURE OF HEALTHCARE



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



The solution is found in bioconvergence





Our unique bioconvergence offering





Large market opportunities for CELLINK



Disease and tissue modelling

Disease and tissue analysis

Disease and tissue diagnostics



The bioconvergence market

Bioconvergence market opportunity				
Bioprocessing	The upstream bioprocessing market size is expected to reach \$12.6Bn by 2026, at a CAGR of 12.9% during the period.	\$45Bn+		
Cell culture	The cell culture market is expected to reach \$33.1Bn by 2025, at a CAGR of 11.8% during the period.			
Single cell analy	The single-cell analysis market is expected to reach \$5.6Bn by 2025, at a CAGR of 17.8% during the period.	\$24Bn+		
3D Cell culture	The 3D cell culture market is expected to reach \$12.6Bn by 2026, expanding at a CAGR of 29.1% during the period.			
Cell line develop	Oment The cell line development market is expected to reach \$6.4Bn by 2025, at a CAGR of 10.8% during the period.			

LIFE SCIENCES



Serving our customers in the best possible way in the era of workflows



Drug screening workflow

Dispensing single cells to ensure specific genetic composition

Bioprint tissue using bioprinter and tissue specific bioink

Dispense drug molecules and compounds onto tissues Monitor the treated tissue for drug response

Annen Alas (1997) Frank Alas (1

NAME OF A





FINANCIAL SUMMARY

na:

1.04

12 🕮

18 27



Continued strong organic growth, positive result and strategic investments for the future

Q5: September – December 2020

- Net sales amounted to SEK 239,216 thousand (SEK 50,189 thousand), which corresponds to an increase of 377% (118%) compared to the corresponding period a year earlier, of which 73% (47%) was organic growth.
- **EBITDA** amounted to SEK 40,414 thousand (SEK -15,567 thousand), corresponding to a margin of 16.9% (-31.0%).
- **Profit for the period amounted to SEK 13,219 thousand** (SEK -18,895 thousand), which generates earnings per share after dilution of SEK 0.26 (SEK -0.48). The result was positively affected by the market valuation of the company's short-term investments of SEK 5,122 thousand (SEK 272 thousand).
- Rolling twelve-month net sales from consumables amounted to SEK 35,091 thousand (SEK 15,699 thousand), an increase of 124%. The share of total product sales was 11.9%, a decrease of 0.2 percentage points (12.1% in the comparison period).

September 2019 – December 2020

- Net sales amounted to SEK 416,009 thousand (SEK 155,646 thousand), which corresponds to an increase of 167% (128%) compared with the corresponding period a year earlier. Of the increase, **48%** (77%) was **organic growth**.
- **EBITDA amounted to SEK 816 thousand** (SEK -12,216 thousand), corresponding to a margin of 0.2% (-7.8%). Operating profit was affected by costs for the listing on Nasdaq Stockholm and acquisitions totaling SEK -18,264 thousand (SEK -6,395 thousand).
- **Profit for the period amounted to SEK -48,994 thousand** (SEK -18,314 thousand), which generates earnings per share after dilution of SEK -1.10 (SEK -0.51). In addition to the listing and acquisition costs, the net result was affected by the market valuation of the company's short-term investments of SEK -1,211 thousand (SEK 1,722 thousand).
- Given the company's current growth phase, which is expected to continue during 2021, the Board of Directors proposes no dividend for the financial year 2019/2020.

Strong organic growth and important strategic acquisition under continued challenging conditions

kSEK	Sep-Dec 2020	Sep-Dec 2019	Sep 2019-Dec 2020	Sep 2018-Dec 2019
Net sales*	239,216	50,189	416,009	155,646
Net sales Laboratory Solutions*	87,040	-	263,833	-
Net sales Industrial Solutions*	152,176	-	-	_ *
Gross profit	174,368	35,073	298,633	110,496
Gross margin, %	73%	70%	72%	71%
Operating profit before depreciation and amortization (EBITDA)	40,414	-15,567	816	-12,216
Operating margin before depreciation and amortization (EBITDA), %	16.9%	-31.0%	0.2%	-7.8%
Operating profit (EBIT)	14,161	-22,508	-51,927	-26,262
Operating margin (EBIT), %	5.9%	-44.8%	-12.5%	-16.9%
Profit for the period	13,219	-18,895	-48,994	-18,314
Diluted earnings per share, SEK**	0.26	-0.48	-1.10	-0.51
Net debt(-)/Net cash(+)	755,738	68,840	755,738	68,840
Cash flow from operating activities	-7,584	-13,847	-79,400	-29,665
Average number of shares**, ***	52,951,049	39,374,416	44,888,273	36,024,097
Number of shares at the end of the period**	51,601,285	38,984,776	51,601,285	38,984,776
Share price on closing day, SEK**	234.5	83.5	234.5	83.5
Market capitalization on closing day, MSEK	12,101	3,255	12,101	3,255
Number of employees at the end of the period	396	183	396	183

* Segment created September 2020, see note 3 in latest report

** CELLINK conducted a 4:1 split on January 10, 2020. Comparison periods have been restated for correct comparison.

For definitions, see latest report on www.cellink.com/investors

*** Average number of shares including potential ordinary shares.

CELLINK

CELLINK Strong organic growth in combination with improved sales levels in Q5, 2020... **NET SALES (LTM) MSEK** Organic 73% Growth Ouarter

FY2019/20







CELLINK

Long-term financial targets 2019-2022

Organic growth

EBITDA margin

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

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

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).

1997 - 19

With Lang Br.

4 10 10 17mmm 81

5-19 VI

Silve El mar m.

-

2000 1000:00

To Balland



101000

1 10 25

M&A AGENDA

AN LI

to a state country is

A 6 14

12 2

:18 12



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 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



Successful integration of acquired companies



- €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
- Acquired at 11x Revenue, now at ~1x revenue. (In 2 years)



- ~€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
- Acquired at ~7x Revenue, now at 3-5x revenue. (In 1.5 years)

SCIENION A CELLINK COMPANY

- ~€80 MM in transaction value
 50% / 50% in shares/cash
 - ~25% EBITDA margin
- Strengthen position in singlecell handling
- Creating a low volume dispensing powerhouse with Dispendix and a dominant player in single-cell dispensing with Cytena
- Acquired at ~3.7x Revenue, now at ~2.7x. (In 4 months)



- ~€70 MM in transaction value
 40% / 60% in shares/cash
 - ~12% EBITDA margin
- Strengthen position in automation and diagnostics
- Creating an automation and low-volume dispensing powerhouse with SCIENION and Business area Bioprinting
- Acquired at ~3.8x Revenue



Acquisition of Ginolis - main synergies

Enabling the next generation of bioprinting systems

The increased demand for microfluidic and lateral flow IVD tests, PoC-tests

Synergies with the Group's existing product offering and future product development

Increased capacity to deliver larger automation workflows



MatTek, a global leader in in-vitro technology and alternative drug testing models

"The rat gave us an answer, the monkeys gave us another answer, and the human testing gave us a third answer."

MatTek customer



Through the strategic acquisition of MatTek, the CELLINK Group will achieve the following:

Provide alternative testing models which enables the reduction, and in some cases elimination, of animal testing. These solutions allow researchers to gather better data through more physiologically relevant models and thus make better predictions.

Several strong synergies by combining cutting-edge bioprinting technology and modular large-scale robotic flows with 3D reconstructed, human-derived tissue models.



Acquisition is in line with CELLINK's commercial, bioconvergence strategy, complementing CELLINK's product offering and brings the Group closer to the patients through cutting edge products used in clinical and pre-clinical studies.



The transaction in brief

Enterprise Value on cash- and debt-free basis of \$68 million (20% in shares and remaining in cash).

MatTek's revenue reached \$16.6 million in 2020, with and EBITDA margin of 21.9 per cent. Pro-forma revenue growth is expected for 2021 and MatTek's historical growth rate has been in the range of 10 per cent.

MatTek will continue as a part of CELLINK's business area Bioprinting. MatTek will remain under current entity and management post-transaction.

The Acquisition's completion and the transfer of MatTek's shares are expected to take place by March 24, 2021 provided that all conditions for completion are met. MatTek will be consolidated in CELLINK's financial statements from second quarter 2021 and in the financial reporting from April 1, 2021.



Why increase tissue model testing?

Reduces animal testing
Faster drug development processes
Human in-vitro tissues are the best models of actual human tissues
Great potential for personalized medicine



Market potential and growth

In 2020 the global in vitro testing market without using animals as test models is estimated at \$9.1Bn and expected to grow at a CAGR of 10.3% during the forecast period to reach \$14.9Bn by 2025.





CONTACT INFO

Erik Gatenholm, CEO <u>eg@cellink.com</u>, phone: +1 (650) 515 5566 Gusten Danielsson, CFO <u>gd@cellink.com</u>, phone: +1 (857) 332 2138 Visit Investor Relations section for more information: www.cellink.com/investors Investor Relations service and requests please contact: <u>ir@cellink.com</u> Next interim report: Q1, 2021 – release on May 12, 2021, 08:00 (CET)



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 shall it or any part of it form the basis of or be relied on in connection with, 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", "expects", "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 forward-looking statement or affect the extent to which a particular projection is realised. Factors 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 potential market and industry, the ability to develop new 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 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 on a guarantee or a guarantee or a guarantee or a guarantee or the statements are outside of the Company's control and could cause its actual results on differ materially from those incurate 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 of any revisions to any of such statements to reflect future events or developme