InDex Pharmaceuticals Holding AB (publ)





First regulatory approvals to start the phase III study CONCLUDE

PERIOD APRIL-JUNE 2021

- Net sales amounted to SEK 0.0 (0.0) million
- Operating loss amounted to SEK –42.6 (–15.6) million
- Result after tax amounted to SEK –42.6 (–15.6) million, corresponding to SEK –0.08 per share (–0.07) before and after dilution
- Cash flow from operating activities amounted to SEK –53.4 (–32.8) million

SIGNIFICANT EVENTS DURING THE QUARTER

InDex announced that patient recruitment for the phase
 III study CONCLUDE is planned to initiate after the summer

SIGNIFICANT EVENTS AFTER THE QUARTER

- InDex received first regulatory approval from the Swedish MPA to start the phase III study CONCLUDE with cobitolimod
- InDex received FDA clearance to start the phase III study CONCLUDE

All comparative amounts in brackets refer to the outcome during the corresponding period 2020.

PERIOD JANUARY-JUNE 2021

- Net sales amounted to SEK 0.0 (0.0) million
- Operating loss amounted to SEK –51.9 (–39.6) million
- Result after tax amounted to SEK –52.0 (–39.6) million, corresponding to SEK –0.11 per share (–0.17) before and after dilution
- Cash flow from operating activities amounted to SEK –62.0 (–54.7) million
- Cash and cash equivalents at the end of the period amounted to SEK 478.8 (70.6) million
- Number of employees at the end of the period was 7 (7)
- Number of shares at the end of the period was 532,687,650

OTHER EVENTS

- InDex got patent for additional DIMS compounds granted in Europe
- The annual general meeting in InDex Pharmaceuticals
 Holding AB was held on June 3, 2021. Board members
 Wenche Rolfsen (also chairman), Marlene Forsell, Uli
 Hacksell and Lennart Hansson were re-elected. Yilmaz
 Mahshid and Stig Løkke Pedersen had ahead of the annual
 general meeting declined re-election

"We will initiate patient recruitment when the clinics are up and running again after the summer as the healthcare situation has started to normalise", says Peter Zerhouni, CEO of InDex Pharmaceuticals.

InDex is a pharmaceutical development company focusing on immunological diseases where there is a high unmet medical need for new treatment options. The company's lead asset is the drug candidate cobitolimod, which is in late stage clinical development for the treatment of moderate to severe ulcerative colitis – a debilitating, chronic inflammation of the large intestine. InDex has also developed a platform of patent protected discovery stage substances, so called DNA based ImmunoModulatory Sequences (DIMS), with the potential to be used in the treatment of various immunological diseases. InDex is based in Stockholm, Sweden. The company's shares (ticker INDEX) are traded on Nasdaq First North Growth Market Stockholm. Redeye AB is the company's Certified Adviser (+46 8 121 576 90 or certifiedadviser@redeye.se).

CEO statement



We are now close to the start of the phase III study CONCLUDE to evaluate cobitolimod as a new treatment for patients with moderate to severe left-sided ulcerative colitis. It is a global clinical study which will include 440 participants and be conducted at several hundred clinics in over 30 countries. The process of applying for and obtaining approval from the relevant authorities in the participating countries is ongoing, and in July we received the first regulatory approval to start the study from the Swedish MPA and yesterday from the US FDA.

We will initiate patient recruitment when the clinics are up and running again after the summer as the healthcare situation has started to normalise. Together with our CRO we are finalising the practical preparations.

To strengthen our clinical development organisation, we have hired two new employees, Anders Bröijersén as Senior Medical Director Clinical Operations and Johan Levin as Project Manager Clinical Operations. Both have long and broad experience of clinical development and conducting clinical studies. I am pleased that we have been able to attract two such qualified employees and welcome Anders and Johan to the team. Much of their time will be devoted to support the clinics in the study as ambassadors for InDex and the CONCLUDE study, which was a successful model for patient recruitment in our previous study CONDUCT.

In parallel with the phase III study, we will conduct a smaller clinical pharmacokinetic study (PK study) with cobitolimod in Sweden. We have received approval from the Swedish MPA to start also this study. The purpose of the

study is to evaluate the systemic uptake of cobitolimod, and the data will support future regulatory applications for market approval of cobitolimod. Previous preclinical and clinical studies have shown that the systemic uptake of cobitolimod is very limited, which likely contributes to the excellent safety profile. This is a significant advantage compared to the current systemically administered drugs for ulcerative colitis that can cause severe off-target effects.

Besides cobitolimod for ulcerative colitis, InDex has a preclinical portfolio of more than 150 DNA-based Immuno-Modulatory Sequences (DIMS), of which we are testing a selected number of candidates in models of other inflammatory diseases. In April, a new European patent was granted that provides protection for 19 different DIMS substances. This patent has previously been granted in the US and Canada, and we are very pleased that also the European Patent Office confirms the novelty of our DIMS platform.

In June, we had the Annual General Meeting and although we had postponed it as much as possible, we were unfortunately not able to welcome the shareholders to a physical meeting this year either. On September 2, I will present the company at the Pareto Securities Healthcare Conference, which will be held virtually also this year.

Soon I hope we can start meeting each other in person again and I look forward to welcoming the first patient in the phase III study CONCLUDE.

Business overview

INTRODUCTION

InDex is a pharmaceutical development company focusing on immunological diseases where there is a high unmet medical need for new treatment options. The company's lead asset is the drug candidate cobitolimod, which is in late stage clinical development for the treatment of moderate to severe ulcerative colitis – a debilitating, chronic inflammation of the large intestine. InDex has also developed a platform of patent protected discovery stage substances, so called DNA based Immuno-Modulatory Sequences (DIMS), with the potential to be used in the treatment of various immunological diseases. InDex is based in Stockholm, Sweden. The company's shares are traded on Nasdaq First North Growth Market Stockholm.

COBITOLIMOD

Cobitolimod is a potential new medication for patients with moderate to severe ulcerative colitis. Ulcerative colitis is a chronic disease caused by inflammation of the colon. Today, about two million people in Europe and the United States suffer from ulcerative colitis, a disease that has a major impact on the patient's quality of life. Ulcerative colitis is characterised by blood- and mucus-mixed diarrhea, frequent stools, pain, fever, weight loss, and anemia. Patients also have a significant elevated risk of developing colon cancer. Most commonly, ulcerative colitis debuts between 15 and 30 years of age and most patients require lifelong medication. Despite the currently available drugs on the market, many patients with ulcerative colitis still suffer from severe symptoms, and current therapies can cause serious side effects. For those patients that do not respond to medical treatment, the last resort is to surgically remove the colon.

Cobitolimod is a local treatment with a novel mechanism of action. It is a so-called Toll-like receptor 9 (TLR9) agonist that can provide an anti-inflammatory effect locally in the large intestine, which may induce mucosal healing and relief of the clinical symptoms in ulcerative colitis. Cobitolimod is administered via the rectum using an enema allowing a rapid onset of action without systemic exposure and off-target effects.

Cobitolimod met the primary endpoint in the phase IIb study CONDUCT and demonstrated an outstanding combination of efficacy and safety. Data from four previous completed placebo-controlled clinical trials support the efficacy and safety demonstrated in the CONDUCT study. Given the outstanding combination of efficacy and safety, InDex is now advancing cobitolimod into phase III, which is the final stage of development before application for market approval.

Based on the sales of recently launched products, as well as the company's proprietary market research and analyses, including the addressable market described above, the annual global peak sales at a successful commercialisation of cobitolimod are estimated by the company to have the potential to reach more than USD 1 billion.

COMPETITIVE EFFICACY EXCELLENT SAFETY NOVEL MECHANISM OF ACTION LOCAL ADMINISTRATION AND LOW DOSING FREQUENCY

Illustrations: Freepik

POSSIBLITY OF COMBINING

WITH OTHER THERAPIES

Phase III study - CONCLUDE

Based on regulatory guidance InDex is planning a sequential phase III program with two induction studies and a one-year maintenance study with patients that have responded to cobitolimod as induction therapy.

The important initial induction study CONCLUDE will include 440 patients and the company estimates that it will take 18 to 24 months to complete from initiation. CONCLUDE is a randomised, double-blind, placebo-controlled, global phase III study to evaluate cobitolimod as a novel treatment for patients with moderate to severe left-sided ulcerative colitis. The primary endpoint will be clinical remission at week 6. Apart from the dosing 250 mg x 2, which was the highest dose and the one that showed the best efficacy in the phase IIb study CONDUCT, the phase III study will also evaluate a higher dose, 500 mg x 2, in an adaptive study design. This higher dose has the potential to provide an even better efficacy than what was observed in the phase IIb study.

When a sufficient number of the participants in the study have been randomised and have eligible data for the primary endpoint, an interim analysis will be performed in a blinded fashion to select the best dose of cobitolimod and the other dose will be dropped. Following the blinded interim analysis, the additional patients to be randomised into the study will receive only the best dose of cobitolimod or placebo. Patients responding to cobitolimod in the induction study will be eligible to continue in a one-year maintenance study, where they will be treated with either cobitolimod or placebo.

InDex has entered into an agreement for services with the leading global clinical research organization (CRO) Parexel Biotech for the phase III study CONCLUDE. Parexel Biotech has considerable experience managing phase III studies in inflammatory bowel disease. Parexel Biotech was the CRO that InDex successfully collaborated with in the phase IIb study CONDUCT.

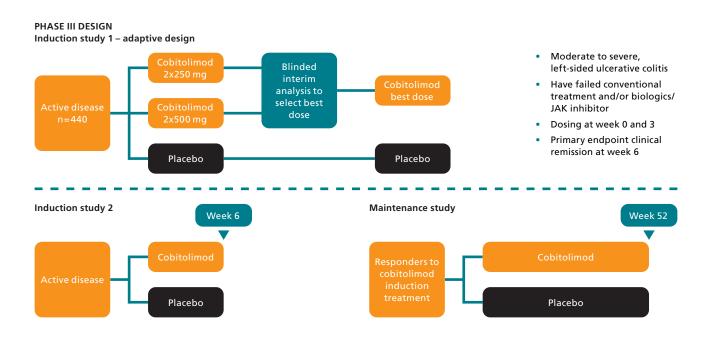
Oral formulation of cobitolimod

InDex has developed a prototype of a novel formulation of its lead drug candidate cobitolimod for oral administration, with targeted drug substance release or delivery to the lower part of the gastrointestinal tract and thus again avoiding systemic exposure. The capsule is a potential follow-on product to the current topical formulation. An oral therapy makes it possible to deliver cobitolimod to parts of the gastrointestinal tract which are inaccessible to an enema and could be more convenient for patients.

This opens the possibility to broaden the therapeutic use of cobitolimod to also include pancolitis and Crohn's disease, where the inflammation can be located higher up in the gastrointestinal tract. The oral formulation development also provides the opportunity to secure additional patent protection for cobitolimod.

OTHER DIMS

InDex has, besides cobitolimod, a preclinical portfolio of more than 150 DNA-based ImmunoModulatory Sequences (DIMS). The DIMS candidates are oligonucleotides that differ in sequence and length but are all TLR9 agonists. DIMS mimic bacterial DNA, without being harmful, and stimulate immune cells to produce beneficial anti-inflammatory cytokines that will help to dampen inflammation. This opens opportunities for the treatment of different inflammatory conditions, in which the immune responses are imbalanced. To capitalise on the substantial historical investments in the DIMS portfolio and to take advantage of the expertise and experience built up during the development of cobitolimod in ulcerative colitis, InDex is testing a selected number of DIMS candidates in models of other inflammatory diseases. Positive signals have been observed, and InDex is now confirming these early results with alternative and complementary methods in order to be able to select a DIMS substance for further development.



SIGNIFICANT EVENTS DURING THE QUARTER

 InDex announced on May 31, 2021 that patient recruitment for the phase III study CONCLUDE is planned to initiate after the summer. The study will evaluate the efficacy and safety of the drug candidate cobitolimod for the treatment of moderate to severe left-sided ulcerative colitis.

SIGNIFICANT EVENTS AFTER THE QUARTER

- InDex announced on July 21, 2021 that that the Swedish Medical Products Agency (MPA) has given approval to start the phase III clinical study CONCLUDE in Sweden.
- InDex announced on August 24, 2021 that the U.S. Food and Drug Administration (FDA) has given clearance to start the phase III clinical study CONCLUDE with cobitolimod in the United States.

OTHER EVENTS

- InDex announced on April 7, 2021 that a patent covering 19 compounds from the company's DIMS platform has been granted by the European Patent Office. The new European patent covers both the composition-of-matter and method-of-use of 19 different DIMS compounds for the treatment of inflammatory diseases, cancer and infectious diseases. The patent provides an exclusivity period until December 2031, with the possibility of up to 5 years term extension after market approval.
- The annual general meeting in InDex Pharmaceuticals Holding AB was held on June 3, 2021. Board members Wenche Rolfsen (also chairman), Marlene Forsell, Uli Hacksell and Lennart Hansson were re-elected. Yilmaz Mahshid and Stig Løkke Pedersen had ahead of the annual general meeting declined re-election. The annual general meeting also resolved, in accordance with the Board's proposal, on the implementation of a long term incentive program in the form of employee stock options to senior executives and other key persons of the group.



Financial overview

FINANCIAL SUMMARY FOR THE GROUP

Because of the nature of the business operations, there may be large fluctuations between different periods.

FINANCIAL DEVELOPMENT DURING APRIL-JUNE 2021

Net sales for the period April to June 2021 amounted to SEK 0.0 (0.0) million. The net sales previous year were related to the sale of DiBiCol test kits up to September 30, 2020. Sale of DiBiCol test kits was then terminated.

Other operating income SEK 0.1 (0.0) million refers to grant received from Vinnova.

Operating expenses for the period amounted to SEK 42.7 (15.6) million. The increase is attributable to higher costs for phase III preparations compared to the costs previous year related to manufacturing of study drug.

The operating expenses during the period refer primarly to costs for phase III preparations and general operating expenses.

Costs for the personnel during the reporting period amounted to SEK 2.9 (2.3) million. The increase is partly related to general salary increases and costs for recruitment of personnel.

Other operating expenses include foreign exchange losses of SEK 8.4 (0) million related to cash and cash equivalents in foreign currency. InDex purchased during the second quarter 2021 USD to be used for future payments related to signed contracts denominated in USD.

FINANCIAL DEVELOPMENT DURING JANUARY-JUNE 2021

Net sales for the period January to June 2021 amounted to SEK 0.0 (0.0) million. The net sales previous year were related to the sale of DiBiCol test kits up to September 30, 2020. Sale of DiBiCol test kits was then terminated.

Other operating income SEK 0.2 (0.0) million refers to grant received from Vinnova.

Operating expenses for the period amounted to SEK 52.1 (39.6) million. The increase is attributable to higher costs for phase III preparations compared to the costs previous year related to manufacturing of study drug.

The operating expenses during the period refer primarly to costs for phase III preparations and general operating expenses.

Costs for the personnel during the reporting period amounted to SEK 5.5 (4.5) million. The increase is partly related to general salary increases and costs for recruitment of personnel.

Other operating expenses include foreign exchange losses of SEK 8.4 (0) million related to cash and cash equivalents in foreign currency. InDex purchased during the second quarter 2021 USD to be used for future payments related to signed contracts denominated in USD.

Cash and cash equivalents as of June 30, 2021 amounted to SEK 478.8 million, which is SEK 425.0 million higher than as of December 31, 2020. The Swedish Companies Registration Office recorded the completed rights issue of 443,906,375 new shares on February 11, 2021. The subscription price was set to SEK 1.20. InDex received approximately SEK 488 million after deduction of the transaction related costs for financial and legal services and for costs for registration and practical management.

FINANCIAL SUMMARY AFTER THE REPORTING PERIOD

The covid-19 pandemic affects the healthcare systems and the investor sentiment globally and must be taken into account in the company's strategic planning. The Board, however, assess that there is no impact on the company's financial position as of June 30, 2021 due to events after the reporting period.

EXPECTED FUTURE DEVELOPMENT

The Board is reviewing the forecasted cash flow on an ongoing basis to determine InDex's capital requirements and resources required to conduct the business activities in accordance with the strategic direction decided by the Board. It is the assessment of the Board that InDex has enough capital to finance all financial commitments InDex has for the coming 12-month period. InDex provides no financial forecast or similar forward-looking statement.

PARENT COMPANY

The net sales amounted to SEK 5.7 (5.8) million during the period January to June 2021 and consisted of invoicing of group wide expenses to InDex Pharmaceuticals AB.

The operating expenses amounted to SEK 8.7 (8.4) million and consisted of personnel expenses and other operating expenses relating to the administration of InDex.

FINANCIAL SUMMARY							
SEK million	Apr-Jun 2021	Apr-Jun 2020	Jan-Jun 2021	Jan-Jun 2020	Full year 2020		
Revenues	0.0	0.0	0.0	0.0	0.0		
Operating result	-42.6	-15.6	-51.9	-39.6	-57,3		
Result after tax	-42.6	-15.6	-52.0	-39.6	-57.4		
Earnings per share before and after dilution, SEK	-0.08	-0.07	-0.11	-0.17	-0.24		
Cash flow from operating activities	-53.4	-32.8	-62.0	-54.7	-70.7		
Cash and cash equivalents at the end of the period	478.8	70.6	478.8	70.6	53.8		

Note: Earnings per share – Net result divided by weighted number of shares (adjusted for the completed rights issue in February 2021).

Other information

EMPLOYEES

The number of employees at the end of the period was 7 (7).

THE SHARE

Nordnet Pensionsförsäkring

Swedbank försäkring AB

S-E-Bankens Utvecklingsstiftelse

Staffan Rasjö

Originat AB

LTIP 2020

Other

Total

The share is listed on Nasdaq First North Growth Market Stockholm since October 11, 2016.

LARGEST SHAREHOLDERS PER JUNE 30, 2021							
	Number of shares	Percentage of capital and votes, %					
Linc AB	69,920,567	13.1					
HBM Healthcare Investments	52,916,667	9.9					
Fjärde AP-fonden	52,314,074	9.8					
Handelsbanken Funds	25,000,000	4.7					
Avanza Pension	23,764,477	4.5					
SEB-Stiftelsen	19,047,617	3.6					
SEB Life International	13,927,350	2.6					
Bengt Thornberg, dödsbo	13,417,394	2.5					
SEB Venture Capital	12,994,367	2.4					
Stiftelsen Industrifonden	12,865,296	2.4					

11,379,525

10,318,953

10,000,000

7,000,000

6,074,144

191,747,219

532.687.650

INCENTIVE PROGRAMMES

At the annual general meeting held on April 20, 2020 it was resolved to issue 3,965,000 warrants to transfer to employees and other key persons within InDex. The warrants had an exercise price of SEK 20 per share and can be exercised during May-October 2023. The Board allocated in July 2020 958,388 warrants to employees and other key persons that were purchased for SEK 0.2522 per warrant. A total of 13 employees and other key persons were offered to subscribe for warrants and 12 of these individuals subscribed for their full allotment.

After the completed rights issue in February 2021 the exercise price and the number of shares that each warrant represents have been recalculated in accordance with the applicable terms. The new exercise price amounts to SEK 7.804 and each warrant entitles the holder to subscribe for 2.5627 shares. The remaining warrants have been terminated.

LTIP 2021

At the annual general meeting held on June 3, 2021 it was resolved to issue 7,200,000 employee stock options to transfer to employees and other key persons within InDex. In addition, 2,262,240 warrants were issued to cover potential cash flow effects from social security costs arising from allotted employee stock options. The options have a strike price of SEK 4 per share and can be exercised during July-December 2024. In July 2021 the Board allocated 5,731,800 options to employees and other key persons free of charge. A total of 13 employees

and other key persons were offered and subsequently subscribed for their allotted employee stock options.

LTIP 2021 will be accounted for in accordance with *IFRS 2* – *Share-based payments*. IFRS 2 stipulates that the employee stock options should be expensed as personnel costs over the vesting period and will be accounted for directly against equity. Personnel costs in accordance with IFRS 2 do not affect the company's cash flow. Social security costs will in accordance with UFR 7 be expensed in the income statement during the vesting period.

REVIEW BY THE AUDITOR

This interim report has not been reviewed by the company's auditor.

FINANCIAL CALENDER

Interim report Q3 November 24, 2021

Stockholm, August 25, 2021 Peter Zerhouni, CEO

FOR MORE INFORMATION, PLEASE CONTACT:

Peter Zerhouni, CEO

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The information in this interim report is information that InDex Pharmaceuticals Holding AB (publ) is obliged to make public pursuant to the EU Market Abuse Regulation. The information was submitted for publication by the contact person stated above on August 25, 2021 at 8:00 CET

This is an English translation of the Swedish interim report. In case of discrepancies between the English translation and the Swedish report, the Swedish report shall prevail.

Condensed consolidated statement of total comprehensive income

	Apr 1-Jun 30,	Apr 1-Jun 30,	Jan 1-Jun 30,	Jan 1-Jun 30,	Full year
SEKk	2021	2020	2021	2020	2020
Revenues					
Net sales	0	9	0	26	35
Other operating income	121	-	156	-	380
Total revenues	121	9	156	26	415
Operating expenses					
Raw material and consumables	-491	-6,726	-825	-14,937	-16,021
Other external expenses	-30,627	-6,208	-36,792	-19,610	-30,990
Personnel costs	-2,937	-2,308	-5,476	-4,489	-9,561
Depreciations/amortisations of tangible fixed assets and right-of-use assets	-317	-338	-634	-573	-1,192
Other operating expenses	-8,368	-	-8,368	-	-
Total expenses	-42,740	-15,580	-52,095	-39,609	-57,764
Operating loss	-42,619	-15,571	-51,939	-39,583	-57,349
Result from financial investments					
Financial income	_	_	_	-	46
Financial expenses	-26	-19	-54	-26	-115
Financial items – net	-26	-19	-54	-26	-69
Earnings before tax	-42,645	-15,590	-51,993	-39,609	-57,418
Taxes for the period	_	-	-	-	-
LOSS FOR THE PERIOD	-42,645	-15,590	-51,993	-39,609	-57,418

Earnings per share, based on the net result attributable to the shareholders of the parent company:

SEK	Note	Apr 1-Jun 30, 2021	Apr 1-Jun 30, 2020	Jan 1-Jun 30, 2021	Jan 1-Jun 30, 2020	Full year 2020
Earnings per share before and after dilution *	6	-0.08	-0.07	-0.11	-0.17	-0.24

^{*} Adjusted for the completed rights issue in February 2021.

In the group there are no items reported in other comprehensive income. So total comprehensive income is consistent with profit/loss for the period. The profit/loss for the period and total comprehensive income are entirely attributable to the equity holders of the parent company.

Condensed consolidated balance sheet

<u>SEKk</u>	Jun 30, 2021	Jun 30, 2020	Dec 31, 2020
ASSETS			
Fixed assets			
Tangible fixed assets			
Equipment, tools and installations	720	872	818
Total tangible fixed assets	720	872	818
Total tangine fixed assets	720	072	010
Right-of-use assets	2,056	3,129	2,593
· ·			
Financial assets			
Other financial assets	1	1	1
Total financial assets	1	1	1
Total fixed assets	2,777	4,002	3,412
Current assets			
Current receivables			
Accounts receivable	_	_	_
Other current receivables	2,010	1,819	907
Prepaid expenses and accrued income	16,719	671	3,031
Cash and cash equivalents	478,792	70,601	53,834
Total current receivables	497,521	73,091	57,772
Total current assets	497,521	73,091	57,772
TOTAL ASSETS	500,298	77,093	61,184
EQUITY AND LIABILITIES			
Equity			
Share capital	10,654	1,776	1,776
Additional paid-in capital	863,174	384,314	384,557
Retained earnings (including profit/loss for the period)	-388,998	-319,196	-337,005
Total equity attributable to the shareholders of the parent company	484,830	66,894	49,328
Liabilities			
Non-current liabilities			
Non-current lease liabilities	1,305	2,110	1,578
Total non-current liabilities	1,305	2,110	1,578
	.,505	2,	.,570
Current liabilities			
Current lease liabilities	516	1,028	763
Account payables	9,127	2,593	3,023
Other current liabilities	1,132	1,173	852
Accrued expenses and deferred income	3,389	3,295	5,640
Total current liabilities	14,163	8,089	10,278
Total liabilities	15,467	10,199	11,856
TOTAL EQUITY AND LIABILITIES	500,298	77,093	61,184

Condensed consolidated statement of changes in equity

Closing balance, June 30, 2021	10,654	863,174	-388,998	484,83
Total transactions with shareholders of the parent company	8,878	478,617	-	487,49
Transaction costs	_	-45,192	-	-45,19
Fransactions with shareholders of the parent company: ssue of shares	8,878	523,809	_	532,68
Fotal comprehensive income for the year	-	-	-51,993	-51,99
Profit/loss for the period equal to total comprehensive income	_	_	-51,993	-51,99
Opening balance, January 1, 2021	1,776	384,557	-337,005	49,32
Closing balance, December 31, 2020	1,776	384,557	-337,005	49,32
Total transactions with shareholders of the parent company	-	243	-	24
Transactions with shareholders of the parent company: Issue of warrants	-	243		24
Total comprehensive income for the year	-	-	-57,418	-57,41
Profit/loss for the period equal to total comprehensive income	-	_	-57,418	-57,41
Opening balance, January 1, 2020	1,776	384,314	-279,587	106,50
Closing balance, June 30, 2020	1,776	384,314	-319,196	66,89
Total comprehensive income for the year	-	-	-39,609	-39,60
Profit/loss for the period equal to total comprehensive income	-	-	-39,609	-39,60
Opening balance, January 1, 2020	1,776	384,314	-279,587	106,50
SEKk	Share capital	Additional paid in capital	Retained earnings, including loss for the period	Total equi
	Equity attribut	holders of the parer	nt company	

Condensed consolidated cash flow

	Apr 1-Jun 30,	Apr 1-Jun 30,	Jan 1-Jun 30,	Jan 1-Jun 30,	Full year
SEKk	2021	2020	2021	2020	2020
Operating activities					
Operating result	-42,619	-15,571	-51,939	-39,583	-57,349
Adjustments for non-cash items:					
Depreciations/amortisations	317	338	634	573	1,192
Interest paid and received	_	-19	_	-26	-70
Income tax paid	-	-	-	-	-
Cash flow from operating activities before changes in working capital	-42,302	-15,252	-51,305	-39,036	-56,227
Changes in working capital					
Decrease/Increase of current receivables	-15,795	-40	-14,791	-669	-2,117
Decrease/Increase of current liabilities	4,669	-17,510	4,133	-15,039	-12,306
Cash flow from changes in working capital	-11,126	-17,550	-10,658	-15,708	-14,423
Cash flow from operating activities	-53,428	-32,802	-61,963	-54,744	-70,650
Investing activities					
Investments in lease assets	_	-3,218	_	-3,218	_
Investments in tangible assets	_	-881	_	-881	-909
Cash flow from investing activities	-	-4,099	-	-4,099	-909
Financing activities					
Amortisation of lease liabilities	-287	-243	-574	-484	-1,639
Lease liability contracted	_	3,138	_	3,138	
Issues of shares, net after transaction costs	_	_	487,495	_	-
Issue of warrants	-	-	-	_	242
Cash flow from financing activities	-287	2,895	486,921	2,654	-1,397
Cash flow for the period	-53,715	-34,006	424,958	-56,189	-72,956
Decrease/increase of cash and cash equivalents					
Cash and cash equivalents at the beginning of the period	532,507	104,607	53,834	126,790	126,790
Currency translation difference in cash and cash equivalents	_	-	_	_	_
Cash and cash equivalents at the end of the period	478,792	70,601	478,792	70,601	53,834

Statement of comprehensive income for the parent company

SEKk	Apr 1-Jun 30, 2021	Apr 1-Jun 30, 2020	Jan 1-Jun 30, 2021	Jan 1-Jun 30, 2020	Full year 2020
Revenues					
Net sales	2,184	2,911	5,661	5,843	11,265
Total revenues	2,184	2,911	5,661	5,843	11,265
Operating expenses					
Other external expenses	-2,592	-3,046	-5,370	-5,648	-11,486
Personnel costs	-1,785	-1,400	-3,273	-2,712	-5,754
Depreciations/amortisations of tangible fixed assets and right-of-use assets	-49	-15	-98	-15	-91
Total expenses	-4,426	-4,461	-8,741	-8,375	-17,330
Operating loss	-2,242	-1,550	-3,080	-2,532	-6,065
Net financial items					
Write-down of financial assets	-100,000	-50,000	-100,000	-50,000	-50,000
Financial income	-	-	-	_	46
Financial costs	-	- 7	-	- 7	-6
Total net financial items	-100,000	-50,007	-100,000	-50,007	-49,960
Profit or loss before tax	-102,242	-51,557	-103,080	-52,539	-56,025
Taxes for the period	-	-	-	_	-
PROFIT OR LOSS FOR THE PERIOD	-102,242	-51,557	-103,080	-52,539	-56,025

In the parent company there are no items reported in other comprehensive income. So total comprehensive income is consistent with profit/loss for the period.

Balance sheet for the parent company

SEKK	Jun 30, 2021	Jun 30, 2020	Dec 31, 2020
ACCETC		·	· · · · · ·
ASSETS			
Fixed assets			
Tangible fixed assets			
Equipment, tools and installations	720	866	818
Total tangible fixed assets	720	866	818
Financial assets			
Shares in subsidiary	247,030	247,030	247,030
Total financial assets	247,030	247,030	247,030
Total fixed assets	247,750	247,896	247,848
Total fixed assets	247,730	247,630	247,040
Current assets			
Current receivables			
Intercompany receivables	286,036	675	779
Other receivables	1,125	633	219
Prepaid expenses and accrued income	521	571	1,247
Total current receivables	287,682	1,879	2,245
Cash and cash equivalents	114,107	69,816	45,491
Total current assets	401,789	71,696	47,736
TOTAL ASSETS	649,539	319,591	295,584
EQUITY AND LIABILITIES			
Equity			
Restricted equity			
Share capital	10,654	1,776	1,776
Total restricted equity	10,654	1,776	1,776
Non-restricted equity			
Share premium reserve	1,108,891	630,031	630,274
Retained earnings	-369,014	-312,989	-312,989
Profit or loss for the period	-103,080	-52,539	-56,025
Total non-restricted equity	636,796	264,503	261,260
Total equity	647,451	266,279	263,036
Liabilities			
Current liabilities			
Accounts payable	428	1,161	1,114
Intercompany liabilities	-	50,098	28,800
Other liabilities	264	534	323
Accrued expenses and deferred income	1,396	1,519	2,311
Total current liabilities	2,088	53,312	32,548
TOTAL EQUITY AND LIABILITIES	649,539	319,591	295,584

Statement of change in equity parent company

	Restricted equity	N	on-restricted equit	у	
SEKK	Share capital	Share premium	Retained earnings	Net result	Total equity
	· · · · · · · · · · · · · · · · · · ·	-			
Opening balance, January 1, 2020	1,776	630,031	-217,005	-95,984	318,818
Disposition of last year's result	-	-	-95,984	95,984	-
Net results and total comprehensive income for the year	_	-	_	-52,539	-52,539
Total comprehensive income for the year	-	-	-	-52,539	-52,539
Closing balance, June 30, 2020	1,776	630,031	-312,989	-52,539	266,279
Opening balance, January 1, 2020	1,776	630,031	-217,005	-95,984	318,818
Disposition of last year's result	-	-	-95,984	95,984	-
Net results and total comprehensive income for the year	_	_	_	-56,025	-56,025
Total comprehensive income for the year	_	-	-	-56,025	-56,025
Transactions with shareholders of the parent company:					
Issue of warrants	_	243	_	_	243
Total transactions with shareholders of the parent company	-	243	-	-	243
Closing balance, December 31, 2020	1,776	630,274	-312,989	-56,025	263,036
Opening balance, January 1, 2021	1,776	630,274	-312,989	-56,025	263,036
Disposition of last year's result	-	-	-56,025	56,025	-
Net results and total comprehensive income for the year	-	-	-	-103,080	-103,080
Total comprehensive income for the year	-	-	-	-103,080	-103,080
Transactions with shareholders of the parent company:					
Issue of shares	8,878	523,809	-	-	532,687
Transaction costs	-	-45,192	-	-	-45,192
Total transactions with shareholders of the parent company	8,878	478,617	-	-	487,495
Closing balance, June 30, 2021	10,654	1,108,891	-369,014	-103,080	647,451

Statement of cash flow for the parent company

SEKk	Apr 1-Jun 30, 2021	Apr 1-Jun 30, 2020	Jan 1-Jun 30, 2021	Jan 1-Jun 30, 2020	Full year 2020
Operating activities					
Profit or loss before tax	-102,242	-51,557	-103,080	-52,539	-56,025
Adjustments for non-cash items:	102,212	31,337	103,000	32,333	30,023
Write downs	100,000	50,000	100,000	50,000	50,000
Income tax paid	_	-	_	_	-
Depreciations/amortisations	49	15	98	15	91
Cash flow from operating activities before changes in working capital	-2,193	-1,542	-2,982	-2,524	-5,934
Changes in working capital					
Changes in current receivables	60,609	-299	-285,438	-892	-1,258
Changes in current liabilities	-3,155	21,477	-30,459	-852	-21,616
Cash flow from changes in working capital	57,454	21,178	-315,897	-1,744	-22,874
Cash flow from operating activities	55,261	19,636	-318,879	-4,268	-28,808
Investing activities					
Shareholder's contribution	-100,000	-50,000	-100,000	-50,000	-50,000
Investment of leases	_	-881	_	-881	-909
Cash flow from investing activities	-100,000	-50,881	-100,000	-50,881	-50,909
Financing activities					
Issues of shares, net after transaction costs	_	_	487,495	_	_
Issue of warrants	-	_	_	_	243
Cash flow from financing activities	-	-	487,495	-	243
Cash flow for the period	-44,739	-31,245	68,616	-55,149	-79,474
Decrease/increase in cash and cash equivalents					
Cash and cash equivalents at the beginning of the period	158,846	101,061	45,491	124,965	124,965
Cash and cash equivalents at the end of the period	114,107	69,816	114,107	69,816	45,491

Development of parent company's share capital

SEK		Change in	Total	Number of	Total number	Paid in
Date	Transaction	share capital	share capital	new shares	of shares	amount
Jun 27, 2016	Inception of the company	500,000	500,000	500,000	500,000	500,000
Sep 7, 2016	Split of shares	_	500,000	45,500,000	50,000,000	-
Sep 7, 2016	Share issue in-kind	601,345	1,101,345	60,134,466	110,134,466	_
Sep 7, 2016	Reduction of number of shares	-500,000	601,345	-50,000,000	60,134,466	-
Sep 7, 2016	Share issue	_	601,345	2	60,134,468	-
Sep 8, 2016	Reversed split of shares	_	601,345	-30,067,234	30,067,234	-
Oct 6, 2016	Share issue for pref. shares	52,685	654,030	2,634,279	32,701,513	52,685
Oct 6, 2016	Share issue	560,479	1,214,509	28,023,969	60,725,482	235,401,340
Oct 12, 2016	Share issue	14,305	1,228,814	715,250	61,440,732	6,008,100
Oct 25, 2016	Share issue	17,969	1,246,783	898,421	62,339,153	7,546,736
Nov 14, 2016	Share issue	1,895	1,248,678	94,725	62,433,878	795,690
Dec 29, 2016	Share issue in-kind	1,300	1,249,978	65,015	62,498,893	_
Jan 13, 2017	Share issue	591	1,250,569	29,540	62,528,433	248,136
Oct 23, 2018	Share issue	125,057	1,375,626	6,252,842	68,781,275	37,642,109
Sep 23, 2019	Share issue	275,125	1,650,751	13,756,255	82,537,530	96,018,660
Oct 10, 2019	Share issue	124,874	1,775,625	6,243,745	88,781,275	43,581,340
Feb 11, 2021	Share issue	8,878,127	10,653,753	443,906,375	532,687,650	532,687,650

Notes

NOTE 1 GENERAL INFORMATION

This interim report includes the parent company InDex Pharmaceuticals Holding AB (publ), Corp. Reg. No. 559067-6820, the subsidiary InDex Pharmaceuticals AB and the sub-subsidiary InDex Diagnostics AB ('InDex', 'the company' or 'the group'). InDex Pharmaceuticals Holding AB (publ) is a parent company registered in Sweden with its registered office in Stockholm with the address Berzelius väg 13, 171 65 Solna, Sweden.

Unless otherwise stated, all amounts are in thousands of Swedish kronor (SEKk). Figures in parentheses refer to the comparative period.

NOTE 2 ACCOUNTING POLICIES

InDex applies International Financial Reporting Standards (IFRS) as adopted by the EU. This report is prepared in accordance with IAS 34 Interim Financial Reporting and the Annual Accounts Act. The parent company prepares financial reports in accordance with the Swedish Financial Reporting Board's recommendation RFR 2 Accounting for Legal Entities and the Swedish Annual Accounts Act.

Applied accounting principles and calculation methods are the same as in the annual report for 2020.

None of the IFRS or IFRIC interpretations that have yet to come into legal effect are expected to have any significant impact on InDex.

RISKS AND UNCERTAINTIES

OPERATIONAL RISK

There is no guarantee that InDex's research and development will result in commercial success. There is no guarantee that InDex will develop products that can be patented, that granted patents can be retained, that future inventions will lead to patents, or that granted patents will provide sufficient protection for InDex's products.

There is no guarantee that InDex will obtain the necessary approvals to conduct the clinical trials that InDex would like to conduct, or that the clinical trials conducted by InDex, independently or in collaboration with partners, will demonstrate sufficient safety and efficacy to obtain necessary regulatory approvals or that the trials will lead to pharmaceuticals that can be sold on the market. It cannot be excluded that the regulatory approval process will require increased documentation and thereby increased costs and delays in projects or lead to projects being shut down. Increased development costs and longer development time

may mean that the risks of a project increase and that the compound's potential to successfully reach the commercial stage decreases or that the time for patent protected sales is reduced.

FINANCIAL RISK MANAGEMENT

InDex may also need to raise additional capital in the future. Both the size and timing of InDex's possible need for capital in the future depend on several factors, including the possibility of entering into collaboration or licensing arrangements and the progress made in research and development projects. There is a risk that the necessary financing of the operations is unavailable at the right time and at a reasonable cost.

For a detailed description of significant risks, refer to InDex's annual report for 2020. The annual report is available on the company's website.

NOTE 4 IMPORTANT ESTIMATES AND JUDGEMENTS

The group makes estimates and assumptions about the future. The resulting accounting estimates will, by definition, rarely correspond to the actual results. The assumptions and other sources of estimation uncertainty where there is a significant risk of material adjustment to the carrying amounts of assets or liabilities within the next financial year are outlined below.

(i) Accrued costs for clinical trials

At each balance sheet date, management estimates the proportion of the coming milestone payments that have been accrued. The accrual for accrued costs is based on external parameters coupled with management's estimate of percentage of completion.

(ii) Tax loss carry-forwards

Deferred tax assets related to loss carry-forwards or other future tax deductions are recognised to the extent it is probable that the deduction can be offset against future taxable profits. Since the group does not report positive results no deferred tax asset related to loss carry-forwards has yet been recognised.

(iii) Estimates and assessments linked to development costs

An important assessment in financial reporting refers to the point in time for capitalizing pharmaceutical development costs. Based on the accounting policies set out under note 2 in the annual report for 2020, no pharmaceutical development costs meet the criteria for capitalisation and have therefore been expensed. Pharmaceutical development costs will be, at the earliest, capitalised after positive results have been achieved in phase III clinical trials or until registration studies have commenced. The reasons being that before that time, it is too uncertain whether the costs will generate future economic benefits and that financing of the asset's completion has not been secured.

NOTE 5 RELATED PARTY TRANSACTIONS

No related party transactions have occured from a group perspective.

InDex Pharmaceuticals Holding AB invoices InDex Pharmaceuticals AB for overall group functions.

NOTE 6 EARNINGS PER SHARE

Earnings per share is calculated by dividing the result for the period by the weighted average number of outstanding ordinary shares during the period.

InDex had potential ordinary shares in the form of warrants. However, these did not give rise to any dilution effect in 2020 or 2021 as a conversion to ordinary shares decreases loss per share.

SEK million	Jan-Jun 2021	Jan-Jun 2020	Full year 2020
JEK IIIIIIIOII	2021	2020	2020
Net result attributable to the equity			
shareholders of the parent company	-52.0	-39.6	-57.4
Total:	-52.0	-39.6	-57.4
Total:	-52.0	-39.6	-57.4
Total: Weighted average number of shares	-52.0	-39.6	-57.4
	-52.0 465,280	-39.6 236,750	-57.4 236,750