

### **Forward-Looking Statements**

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 regarding our expectations for future performance, including but not limited to our future revenue, gross margin, operating expenses, and cash flow. Forward-looking statements are neither historical facts nor assurances of future performance and are based only on our current beliefs, expectations and assumptions. Forward-looking statements 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. Therefore, you should not rely on any of these 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, the following: impact of COVID-19 related to new SCIg patient starts, plasma supply, clinical trial activity, supply chain and labor availability; inflationary impacts; effects of war and other global conflict; introduction of competitive products; availability of insurance reimbursement; changes in U.S. Food and Drug Administration regulations; 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 and any new product we introduce; 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 personnel; and general economic and business conditions, as well as those risks and uncertainties included under the captions "Risk Factors" in our Annual Report on Form 10-K for the year ended December 31, 2021 and our Quarterly Report on Form 10-Q for the quarter ended June 30, 2022, which are available on the SEC website at <a href="https://www.sec.gov">www.sec.gov</a> [sec.gov] and on our website at <a href

Sales: All references to sale(s) within the presentation refer to net sale(s).





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# KORU Medical Well Positioned to Create Long-Term Value as a Drug Delivery Player

- Manufacture and sell innovative, easy-to-use Freedom mechanical infusion systems
- Leading share position in growing large volume SCIg drug therapy market for patients suffering from Primary Immunodeficiency (PIDD) and Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) – \$300M<sub>(1)</sub> U.S. market potential with existing approved drugs
- Attractive economics: high product loyalty with low churn and recurring revenue model
- Capitalizing on accelerating shift from institutional care to lower-cost home and alternative site settings
- Expanding leadership position into new drug therapies \$1B market opportunity

Market Leading Home infusion platform

25k+

Patients on Freedom Platform

Doubled label Indications Since 2021

12

Commercialized drugs/indications on KORU Medical pumps Growing SCIg market

<20%

SC penetration \$300M<sub>(1)</sub> potential

Expanding to New Drug Therapies

\$1.0B<sub>0</sub>

Total addressable market



### **Q2 2022 Highlights**



Q2 2022 marked third consecutive quarter of double-digit year over year growth



Continued **progress within our novel therapies pipeline** and new deal advancements



Core domestic growth continued to outpace the overall U.S. SCIg market



Q2 gross margins affected by **first half 2022 supply chain disruptions and labor shortages** 



**Continued to solidify our foundation** with new corporate headquarters, outsourced manufacturing, and strengthened executive team



Raised revenue guidance to \$27.0 to \$27.5M

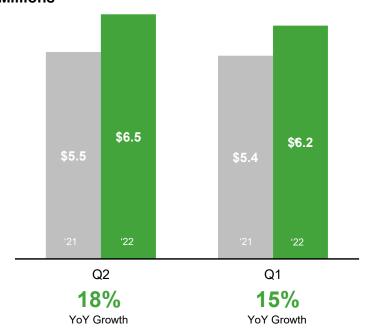


### **Strong Q2 Sales Momentum**

\$6.5M\*

Q2 2022 net sales

YoY Quarterly Net Sales; in Millions



- 18% year-on-year sales growth
  - Three consecutive quarters of double-digit growth
- Strength across all three businesses
  - Novel therapies 747% growth driven by pipeline expansion
  - Domestic core market growth of 8.7% outpacing a rebounding U.S. SCIg market
  - International business growth of 10.7% driven by key tender wins multiple European markets



<sup>\*</sup> Does not include \$0.3M backorders in Q2 arising from supply chain issues

### **Executing on our Strategic Milestones**

## Increase Core SCIg Penetration \$300M<sub>(i)</sub> U.S. TAM

## New SCIg Patient Starts

- U.S. SClg market growth 6.9%<sub>(2)</sub>QTD
- KORU Medical domestic core growth 8.7%<sub>(3)</sub> Q2 2022
- YTD Freedom pumps +21% growth

## Win SCIg Prefills

- Prefill market growth of 437%<sub>(2)</sub> QTD, 10%<sub>(2)</sub> penetration
- FreedomEdge® only 510(k) clearance with prefills

## **Geographic Expansion**

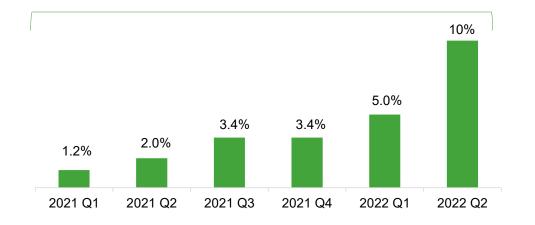
- 10.7% international growth
- Strength in several EU markets – driven by label expansion and tender wins
- Hired new international sales leader



# Prefilled Syringes Growth Creates Short- and Long-Term Upside Potential

Prefilled syringes are rapidly gaining market penetration

## Prefilled Penetration % of SCIg grams<sub>(1)</sub>



### KORU Medical upside share and market expansion opportunity

- As the only FDA cleared platform FreedomEdge® for use with prefilled syringes, a prefill patient is a KORU Medical patient
- Easier infusions via prefills drive market expansion. KORU Medical benefits as share leader
- Focused KORU Medical innovation effort on prefills



### **Continued Momentum in Novel Therapies**

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**Total Closed** 

## **Extend to Novel Therapies** \$1B<sub>(1)</sub> U.S. TAM

#### Focus on large volume >10ml

- Q2 Novel Therapies Sales Growth of 747% reflects momentum from 2021 deals
  - Sales for Novel Therapies includes finished product sale for use in clinical trial, and/or developmental services sales in the form of non-recurring engineering (NRE) fees
- Two new closed deals in Q2:
  - 1 new phase III Ig deal
  - 1 new phase II new drug deal in nephrology
  - Expanded scope of previously announced innovation agreement
- Nine total closed agreements across 6 drug categories
  - 10-15 opportunities in pipeline

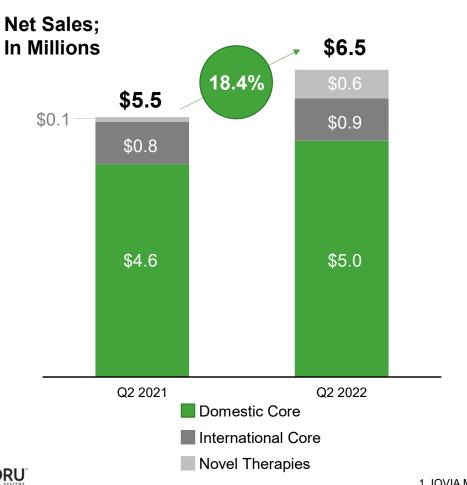
Q2 2022 New Cl	<b>2</b> osed in Q2 2022		aa aa	
	Development Stage	lg	New Drugs	Total
Closed	Phase III	1	0	1
	Phase II	0	1	1
Total Closed		1	1	2

Closed Agreement N	s 9 ew Closed Since 2021	•	Respiratory • Oncology • Immunology •	Hematology Neurology Nephrology	
	Development Stage	lg	New Drugs	Total	
Closed	Phase III	3	1	4	
	Earlier	1	4	5	





#### **Q2 2022 Sales**



#### **Novel Therapies**

- Increased 746.6% year on year
- Driven by NRE development milestone and pipeline expansion

#### **Domestic Core**

- Increased 8.7% year on year
- Rebounding SCIg market growing 6.9%<sub>(1)</sub> through Q2 2022
- Reported growth excludes \$0.3M backorders arising from supply chain issues

#### **International Core**

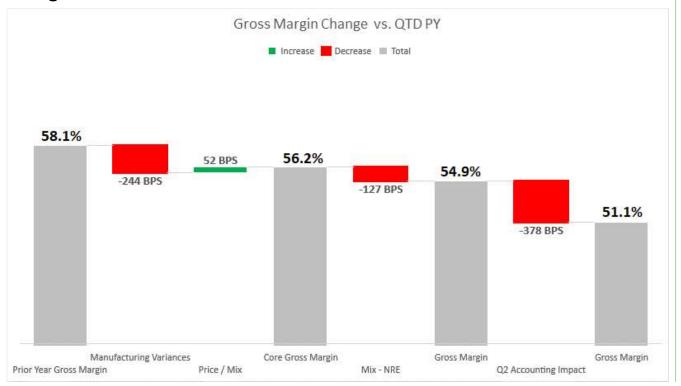
- Increased 10.7% year on year
- Due to growth in several European countries driven by key tender wins

( KORU™

1. IQVIA Market Data

### Q2 2022 Gross Margin

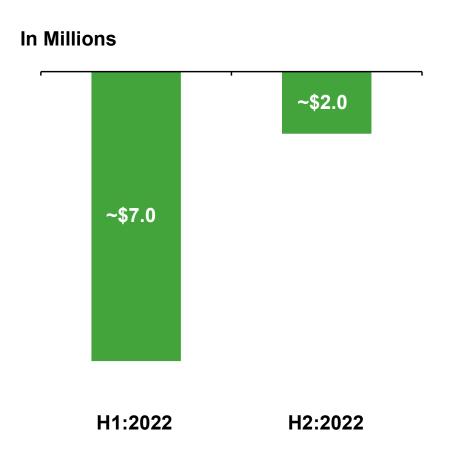
#### Change vs. Q2 2021



- 56.2% Core Product Gross
   Margin driven by supply
   shortages from Q1 and
   favorable average selling prices
- 54.9% Gross Margin inclusive of higher service revenue mix, having a dilutive impact
- 51.1% Reported Gross Margin post accounting impact of
  manufacturing variances
  accelerated and amortized in Q2
  for supply issues and labor
  shortages contributing to lower
  finished goods inventory



### 2022 Forecasted Usage of Cash



#### Ending cash balance of ~\$16.0M

#### First Half 2022

- (-) Net Loss
   Lower gross margins due to higher production variances and NRE revenue
- (-) HQ relocation to Mahwah, NJ ~\$2.0M
- (-) Higher raw material and work in process (WIP) inventory

#### Second Half 2022

- (+) Net Loss lower net loss expected than 1H Higher gross margins due to improved operating efficiencies
- (+) Decreased Working Capital
  Lower raw material & WIP inventory levels
- (+) Employee Retention Credit (ERC), equipment financing & leasehold improvement reimbursements ~\$1.5M





## **Guidance and Closing Comments**

Linda Tharby

President and Chief Executive Officer

#### 2022 Guidance

#### Sales Growth

Raising range to \$27.0 to \$27.5 million net sales in 2022

#### **Key Drivers**

- Expanded Novel Therapies pipeline
- Increased core SCIg market growth

## **Gross Margin Profile**

Updating to **55-60%** exit rate for 2022

#### **Key Drivers**

- Q2 2022 supply chain disruptions resulted in unfavorable manufacturing variances in 1H22
- Outsourcing initiative delayed to Q1 2023
- Expect steady margin improvement for remainder of 2022

## Operating Expenses & Cash Flow

Reiterating OpEx range of \$27 to \$28 million in 2022: ending the year with cash balance of ~\$16.0 million

#### **Key Drivers**

- Expect OpEx to stabilize at \$7M per quarter
- Lower Working Capital in 2H
- ERC, Financing & Tenant Improvement Credits



### **Q2 2022 Highlights**



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Core domestic growth continued to outpace the overall U.S. SCIg market



Q2 gross margins affected by **first half 2022 supply chain disruptions and labor shortages** 

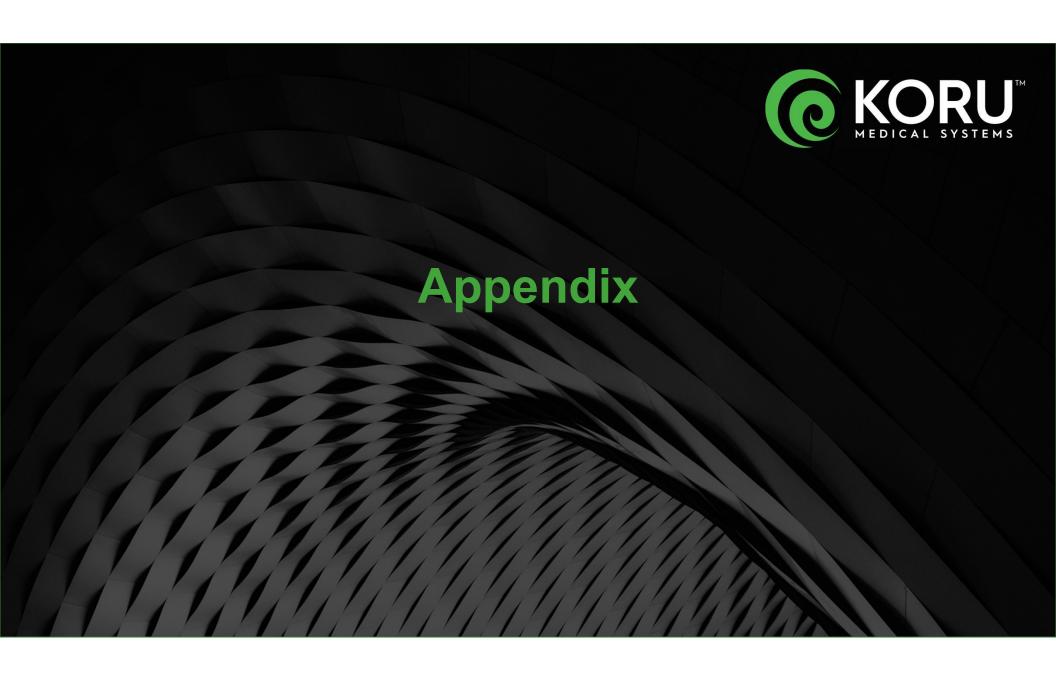


**Continued to solidify our foundation** with new corporate headquarters, outsourced manufacturing, and strengthened executive team



Raised revenue guidance to \$27.0 to \$27.5M





### **GAAP Reconciliation**

Reconciliation of GAAP Net (Loss) to Non-GAAP Adjusted EBITDA:		Three Months June	69504350360	Six Months Ended June 30,			
		2022	2021	esti-	2022	2021	
GAAP Net Loss	\$	(2,921,341)	(1,124,549)	\$	(5,458,855) \$	(2,400,687)	
Tax (Benefit)/Expense		(710,260)	(245,316)		(1,307,859)	(1,187,677)	
Depreciation and Amortization		125,882	118,415		235,134	233,888	
Interest (Income)/Expense, Net		(3,566)	(9,950)		(2,103)	(19,721)	
Reorganization Charges		270,433	224,605		565,433	1,193,880	
Discontinued Product Expense		<del>-</del>	_		— ·		
Litigation Expenses		_	<del>-</del>		<del>-</del>	-	
Manufacturing Initiative Expenses		50,344	149,718		88,349	201,441	
Stock-based Compensation Expense		821,513	605,172		1,659,070	1,339,356	
Non-GAAP Adjusted EBITDA	\$	(2,366,995)	\$ (281,905)	\$	(4,220,831) \$	(639,520)	

Reconciliation of Reported Diluted EPS to Non-GAAP Adjusted Diluted EPS:		Three Months Ended June 30,				Six Months Ended June 30,			
		2022		2021		2022		2021	
Reported Diluted Earnings Per Share	\$	(0.07)	\$	(0.03)	\$	(0.12)	\$	(0.05)	
Reorganization Charges		0.01		0.01		0.01		0.03	
Discontinued Product Expense				_				_	
Litigation Expenses		9.2		1000		85 <u>—2</u> 7		_	
Manufacturing Initiative Expenses						_			
Stock-based Compensation Expense		(502)		<u></u>		2-		0.01	
Tax (Expense) Adjustment		_		-		_		(0.01)	
Non-GAAP Adjusted Diluted Earnings Per			50.0		\$		\$		
Share	\$	(0.06)	\$	(0.02)	52	(0.11)	- 55	(0.02)	



#### **GAAP Reconciliation**

Reorganization Charges. We have excluded the effect of reorganization charges in calculating our non-GAAP measures. In 2021 we incurred significant expenses in connection with the departure and replacement of our chief executive officer and the recruiting of two new board members, which we would not have otherwise incurred in periods presented as part of our continuing operations. In 2022 we incurred further severance expense related to the reorganization of the leadership team and the departure of our chief financial officer, which we would not have otherwise incurred in periods presented as part of continuing operations.

Manufacturing Initiative Expenses. We have excluded the effect of expenses related to creating manufacturing efficiencies, in calculating our non-GAAP measures. We incurred expenses in connection with these initiatives which we would not have otherwise incurred in periods presented as part of our continuing operations. We expect to incur related expenses for the next six to nine months.

Stock-based Compensation Expense. We have excluded the effect of stock-based compensation expense in calculating our non-GAAP measures. We record non-cash compensation expense related to grants of options for executives, employees and consultants, and grants of restricted shares to our board of directors. Depending upon the size, timing and the terms of the grants, the non-cash compensation expense may vary significantly but will recur in future periods.

