Bukwang Pharmaceutical

Business results 2023



Making Tomorrow Better

This presentation contains forward-looking statements about Bukwang Pharmaceutical Co., Ltd. Such statements are related to incidents of the future and not of the past, and include business status and financial performance that the company predicts for the future. The predictions and estimations have inherent uncertainties and risks, so please keep in mind that the company's actual future performance may differ from the estimated performance. The inherent uncertainties and risks include changes in relevant rules and regulations, changes in general business environment and fluctuations in the financial market.

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2 R&D updates

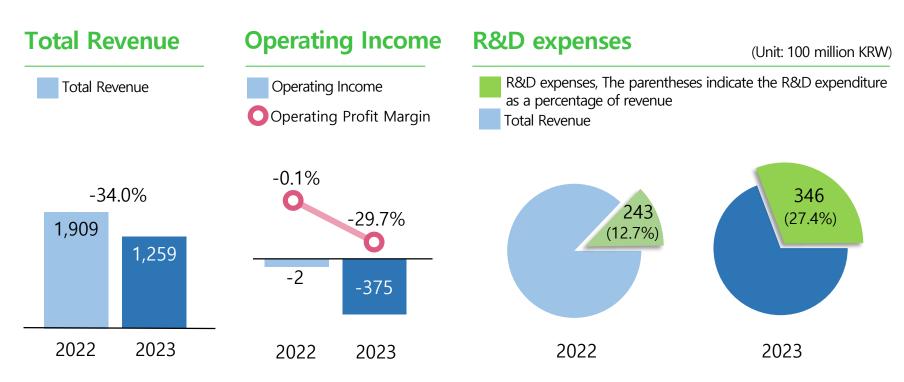
3 Appendix



Key Management Status

Summary of the annual consolidated performance for 2023

Key Management Status

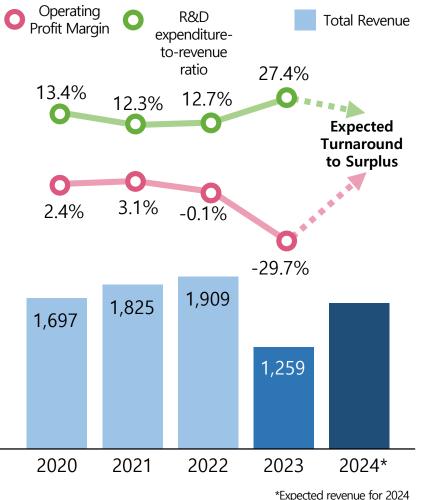


- The decrease in revenue and operating profit is attributed to management improvement measures implemented in the third and fourth quarters of 2023
 - Efforts to enhance financial soundness by reducing credit sales, shortening accounts receivable cycles, and reducing distribution inventory
 - Tight adjustment of distribution margins that had been managed loosely
 - Reconfiguration of product portfolios through careful profitability analysis, including adjustments to underperforming products
- The sales volume of prescription drugs, a key revenue factor, is showing an increasing trend
- R&D expenses have increased due to the final stages of the JM-010 European Phase 2 clinical trial



Key Issues and Business Outlook

Revenue, operating profit margin, <u>R&D expenditure ratio trend</u>



Highlights

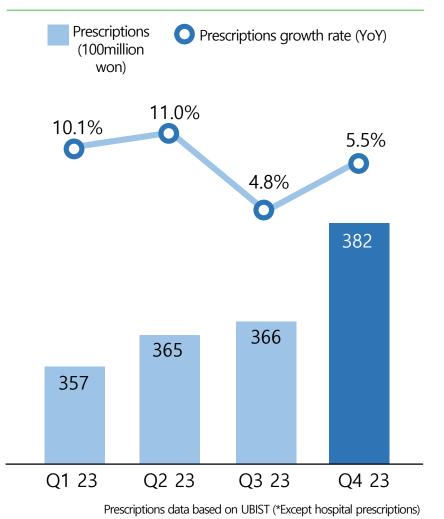
Highlights of 2023

- During the process of restructuring our client base focusing on key trading partners, there have been some returns and discontinuations of shipments
- Supply issues have arisen with certain items such as antiviral agents
- Reverse-base effect on COVID-19 beneficiary items

Business Outlook for 2024

- The release of the new CNS drug Latuda is expected to increase revenue in the CNS sector
- Efforts to reduce return rates through efficient wholesale supply and inventory management have been completed, and this is expected to lead to improved profitability
- Successful stabilization of the sales department's organization is enabling exploration of various growth opportunities

Quarterly Trends in outpatient prescriptions



Highlights

Q4' 23 Highlights

- Prescriptions-driven growth in focused management items (YoY)
 - Legalon (+8.7%), Dexid+Thioctacid (+6.1%),
 Feroba-You (+6,1%), Respiratory products (+14.4%),
 CNS products (+2.1%)

Q1' 24 Highlights

- New prescriptions of Zaledeep
- Sustained growth in key strategic items such as Dexid, Legalon
- Strengthening of Feroba-You in hospital's channel activities
 - reinforcing activities in Gastroenterology and Nephrology
- Preparation for the release of Latuda establishment of a new organization in the medical department

Key Management Status

Business/R&D/Sales Objectives and Strategies

1. Profit-driven sales and marketing

Achieving a surplus in 2024

- Expansion of market share through growth in strategic product categories for ETC
- Increase in CNS sales
 Successful launch of the new CNS drug Latuda
 Activation of existing CNS product sales such as Jaledeep
- Securing competitiveness through efficient distribution strategies and restructuring of sales networks

3. Securing competitive production costs

Cost-competitive production of products and effective inventory management

- Establishment of a competitive production system
- Company-wide production management
- Compliance with production, returns, and inventory management regulations

2. Building a profitable product portfolio and maintaining competitive R&D efforts

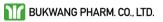
Acquisition of products and new drug development to improve operating profit

- Reconstruction of product portfolios based on contribution margin
- Continual pipeline development for the future including new drugs, improved drugs, and generic development
- Sustaining global open innovation strategies
- Establishment of strategies for early realization of appropriate drug values

4. Driving the enhancement of business processes and stabilizing the organization

Implementation of business systems and structures to ensure transparency and competitiveness

- Establishment of organizational operating systems for standardization and enhancement of business processes
- Implementation of business systems and structures to ensure transparency and competitiveness



R&D updates

Indication & Mode of action

- Antipsychotic drug developed by Sumitomo Pharma in Japan
- Treatment of Schizophrenia/bipolar depression
- Lurasidone acts as an antagonist of dopamine D₂ receptor, serotonin 5-HT₂ receptor, and 5-HT7 receptor

Market Size

- After FDA approval in October 2010, achieved North American sales of USD 1.5 billion in 2022
- Marketed in over 53 countries (US, Canada, EU, UK, Switzerland, Russia, Japan, China, Singapore, Thailand, Hong Kong, Taiwan, Australia, United Arab Emirates (UAE), Kuwait, Saudi Arabia and Brazil)

Contract

- Entered into a License agreement with Sumitomo Pharma (Apr, 2017)
- Bukwang has exclusive development and commercialization right in Korea

Progress Situation

- Approval of phase 3 study in Korea (Aug, 2017)
- Phase 3 study patient enrollment completed (June, 2022)
- Positive phase 3 study top-line results reported. (July, 2022)
- NDA submission to the Ministry of Food and Drug Safety (MFDS) in Korea (Oct, 2022)
- Application for health Insurance coverage (Sep, 2023)
- NDA approval by the MFDS (Nov, 2023)

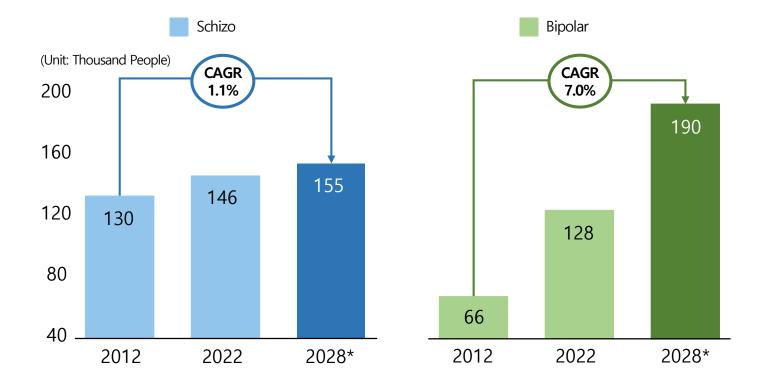
Upcoming Schedule



Latuda(Lurasidone) : The number of patients by indication

R&D updates

- According to HIRA data, As of 2022, there were a total of 146,000 patients with schizophrenia, and it is predicted to grow at a rate of approximately 1% annually in the future.
- For patients with bipolar disorder, the predicted number is approximately 128,000 as of 2022, with an anticipated growth rate in the 7% range
- It is forecasted that from 2025 onwards, the number of patients with bipolar disorder will surpass those with schizophrenia



Source: Self-prediction by Bukwang Pharmaceutical based on data from the Health Insurance Review and Assessment Service (HIRA)

JM-010 (Contera Pharma) : Treatment for Dyskinesia in PD

Indication

Dyskinesia in Parkinson's Disease

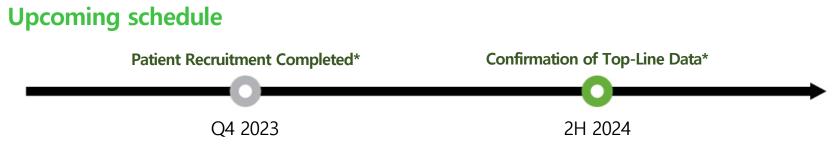
Development Partner

 Contera Pharma A/S, a CNS focused Danish subsidiary, is currently undergoing joint development with Bukwang where Contera has the development and commercial rights in Europe and U.K while Bukwang has the Rest of the World (ROW) right

Progress

- Phase 2a (PoC) study completed in South Africa (Nov 2015)
- Phase 1 study completed in Germany (2016)
- toxicity testing for Phase 2 clinical trials was completed (2017)
- From 2019 to the present, Phase 2 studies are ongoing in the United States and Europe/Korea
- From 2022 to 2023, expansion of clinical sites to include South Korea and Slovakia for European clinical trials
- As of January 2024, patient recruitment for phase 2 EU/KR study* and the first part of the phase 2 study in the United States have been completed

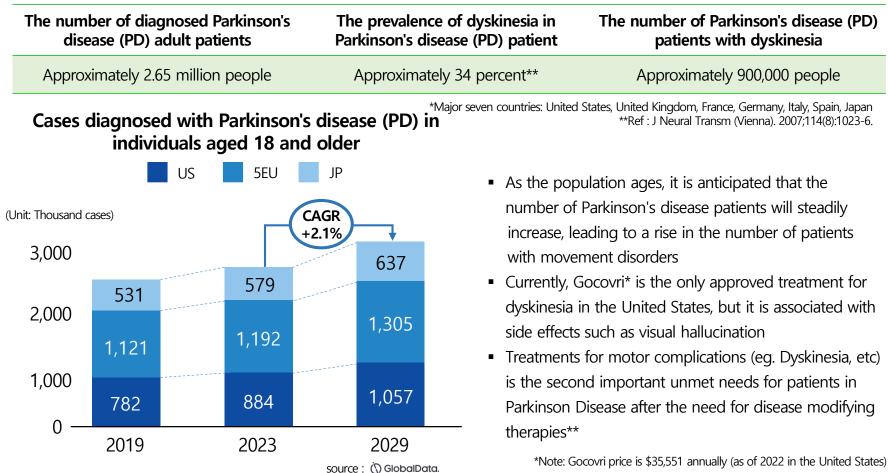
*Countries where clinical trials are conducted in Europe and Korea: Germany, France, Spain, Italy, Slovakia, South Korea



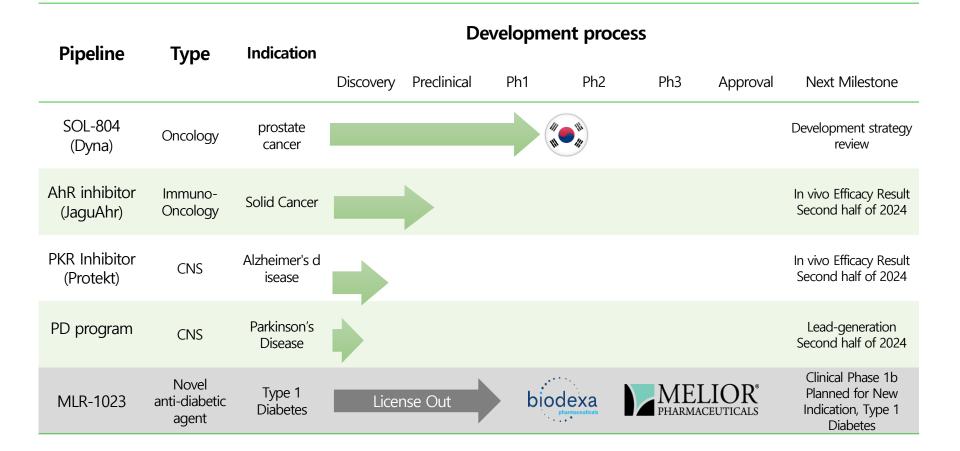
MUKWANG PHARM. CO., LTD.

R&D updates

It is estimated that there are approximately 900,000 patients with Parkinson's disease worldwide across the seven major countries*, and this number is expected to steadily increase









APPENDIX - Profit and Loss of fourth Quarter in 2023

Separate	Unit: 100 million KRW	2023	YoY	2022	2021
	Sales Revenue	1,252	-34.2%	1,903	1,816
	COGS ratio(%)	63.1%		58.7%	59.0%
	R&D expenses	155	29.2%	120	149
	R&D ratio(%)	12.4%		6.3%	8.2%
	Operating Income	-172	swing to loss	123	132
	Margin(%)	-13.7%		6.5%	7.3%
	EBITDA	-135	swing to loss	163	177
	EBITDA (%)	-10.8%		8.6%	9.7%
	Net Income	-191	swing to loss	20	24
	Margin (%)	-15.2%		1.1%	1.3%

Consolidated Unit: 100 million KRW	2023	YoY	2022	2021
Sales Revenue	1,259	-34.0%	1,909	1,825
COGS ratio(%)	62.6%		58.4%	58.5%
R&D expenses	346	42.4%	243	225
R&D ratio(%)	27.4%		12.7%	12.3%
Operating Income	-375	loss continued	-2	56
Margin(%)	-29.7%		-0.1%	3.1%
EBITDA	-334	swing to loss	40	103
EBITDA (%)	-26.5%		2.1%	5.6%
Net Income	-344	loss continued	-42	-28
Margin (%)	-27.3%		-2.2%	-1.5%

- The difference between operating profit and net profit attributable to non-controlling interests is mainly due to variances in Contera Pharma's research and development expenses

APPENDIX - Summary of Consolidated Financial Position

Appendix

Unit (100million won)	2023	2022	changes
Current Assets	2,542	2,187	356
Cash & Cash equivalents	1,512	855	657
Account Receivables	357	880	-523
Inventories	568	386	182
Others	105	66	39
Non-Current Assets	1,748	1,822	-74
Fellow subsidiary & Investments in Associates	131	153	-22
Investments	176	198	-22
Tangible Assets	1,210	1,219	-9
Intangible Assets	164	70	94
Others	67	182	-115
Total Assets	4,290	4,009	281
Total Liabilities	1,948	1,291	657
Accounts Payables & Others	24	165	-141
Contract Liability	185	171	14
Borrowings	800	-	800
Others	939	955	16
Total Equity	2,343	2,718	-375
Leverage ratio	83%	47%	36%

Key changes

- Account Receivables
 Decrease in sales leading to a
 reduction in accounts receivables
- Inventories
 Increase in inventories in sales reduction
- Accounts Payables & Others
 Decrease in production volume due
 to increase in inventory assets

Borrowings

Increase in long-tem borrowings

Thank you!

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