

Investor Presentation

November 2020

NASDAQ: KRMD

FORWARD-LOOKING STATEMENTS / NON-GAAP MEASURES



This presentation contains "forward-looking statements" within the meaning of the safe harbor provisions of the U.S. Private Securities Litigation Reform Act of 1995. Forward-looking statements can be identified by words such as : "expect," "plan," "goals," "believe," "intend," "see," "could," "should," and similar references to future periods. Examples of forward-looking statements include, among others, statements we make regarding our long-term growth potential and sustainability, our strategic growth initiatives and long-term financial goals, issues expected with U.S. plasma supply, expected increase in IG supply, and the potential impact of COVID-19 in the market. Forward-looking statements are neither historical facts nor assurances of future performance. Instead, they are based only on our current beliefs, expectations and assumptions regarding the future of our business, future plans and strategies, projections, anticipated events and trends, the economy and other future conditions. Because forward-looking statements relate to the future, they are subject to inherent uncertainties, risks and changes in circumstances that are difficult to predict and many of which are outside of our control. Our actual results and financial condition may differ materially from those indicated in the forward-looking statements. Important factors that could cause our actual results and financial condition to differ materially from those indicated in the forward-looking statements include, among others, changes to health care policies; success of our research and development efforts; our ability to raise capital if or when needed; acceptance of and demand for new and existing products; expanded market acceptance of the FREEDOM Syringe Infusion System; our ability to obtain required governmental approvals; success in enforcing and obtaining patents; continued performance by principal suppliers; continued customer preference to work through distributors; continued service of key personnel and attracting and maintaining new pe

Non-GAAP Adjusted EBITDA

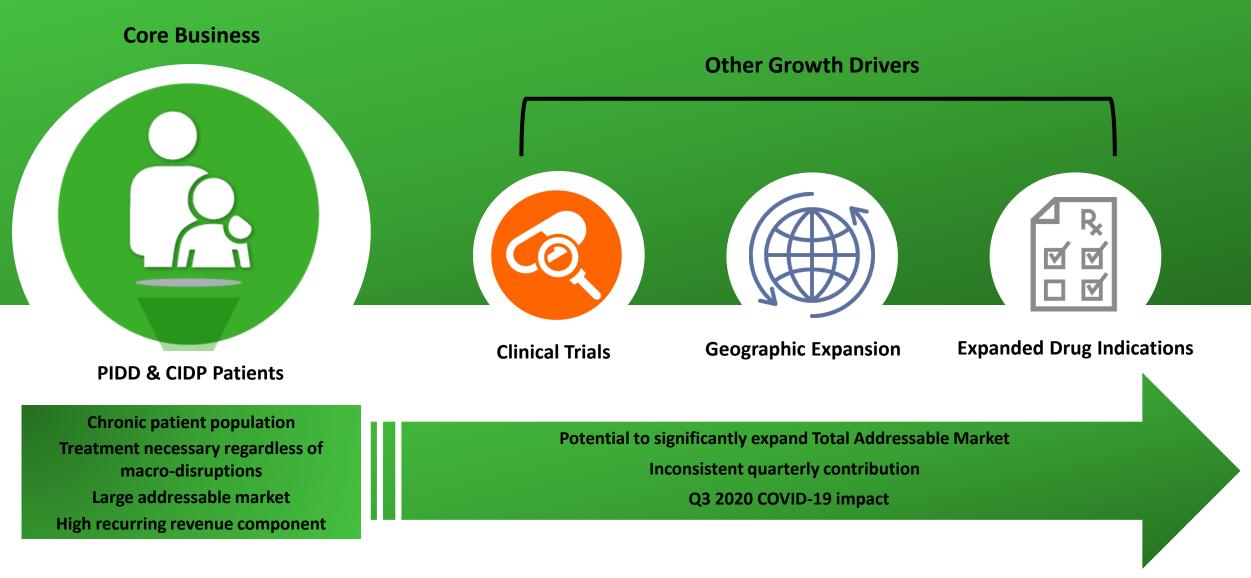
Adjusted EBITDA excludes from net income / (loss): income tax expense, depreciation and amortization, interest income, net, reorganization charges, discontinued product expense, litigation expenses including stock-based settlement expense, manufacturing initiative expenses, and stock option expense

Non-GAAP Measures

This presentation includes the non-GAAP financial measure of "Adjusted EBITDA," that is not in accordance with, nor an alternate to, generally accepted accounting principles and may be different from non-GAAP measures used by other companies. In addition, this non-GAAP measure is not based on any comprehensive set of accounting rules or principles. Non-GAAP financial measures should not be considered a substitute for, or superior to, measures of financial performance prepared in accordance with GAAP. They are limited in value because they exclude charges that have a material effect on our reported results and, therefore, should not be relied upon as the sole financial measures to evaluate our financial results. The non-GAAP financial measure is meant to supplement, and to be viewed in conjunction with, GAAP financial results. A reconciliation of our non-GAAP measure is included in this presentation.

BUSINESS PROFILE



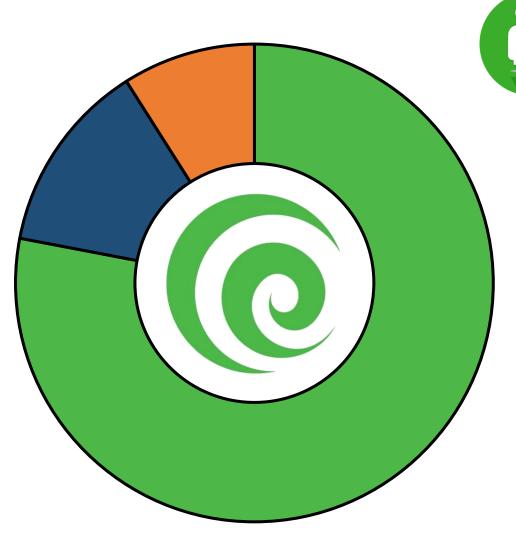


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New Life, New Beginnings -3-

BUSINESS PROFILE





Representative YTD 2020 Revenue Profile

Core Business / U.S. PIDD + CIDP Patients

- Chronic disease state
- Large addressable markets
- High recurring revenue component; minimal patient churn
- Growth aligned with that of Ig market



Clinical Trials

- Potential to significantly expand Total Addressable Market
- Approvals add to Core Business
- Utilize "off-the-shelf" product; customize where necessary
- Creates new and strengthens existing pharmaceutical relationships
- Position us to participate in new clinical trials



International Sales

- A developing revenue stream
- Potential to significantly expand Total Addressable Market

KORU MEDICAL SYSTEMS



PUTTING THE PATIENT FIRST WITH EASY-TO-USE HOME INFUSION SOLUTIONS

KORU Medical Systems manufactures and sells the Freedom Integrated Infusion System that allows chronically-ill patients to self-administer *subcutaneous infusion therapy* in their homes





Supporting the Migration to At-Home Healthcare

Benefitting from

Secular Growth Trends



Pursuing Multiple Growth Pathways

Lowering Costs;

Delivering Improved

Outcomes





Clinical Trial Activity



Strong Core Business





KORU'S POSITIONING



LEVERAGING OUR CORE STRENGTHS TODAY TO DRIVE GROWTH TOMORROW



Significant Market Share and Growth Potential



Differentiated Technology



Strong Pharma Relationships



Premium Customer Retention / Recurring Revenue Component



Value Proposition Across The Care Continuum

Growing Adoption and Opportunity in Current Space (Ig)	Expanded Indications for Existing Therapies (e.g. Secondary Immunodeficiency)
K K N	Rx M M M M M M M
Expand Outside the United States	Support Drug Development for New Disease States

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OUR CORE BUSINESS OPPORTUNITY UNDERPENETRATED MARKETS = COMPELLING GROWTH OPPORTUNITY EXCLUDES OTHER DISEASE STATES AND SECONDARY IMMUNE DEFICIENCY DISEASES



Primary Immunodeficiency Disease

400+ immune diseases

\$750 Annual Recurring Revenue Per PIDD Patient

Chronic Inflammatory Demyelinating Polyneuropathy

Expanded Indication

\$1,500 Annual Recurring Revenue Per Patient (Assumes 2 treatments per week)

US PIDD patient population data from https://www.cslbehring.com/patients/find-your-disease/immunodeficiency-and-autoimmune-diseases

Global PIDD patient population data: <a href="https://www.businesswire.com/news/home/20200915005293/en/European-Medicines-Agency-Approves-Label-Update-for-HYQVIA®-Human-Normal-Immunoglobulin-10-and-Recombinant-Human-Hyaluronidase-Expanding-its-Use-to-a-Broader-Group-of-Patients-with-Secondary-Immunodeficiencies CIDP patient population (2018) from https://www.ajmc.com/journals/supplement/2018/examining-therapies-cidp/chronic-infammatory-demyelinating-polyneuropathy-considerations-for-diagnosis-management-and-population-health. Other figures are KORU Medical estimates.



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\$37.5 M Total U.S. Addressable Market

\$375 M

Total U.S. Addressable Market

> **25,000** U.S. CIDP Population

Market leader High patient retention

500,000

Undiagnosed U.S.

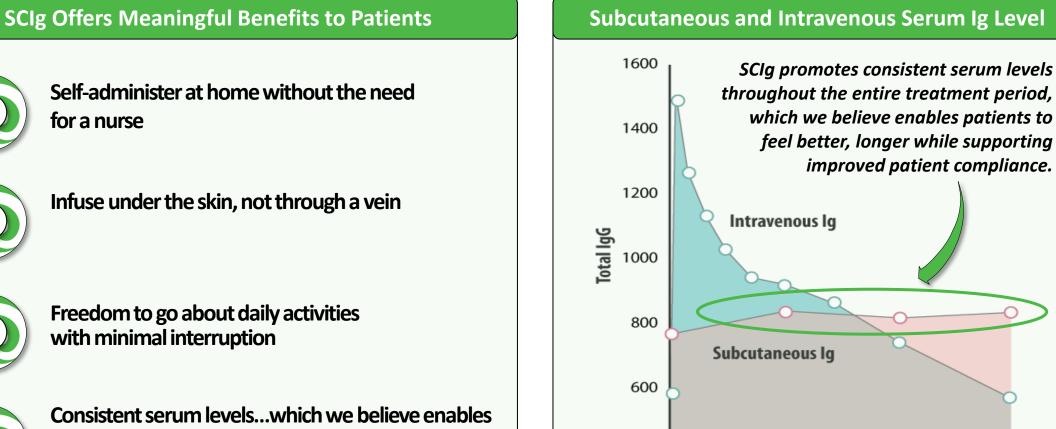
PIDD Population

6,000,000

Estimated Global PIDD Patient Population; High Potential for Additional Diagnosis

5% Total U.S. Market Penetration (KRMD)





400

0

2

6

8

Days

patients to feel better, longer

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10 12 14 16 18 20 22

AT-HOME INFUSION: THE SAVINGS OPPORTUNITY (1)



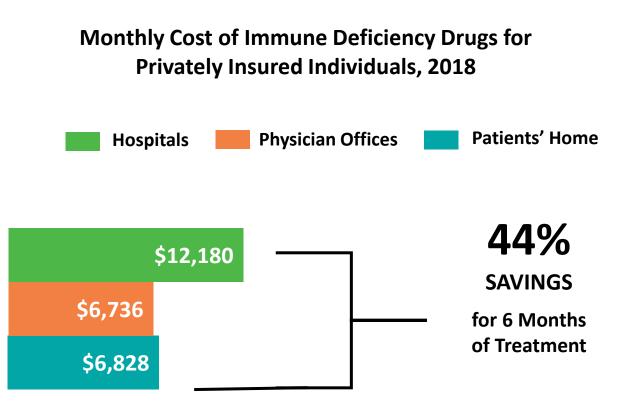


"For certain administered specialty drugs, treatment at home can improve patients' physical and mental well-being and reduce disruption of work schedules and family responsibilities, all without increasing the likelihood of adverse drug events or side effects."

UNITEDHEALTH GROUP® Reducing Specialty Drug Costs / September 2019

(1) UnitedHealth Group®: Reducing Specialty Drug Costs / September 2019.

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CORE BUSINESS: GROWING NUMBER OF SCIg DRUGS



CSL Behring

First 20% SCIg therapy cleared to treat PIDD Cleared to treat CIDP (2018)

Collaborative approach - KORU + Pharma companies



KORU Medical's Freedom60® is featured at <u>www.hizentra.com</u> and in a national advertising campaign for Hizentra® SCIg therapy

Source: <u>www.hizentra.com</u>

GRIFOLS







octapharma



CLINICAL TRIALS DEVELOPMENT PIPELINE*



Therapeutic Focus



Hematology



Nephrology



Neurology



Respiratory



Oncology



Acute Care Instrumentation

Immunology / Inflammation



Primary + Secondary Immunodeficiency

Total Addressable Patient Market Potential +100 M Worldwide"



*October 2020. Some of these agents and applications are investigational and have not been evaluated or approved by the US Food and Drug Administration (FDA) or any other regulatory agency worldwide for the uses under investigation. There is no guarantee any of the agents or applications will demonstrate clinical/commercial success.

**Based on management analysis

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CLINICAL TRIAL INITIATIVES: COMPLETED PHASE III HEMATOLOGY CLINICAL TRIAL USING FREEDOM SYSTEM





Source: https://twitter.com/alexionpharma/status/1082966719524732928?lang=da

Paroxysmal Nocturnal Hemoglobinuria (PNH)

This new PNH drug is advancing towards its planned 2021 U.S. / global launch

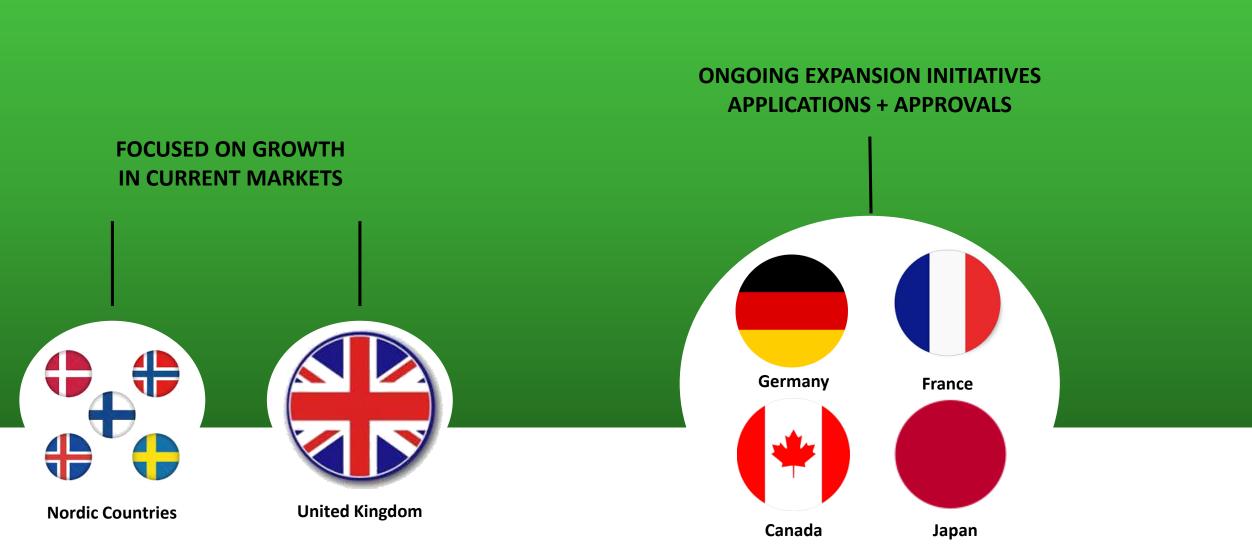
KRMD believes that its Freedom System will be used in several additional upcoming clinical trials focused on expanding indications and disease states for this same drug.

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GEOGRAPHIC EXPANSION



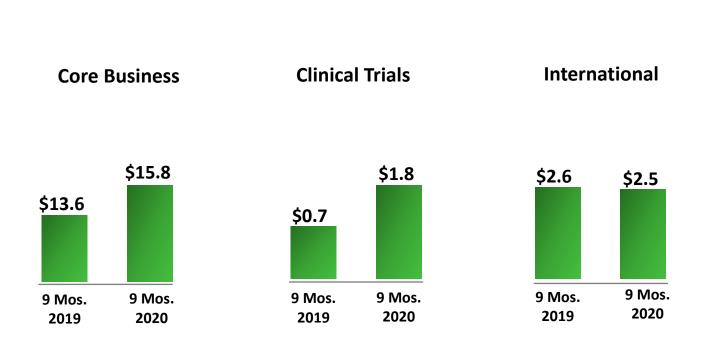


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2020 NINE MONTH NET SALES (\$ in MMs)



Net Sales +19% \$20.1 \$16.9 9 Mos. 9 Mos. 2019 2020



BALANCE SHEET AND CAPITAL STRUCTURE





\$32.4 M Cash & Cash Equivalents September 30, 2020



48.0 M Diluted Shares Outstanding September 30, 2020



\$10 M Share Repurchase Plan Authorized

(\$ in millions)	September 30, 2020	December 31, 2019
Cash & Cash Equivalents	\$ 32.4	\$ 5.9
Current Assets	\$ 42.6	\$ 11.9
Total Assets	\$ 45.4	\$ 13.9
Total Liabilities	\$ 5.5	\$ 2.7
Shareholders' Equity	\$ 39.9	\$ 11.2

INVESTMENT CONCLUSIONS





Leading Product Market Share



Expandable Patient Base in Core Addressable Markets



Exploring Multiple Growth Pathways



Clinical Trial Participation



Supporting Migration to At-Home Care



Well Capitalized



YTD 2020 Adjusted EBITDA \$3.9 M Cash Flow Positive RECONCILIATION OF GAAP NET INCOME TO NON-GAAP ADJUSTED EBITDA



		Nine Months Ended		
		<u>September 30,</u>		
		<u>2020</u>		<u>2019</u>
GAAP Net Income/(Loss)		(377,435)	\$	644,606
Income Tax Expense		316,200		189,265
Depreciation and Amortization		297,801		252,594
Interest Income, Net		(23,690)		(59,091)
Reorganization Charges		_		354,926
Discontinued Product Expense		71,318		
Litigation*		2,446,747		2,481,471
Manufacturing Initiative Expenses		194,804		120,386
Stock Option Expense		<u>1,011,140</u>		<u>640,775</u>
Non-GAAP Adjusted EBITDA**		3,936,885	\$	4,624,932

*For the nine months ended September 30, 2020, litigation consisted of a \$2.2 million non-cash, stock-based settlement expense.

**Adjusted EBITDA excludes from net income / (loss): income tax expense, depreciation and amortization, interest income, net, reorganization charges, discontinued product expense, litigation expenses including stock-based settlement expense, manufacturing initiative expenses, and stock option expense.

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THANK YOU



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