Better health. Within reach. Every day.

Hikma US Meet the Management, Columbus Ohio 15 May 2025

DEGI TAN



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Hikma US Meet the Management Hikma Rx

2023

hikma.

Presenting team



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Introducing Hikma Rx

We are renaming our Generics business division As of today, Generics will become:

Hikma Rx

Rationale:

To better reflect our focus on providing differentiated and complex prescription (Rx) medicines.

Hikma's three business divisions:







Columbus, Ohio



Why Columbus?

- Within a one-day drive over 150 million people can be reached from Columbus 46% of the US population and 48% of headquarters operations One Columbus
- First among the 10 largest Midwest Metro for population, job and GDP growth One Columbus
- Ohio ranked 10th for America's States for Businesses CNBC
- Foreign Trade Zone 138 has ranked in the **Top 10** in the nation for warehousing and distribution for the last **8 years** *FTZ Board Annual Report, 2019*
- Home to one of the world's only cargo-dedicated airports One Columbus
- Ohio ranked #2 in the nation for corporate investment, with the Columbus region in the Top 10 for cities BizJournals
- Ninth Best State for Business Chief Executive Magazine

1 Introduction

2 Manufacturing

3 • R&D and Business Development



5 • Wrap-up

1 Introduction

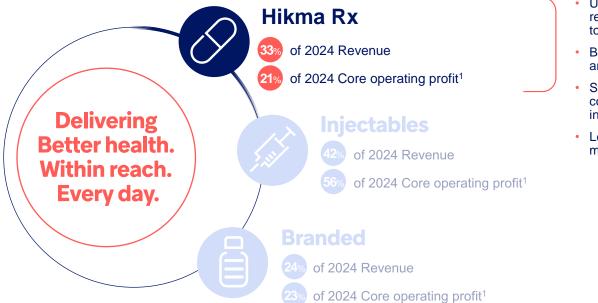
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Our high-quality Hikma Rx business



- US-based manufacturer and supplier of oral, nasal, respiratory and other generic and specialty products to the North American retail market
- Broad and increasingly complex generic and specialty product portfolio
- State-of-the-art US manufacturing facility with complex capabilities including nasal sprays, inhalation and high containment technology
- Leveraging capabilities to increase contract manufacturing partnerships

¹ Core operating profit is \$719 million. Before unallocated corporate costs of \$99 million and operating loss from Other business of \$9 million, core operating profit contribution from business segments is \$827 million Hikma Pharmaceuticals PLC

A differentiated platform to drive long-term success

Building on Hikma Rx competitive strengths





A talented and highly capable team that knows how to compete and win

Strong and expanding R&D team with multiple new programs launched



Large and expanding US-based manufacturing and development capabilities



Excellent record of successful FDA quality inspections



Deep industry experience and strong customer relationships with industry-leading service levels

Delivering on promised strategic and tactical initiatives



2025 FDA inspections; FDA training partner

 Quality: continuously improve to maintain industry-best standards/performance

Introduction

2 Manufactruing

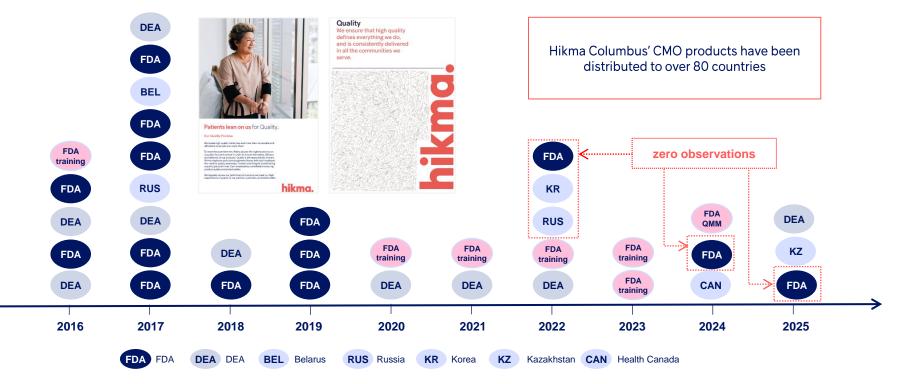
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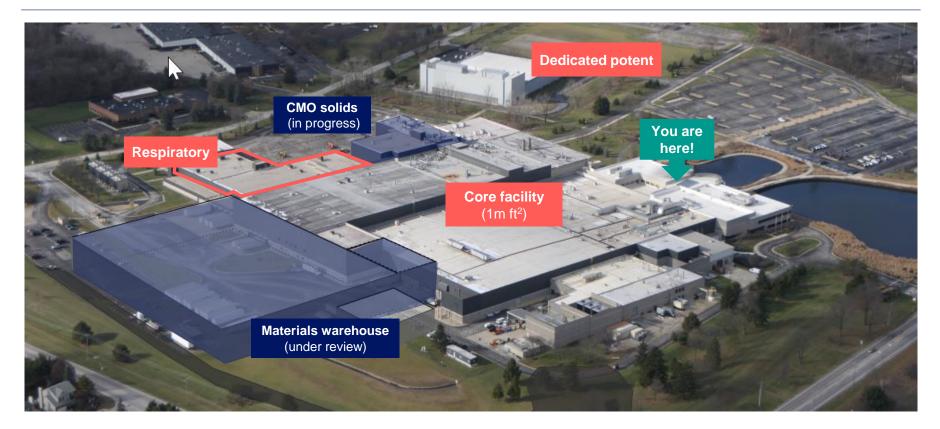
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Uncompromising quality culture with proven track record

Over a decade of excellent audit and inspection results



Our facility and expansion plans (recent and future)

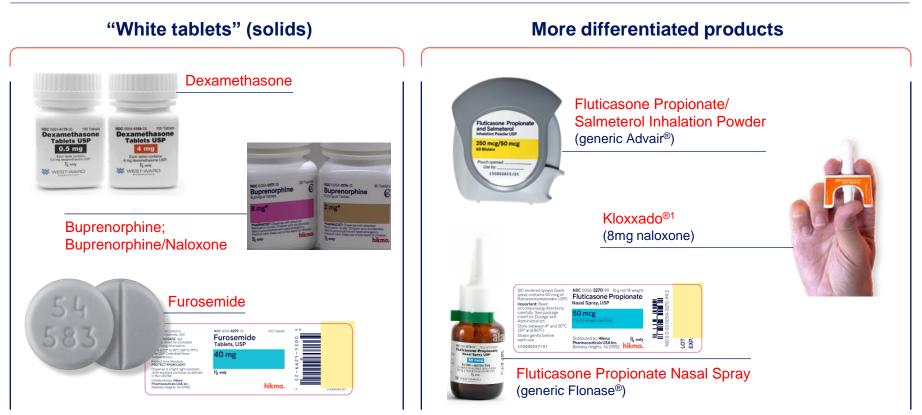


Significant capacity and differentiated capabilities

	Solid Dose ("White Tablet")	Nasals / Liquids	High Containment (HCO)	Dry Powder Inhaler (DPI)
Manufacturing area makeup	 Charging suites Zanchetta granulators Fluid bed granulators/dryers Roller compactors Bin to bin transfer rooms Bin tumble rooms Tote bins (50-2000L) 	 1000-4000L stationary processing tanks Isolated processing suites 50-1000L portable processing tank Clean in Place (CIP) 	 Fully-enclosed equipment Ribbon blenders Roller compactors Single pot granulators Tablet presses Encapsulators Bin tumble rooms Tote bins (150-1100L) 	 Fluticasone/salmeterol Fully-enclosed equipment Conical mill GEA blender Tote bins (30L & 50L)
Key technologies	 Direct blend Aqueous/solvent granulation Roller compaction 	SolutionsSuspensions	 Direct blend Aqueous/solvent granulation Roller compaction 	Direct blendPowder filling in isolationDevice assembly
Packaging area makeup	 Solid tablet dosages Solid capsule dosages Solid powder dosages Foil/foil unit dosages Foil/PVC/foil unit dosages 	 Multi-dose nasal packaging lines Unit-dose nasals Isolation filling line with secondary packaging 	Solid tablet dosagesSolid capsule dosagesFoil/foil unit dosages	 Thermoforming technology Blister strip handling system Mikron coiling technology

Specialised / increasing layers of differentiation

Select products manufactured at our Columbus facility

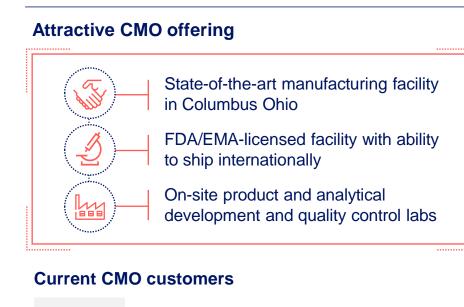


¹ Hikma produces as part of exclusive commercial partnership with Emergent Biosolutions Hikma Pharmaceuticals PLC

Our distribution centers are strategically located to serve key customers

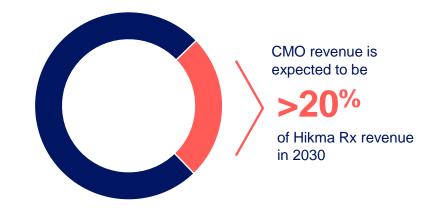
	Creekside	Wilson Rd.
Material types	Finished goods Room temp Cold Chain 	& ram materials erature
Size	275,000 ft ²	Vault 18,000 ft ² ; Starting material 50,000 ft ²
Temperature controls	Mapped with full generator backup	Mapped with electric backup
Regulatory & industrial certified	FDA; OSHA; CTPAT; Hazmat	FDA; DEA; OSHA; CTPAT; Hazmat
Security	U U	; interior and exterior deo
Other highlights	Purpose-built in 2016; centrally located in US	

Growing CMO business





Expect CMO contribution to grow over the next 5 years



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Leveraging our global footprint and highly skilled teams to advance our R&D programmes

Columbus, Ohio, US

- R&D + RA + BD/Pipeline
- Focus on complex products
- Nasal, Orally inhaled, MR/ER tablets
- · State of the art facility
- Leader with >25Y experience (Teva, GSK, Roche, Sanofi)



Berkely Heights, New Jersey, US

- RA, BD/Pipeline
- · Segment leader, commercial





Zagreb, Croatia

- Newly established R&D
- Initial focus on solids
- Leader with >25Y experience (Xellia, Teva, Pliva)



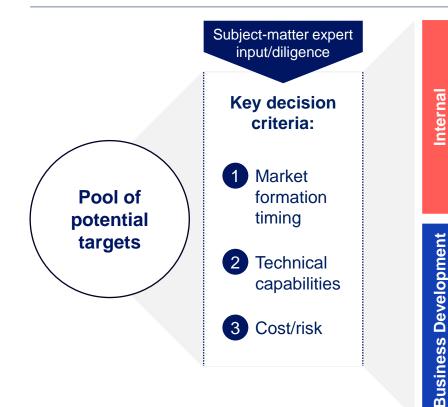
Amman, Jordan

- R&D in Bayader and Sahab
- High potency capabilities
- Hikma Chemicals
 - Low volume, high value API
 - Towards vertical integration
- IPRC
 - Fully equipped CRO
 - >4.5k bioequivalence, bioavailability and PK studies
 - US FDA, EMA plus other regulatory inspected
 - 51% Hikma owned



Hikma Pharmaceuticals PLC

Product selection: utilising internal capabilities and business development to build a robust pipeline



Utilize network of three development sites

- Columbus: complex dosage forms (e.g., DPIs, nasals), controlled substances
- Amman: global programs (US + MENA)
- Zagreb: solid orals (tech transfer to commercial mfg. sites)

Leverage **Hikma Chemicals** for targeted backward integration Utilise **IPRC** for quicker, more flexible BE studies

Deal rationale:

- 1. Market formation: partner's application allows for **earlier launch**
- 2. Technical capabilities: access to **unique technology** from partner
- 3. Cost/risk: share investment burden and upside with partner to **de-risk portfolio**



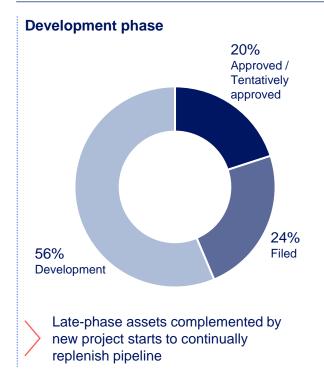


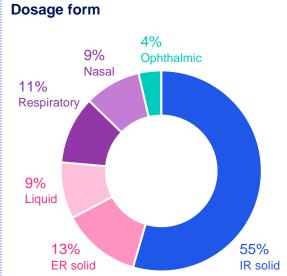
Respiratory portfolio



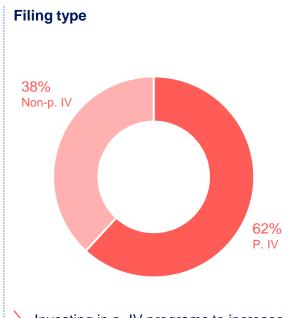
Our pipeline

Key pipeline metrics





Balance of solid orals (comprise majority of future LOEs) and complex respiratory and nasal drug-device combinations



Investing in p. IV programs to increase first-to-market launches

Expanding our pipeline of higher-value first wave launches

First-to-file paragraph IV applications

Reference listed drug (molecule)		Market size (\$m) ¹	Number of filers ²	Status
Vyndamax [®] (tafamidis)	155	\$2,788	3	Active litigation
Spravato® (esketamine)	4 0	\$773	3	Active litigation
Wakix® (pitolisant)	455	\$71 4 ³	7	Active litigation
Mekinist® (trametinib)	455	\$449	1	Settlement (confidential)
Nucynta [®] IR (tapentadol)	\bigcirc	\$171	4-5	AG agreement (30 days prior to LOE)
Sunosi [®] (solriamfetol)	\bigcirc	\$165	6	Settlement (2040)
Steglatro [®] (ertugliflozin)	155	\$147	3	Settlement (confidential)
Fetzima® (levomilnacipran)	\oslash	\$72	8	Settlement (confidential)
Xeljanz® (tofacitinib)	455	\$19	3	Settlement (confidential)

¹ IQVIA US sales, Feb 2025 MAT

² Known ANDA filers (including Hikma) as of Apr 21, 2025

³ Harmony Biosciences 2024 10-K

⁴ Incyte 2024 10-K

⁵ Corcept Therapeutics 2024 10-K Hikma Pharmaceuticals PLC

Follow-on paragraph IV applications

Reference listed drug (molecule)		Market size (\$m) ¹	Number of filers ²	Status
Jakafi [®] (ruxolitinib)	\bigcirc	\$2,792 ⁴	2	Active litigation
Xtandi [®] tablets (enzalutamide)	\bigcirc	\$1,835	8	Settlement (confidential)
Xtandi [®] capsules (enzalutamide)	455	\$904	7	Settlement (confidential)
Korlym [®] (mifepristone)	\bigcirc	\$675 ⁵	4	Settlement (2034 or earlier)
Dificid [®] (fidaxomicin)	\bigcirc	\$399	3	Settlement (confidential)
Nayzilam [®] (midazolam)	₽	\$186	2	Active litigation
Fanapt [®] (iloperidone)	\bigcirc	\$156	6	2027
Nucynta [®] ER (tapentadol)	\bigcirc	\$121	3	AG agreement (30 days prior to LOE)



Investing in highly-differentiated generic respiratory platforms

Generic Respimat[®] SMI¹

- Strategic rationale: few firms capable of successfully developing device; differentiated delivery mechanism
- Market opportunity⁴:



Status: Advanced development; 2H 2026 submission



2 Generic Ellipta[®] DPI²

- Strategic rationale: ability to leverage gAdvair development, regulatory, and manufacturing expertise
- Market opportunity⁴:
 ~\$11b

Status: Device identified; feasibility studies ongoing



- Strategic rationale: propellant transition represents unique opportunity to capture first-mover advantage
- Market opportunity⁴:
 ~\$9b
- Status: Finalising agreement with strategic partner

¹ Soft mist inhaler

² Dry powder inhaler

³ Low global warming potential pressurized metered dose inhaler

⁴ US IQVIA sales Feb 2025 MAT

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Manage risk with success-based milestones

Increased emphasis on complex generics to add sustainable value

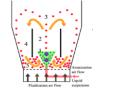
New technologies

- Liquid-filled capsules
- Soft-gel capsules
- Hot melt extrusion
- Multi-particulate delivery systems
- Recruited key talent to implement and deliver increased complexity pipeline

Hot melt extruder

Fluid bed granulator

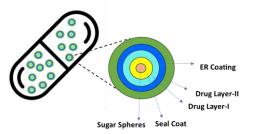




Modified and extended release

- Higher barrier-to-entry products
- Expanding internal technology capabilities to increase differentiation
- Product examples:
 - Generic Ferriprox[®] (Deferiprone b.i.d.)
 - Generic Tarpeyo[®] (Budesonide DR)
 - Generic Entereg[®] (Alvinopan)

Delayed release beads in capsule



Nasal sprays

- Building on solid commercial foundations
- Installing new unit-dose line
- Product examples:
 - Generic Nayzilam[®] (Midazolam, filed)
 - Generic Spravato[®] (Esketamine, filed)
 - Epinephrine (H2 2025 submission)
 - Other novel 505(b)(2) nasal products in development

Kloxxado[®] unit-dose nasal spray



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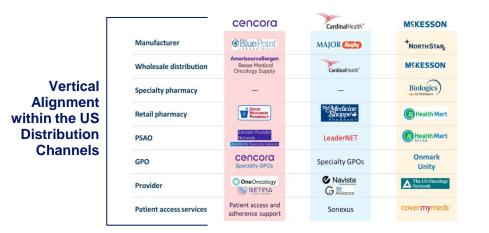
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Competing in a changing regulatory and market landscape



Evolving regulatory and market dynamics

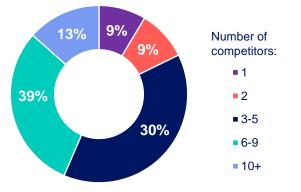
- DSCSA implementation: nearing full implementation of DSCSA with exemptions for smaller customers until November 2026
- Enhanced market position: leveraging 340B program and Inflation Reduction Act (IRA) provisions to strengthen market position
- Price negotiations: IRA's Medicare price negotiations can lead to predictable pricing structures
- Out-of-Pocket Caps: Caps on out-of-pocket spending for Medicare Part D beneficiaries can increase demand for affordable generic alternatives

Hikma in a strong position

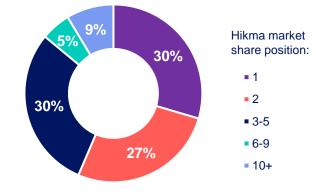
- Strong reputation as a high-quality, consistent supplier
- US based manufacturing within close proximity to customers
- Increasing diversity of revenue stream through geographic expansion

Customers continue to select Hikma as their supplier of choice

Percentage of portfolio by number of competitors



Percentage of portfolio by Hikma market share position



Hikma established and maintains a competitive edge in the marketplace

- Hikma is a top supplier of generics (non-injectables) in the U.S. with a growing portfolio of over 100 product families including differentiated technologies such as inhalation, nasal sprays, and high containment capabilities.
- Successfully sets the standard of excellence by establishing and maintaining best in class supply practices across a multitude of product families

Hikma is a top two supplier for 57% of its portfolio

- Stellar reputation serves as a foundation for developing relationships with key customers that lead to stable and reliable demand
- Portfolio optimization and Lifecycle Management ensure that Hikma remains and flourishes in sustainable markets while differentiating from the competition

Select products contributing to Hikma's success

Sodium Oxybate	Fluticasone Salmeterol (generic Advair®)	Fluticasone (OTC + Rx)	Albuterol (Proventil [®])
 Current authorized generic for Jazz, Xyrem. Approved for the treatment of cataplexy or excessive daytime sleepiness in patients 7 years of age and older with narcolepsy. REMS program ensures Sodium Oxybate is dispensed only through a Certified Pharmacy Utilized Hikma expertise in market 	 Proactively capitalized on accelerated brand-to-generic conversion Significant increases in volume offset natural erosion elsewhere Between 2023 to 2024, Hikma experienced product growth of 135% and over the past 12 months, market share grew from 21% to 34% 	 Continuing to see increased OTC demand from current CMO partner Pricing and supply optimization strategy have continued to establish Fluticasone as a cornerstone product Customers continue to value continuity of supply over price 	 Growth through conversion as the 6.7G (Proventil[®]) continues to increase at the expense of the 8.5G (ProAir[®]) and 18G (Ventolin[®]) Hikma secured significant incremental formularies Proactively capitalized on market supply issues
access to successfully launch and increase unit conversion of Xyrem.	Market Share Trend (per IQVIA)	Hikma YoY % Growth	Market Distribution by Product
 Increasing prevalence of sleep disorders. Market expected to grow at a CAGR of 10% 2023 to 2030. Royalties to Jazz increased 	● GSK ● Hikma ● Prasco ● Teva ● Viatris 39% 37% 36% 38% 41% 40% 39% 33% 37% 34% 35% 32% 31% 30% 24% 24% 25% 25% 23% 19% 18% 17% 17% 18% 18% 17% 17% 17% 17% 14% 11% 12% 12% 11% 11% 11% 11% 11% 11% 10% 10% 5% 5% 5% 4% 5% 5% 0% 0% 0% 0% 0% 0% 0% 0%	+12%	52% 50% 46% 44% 34% 29% 33% 20% 21% 21% 25% 2021 2022 2023 2024
	Jan Feb Mar Apr May Jun Jul Aug Sep Oct Nov Dec Jan Feb Mar 2024	2022 2023 2024	 6.7G 8.5G 18G

Leveraging our US capabilities to drive synergies and success

Integrated market approach: Adapting to evolving market dynamics by merging traditional channel lines, creating a unified and powerful strategy.

Unified strategy: Utilizing cross-functional collaboration to pinpoint & leverage the most effective channels to influence buying decisions and adoption by broad array of market.

Continued market leadership and customer engagement **Growth and relationship building:** Driving growth through diverse and specialty launches, while strengthening relationships with unique customers and limited distribution networks.

Creative positioning: Enhancing market penetration and product reach through a multi-channel approach, firmly establishing Hikma as a dominant leader in the pharmaceutical industry.

Coordinating Efforts to Maximize and Drive Market Leadership



- Hikma Rx impact: Established published pricing, optimized retail price calculations, and enhanced customer engagement for Injectables.
- Strategic execution: Harnessed Hikma Rx expertise and retail market relationships to drive a highly successful launch.
- Market penetration: Captured a remarkable 16% market share within the first quarter following the December 2024 launch.



- Impactful collaboration: Hikma Rx leveraged US Injectables' expertise in the Institutional market to execute a highly successful launch.
- Enhanced outreach: Injectables facilitated outreach to institutional customers, securing product inclusion in Benefits contracts to expand reach across GPOs.
- Market penetration: Achieved a notable 15% market share within the first quarter following the December 2024 launch.

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Hikma Rx – well positioned for the future

Our strategy positions Hikma Rx for continued near-term success, with an acceleration in profitability over the longer-term



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Hikma US Meet the Management Injectables

Presenting team



Dr Bill Larkins

President, Injectables



Joel Rosenstack

Chief Commercial Officer, US Injectables



Nassim Rahmani

Associate VP, Commercial Europe



Natheer Masarweh SVP, Operations



Ragheb Abu Rmaileh



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Three high quality businesses

Diversity through three distinct business units



1. Core operating profit is \$719 million. Before unallocated corporate costs of \$99 million and operating loss from Other business of \$9 million, core operating profit contribution from business segments is \$827 million

Hikma Pharmaceuticals PLC

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2. IQVIA MAT December 2024, generic injectable volumes by eaches, excluding branded generics and Becton Dickinson

A world-class, high growth, defensible Injectables business

Through consistent delivery of profitable growth, we are striving to become **second** largest injectables company by value and volume in the US, **top five** injectables company in Europe and **leading supplier** of injectables in MENA.



Pipeline

- Utilise global network of R&D sites (US, Croatia, Jordan) to increase productivity and complexity
- Endeavor to become leader in ease-of-use products that bring value to customers
- Further enhance our pipeline through value-creating partnerships



- Maintain best in class manufacturing footprint, with industry leading capabilities and quality
- Expand, and in some cases establish, manufacturing capacity across our markets, focusing on adding new technologies and automation



- US reorganised teams into three focus areas: base business, specialty and 503B
- EU expand share through market expansion and tailored R&D for local needs
- MENA leverage local sites to strengthen market access and increase commercial specialisation to support differentiated portfolio

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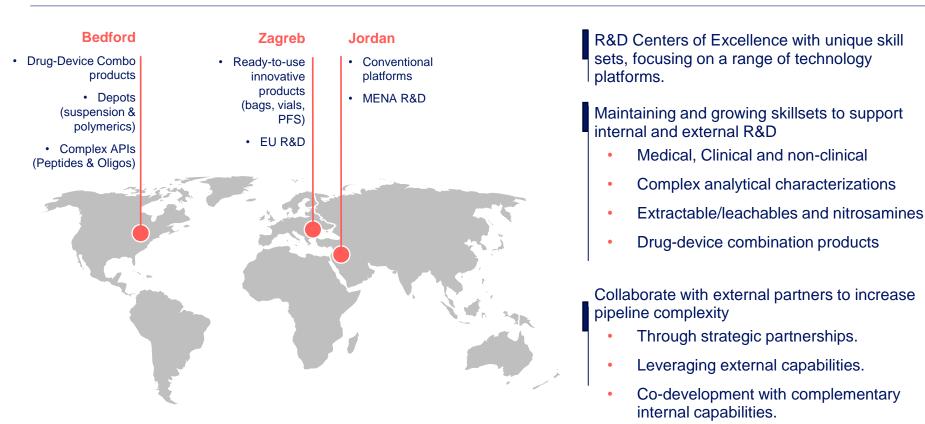
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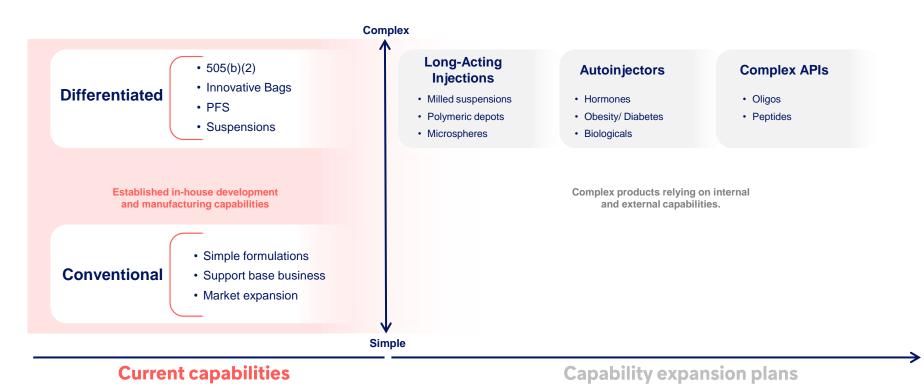
A global footprint with a skilled team of R&D experts



R&D strategic pillars

Strategic pillars	Strategy in action	Diversified pipeline of >160 projects globally ¹	
Increase breadth	Continue to expand pipeline of conventional products, which enhance portfolio breadth.	Differentiated BLA Complex	
Increased focus on RTU products	 Adding differentiated Ready to Use (RTU) products through Zagreb R&D to bring value to our end-users. 	^{3%} 17%	
	 Maximise our manufacturing capabilities in bags and PFS. 		
	 Over 20 RTU products currently in development, including 10 aseptic bag products. 		
Utilise global reach	Leverage global portfolios and R&D capabilities for market expansion.		
	Enhance ex-US pipeline.		
Enhance differentiation through partnership	 Complement internal capabilities through partnerships, including co- development and/or licensing opportunities. 	64% Conventional	
	Peptides, Oligonucleotides, Biosimilars.	Conventional	
	 Long-acting Depots, Drug-Device combo products. 		
	Sharing Risk, Expertise and Cost.	1. Pipeline projects are by market	

Expanding our R&D capabilities by adding new technologies



Increased emphasis on complex injectables to add sustainable value

Ease of use

Ready-to-Use

- Vancomycin RTU (launched)
- Daptomycin RT (launched)
- Autoinjectors
 - Liraglutide (launched)
 - Oils/Hormonals (development)

Long acting depots

- Suspensions
 - Triamcinolone, Betamethasone, methylprednisolone, medroxyprogesterone (approved)
 - Milled suspensions (development) –market size: >\$2bn. From 2032.
- Microspheres
 - Oncology (filed)
- Polymeric Depots
 - Oncology (development)

Complex molecules

- Biosimilars
 - Ustekinumab (filed)
 - Denosumab (filed)
- Peptides
 - Cardiovascular (filed)
- Oligonucleotides
 - Partner identified

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Our global footprint is expanding as we invest in increasing capacity and enhancing capabilities to support future growth embitions

Injectables Manufacturing sites (Portugal and Cherry Hill being the largest sites in the Injectable Division)



These investments allow us to meet customer and patient needs, enhance our quality, efficiency and flexibility and deliver on our growth ambitions

Local capacity and advanced manufacturing technologies enable us to respond to increase in market demand and customer needs

	Europe	United States	MENA		
Small Liquid Parenteral	٠	•	•	- Furana	
Terminal sterilization bags	٠	•		Europe Injectable Hub with all technologies being	
Aseptic bags		•		Portugal is the largest Injectable facility	
Lyophilization	٠	•		02 United States	
High Containment	•			Producing for the US in the USA	
Large Liquid Parenteral	٠	•		03 MENA Meets the localization requirements and tag	
Aseptic Powder Filling	٠			into fast growing markets, largest investm in Saudi Arabia	
Suspensions	٠				
Emulsions		٠			
Pre-filled syringes	٠				
Serviced Market	US, MENA, EU	US	MENA	Available Available Available and expanding	

We are expanding capacity and adding advanced technologies to drive high quality growth

Capacity expansion:

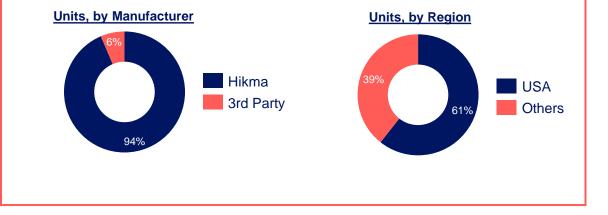
- Increasing liquid filling capacity by c.20% through addition of new lines
- Lyophilization capacity (measured by shelf space) will triple
- We currently operate 266 sqm of lyo:
 - The Bedford expansion (Xellia) will add 180 additional sqm
 - Portugal expansion projects will add 160 additional sqm

New technologies

- Additional technology added through Xellia acquisition – aseptic bag filling
- Aseptic filling technology will support the current pipeline and Ready to Administer (RTA) strategy

Injectables Manufacturing

- US Injectables products are largely produced at Hikma sites, with just 6% produced by 3rd parties
- · 61% of Hikma injectables volumes sold in the US are manufactured in the US



Leverage core expertise, high efficient facilities and global presence to attract CMO customers

As we investing in expanding our manufacturing footprint, and with the additional capacity coming online, we are in active discussions with several potential future partners to sell the extra capacity while transferring our own products into the market.

Our Injectables CMO offering:



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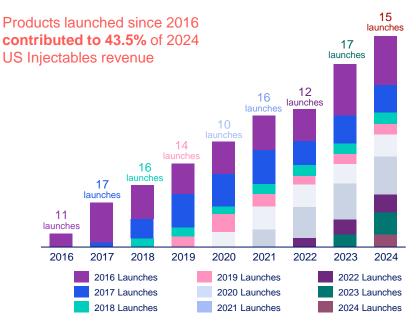
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A top supplier with great relationships and broad portfolio

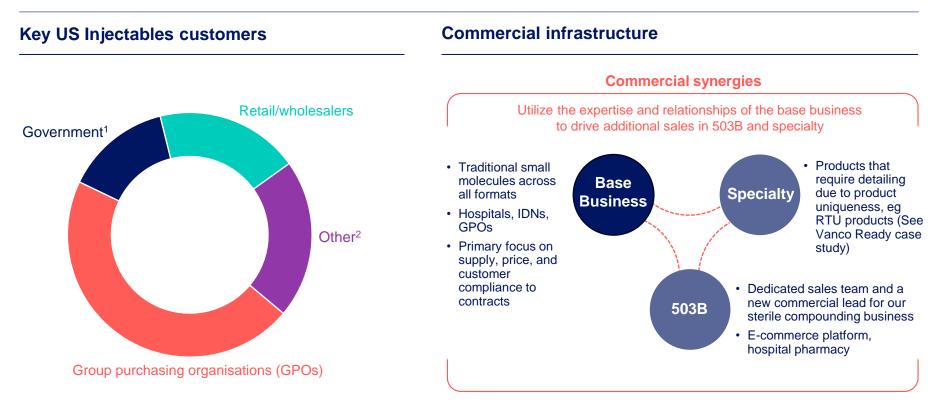
Top 3 supplier by volume Pfizer Top three manufacturers >100 active make up **US** generic 50% injectable Fresenius suppliers of the market Hikma

A broad and expanding portfolio



¹ IQVIA MAT through December 2024, generic injectable volumes by eaches, excluding branded generics, Becton Dickinson and Henry Schein

Customer-centric go-to-market approach



Expanding our portfolio with dosage forms that bring value to our customers

Broad portfolio of products today + planned RTDs

- Top three injectable portfolio in the US market
- Over 170 medications offered in multiple formats
- Addressing over 60 therapeutic categories
- Adding Ready-to-dilute (RTD) formulations



Ready-to-administer (RTA) formulations

- Improves efficiency in hospitals minimizes steps to administration of our products
- Reduces risk of medication errors
- Lessens need for in-house or external compounding services



Case study: VANCO READY – the first and only vancomycin RTU bag, with significant benefits vs. existing formulations. Patent protected to 2035

Product profile

- The first and only commercially available ready-to-infuse solution of vancomycin with no need to compound, dilute, activate or thaw
- Developed by the Croatian R&D team
- Received FDA approval in 2019 with a black box warning against use for patients in early pregnancy
- The product has been reformulated and resubmitted and we expect FDA approval with no black box later in 2025



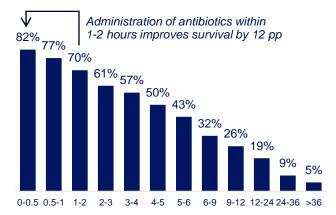
VANCO READY is available in 7 doses covering the needs for most patients

Benefits

- No need for dedicated compounding staff or nurses for preparation
- >12-month shelf life at room temperature allows storage in medical dispensing cabinets and minimizes waste
- Streamlined supplier management, patient operations and training for medical staff
- GMP-status product with lower potential risk for compounding errors vs. inhouse or 503b suppliers
- Reduced time to administration for hospitals leading to improved survival rate for sepsis patients

Survival rate of sepsis patients by time to antimicrobial administration

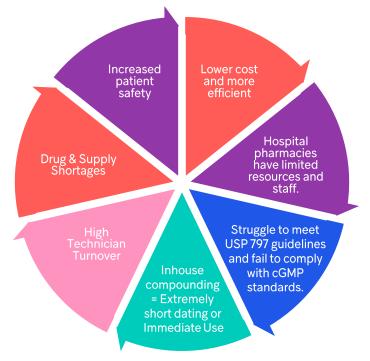
Time calculated as hours between onset of hypotension and IV administration of drug



Source: 1. Kumar A, Roberts D, Wood KE, et al. Duration of hypotension before initiation of effective antimicrobial therapy is the critical determinant of survival in human septic shock. Crit Care Med. 2006;34(6):1589-96 2. Carroll D, Popa A, Hejal R, et al. 1601: Evaluation of timing of first dose antibiotic administration in patients with sepsis or septic shock. Crit Care Med. 2019;47(1):776.

503B Sterile Compounding – a compelling future growth opportunity

Why 503B is important to customers



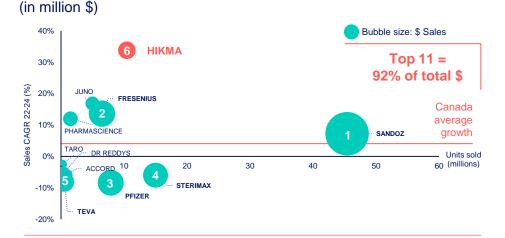
Hikma's unique offering – A vertically integrated 503B compounder, backed by a sterile manufacturing company

- Hikma purchased its Dayton 503B site in 2020 and has upgraded the facility to include 19 compounding suites, two state-of-the-art VHP chambers, and a fully revamped lab room
- Fully FDA inspected and DEA audited with approval in 46 key states
- Dedicated sales team
- Ability to leverage core Hikma capabilities and expertise:
 - Track record in cGMP compliance
 - Extensive access to raw materials
 - Established relationships with hospital customers
 - Quality track record

503B Today:

- Increasing month-on-month growth as new customers are added and portfolio develops
- Expect to be profitable by end of 2025
- Currently reported in 'Others' line

Hikma in Canada – leading growth in the generic hospital market



2024 generic Injectable sales (ex biotech)^{1,2,3} top 11

Top Canadian incumbents are underperforming

Canadian incumbents either growing marginally or declining in the Gx space.
This is due to an increasing focus on biosimilars as opposed to generics.

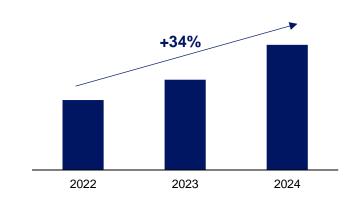
Hikma's strong growth trajectory

- Became the 6th largest company in the Canadian Injectable market by revenue
- With a 34% year-over-year growth, Hikma, surpassed by far, the market average

Hikma well positioned

 Companies smaller than Hikma, some of them virtual and focused on low-volume products

Hikma historical sales growth





đ

Consistent Double-digit Growth

Reflects Hikma's effective launch management, agility responding to market shortages and strong market expertise.

Strategic Focus on Expansion

Momentum is expected to continue as Hikma strengthens its position with new product launches and asset acquisition.

On track to enter Top 5



Hikma's robust supply chain and portfolio expansion are expected to promote our position as a top 5 injectables player

Hikma Pharmaceuticals PLC

(1) IQVIA data December MAT 2024; (2) Pfizer/ Hospira includes Brand and Generics (3) Rest of the market \$62M

Agenda

Introduction

2. R&D

3 Manufacturing

4 • North America Commercial

5 • Europe

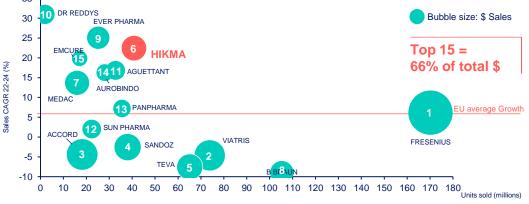
6 • Wrap-up

Europe – Hikma showing strong growth in this large and highly fragmented market

(in billion \$) 10 15 20 25 30 35 40 45 50 55 Germany France UK Italy Spain Hikma coverage = 83% of Europe vol* Nordics Poland Netherlands Belgium Austria Portuga Rest of EU* Currently Operating New Markets Planned Expansion

European Pharma Market – Total Value¹

2024 Generic Injectable Sales (ex Biotech)^{2,3} – TOP 15 (in million \$)



Top 5 EU companies are underperforming

- · European market leaders are growing below the EU average or declining
- This is largely due to a strategic focus on biosimilars and divestment from standard generics

Hikma's strong growth trajectory

- Became the 6th largest player in the EU Injectable Hospital market in 2024
- With 22% year-over-year growth, Hikma well above the market average growth rate

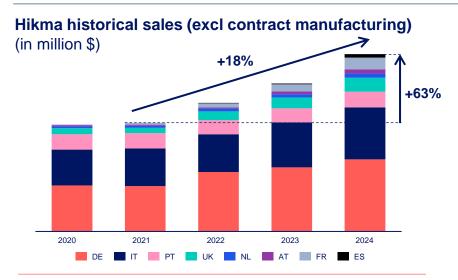
Well positioned

5

- · Several of the top 15 players are growing through niche and lower-volume products
- Several competitors are lacking the scale of being a Pan European player.

¹ EFPIA 2024 Data (*except Russia)
 Hikma Pharmaceuticals PLC
 ² Central and Western Europe, Nordics and Poland
 ³ IQVIA Injectable generic products, Hospital + Germany Retail, 2024 USD sales

Hikma's growing presence across Europe



Consistent double-digit growth

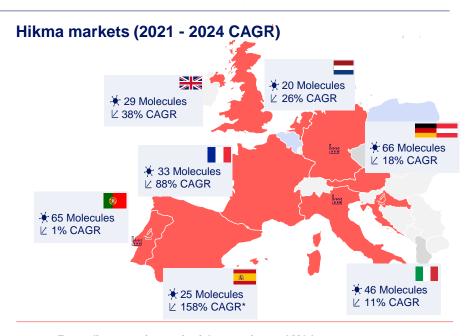
Reflects Hikma's efficient supply chain, agility in responding to market shortages, and focus on higher-revenue, more profitable channels.

Strategic focus on newly launched territories

Momentum is expected to continue as Hikma strengthens its presence in France and Spain and enters new markets.

On track to enter the top 5

Sustained growth and expansion position Hikma to surpass key competitors and enter EU top 5 generic injectable hospital players in the next 24 months.





Expanding capacity, territorial expansion and M&A Advancing into Poland and Belgium while pursuing M&A to accelerate

EU dedicated R&D



The recently acquired R&D center in Zagreb will allow the introduction of strong additions to our EU portfolio and sustain our growth momentum in the region.

Agenda

Introduction

2. R&D

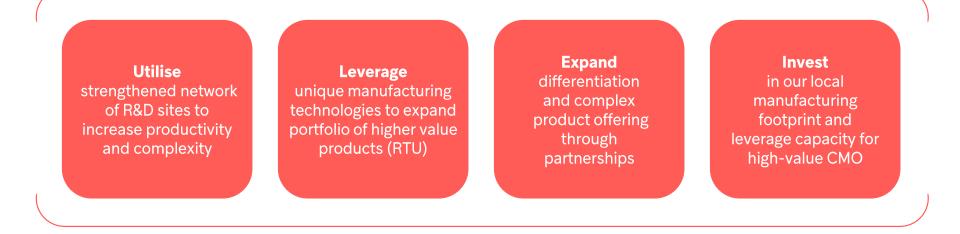
3 Manufacturing

4 • North America Commercial

5 • Europe

6 Wrap-up

Our strategy sets us up for continued success and supports our plan to deliver consistent profitable growth and our ambition to become leaders in all our markets



Better health. Within reach. Every day.

Hikma US Meet the Management Group

Our strategy for growth



Enhance

operational efficiencies and embrace new technologies, maintaining our high-quality levels

Leverage

our broad portfolio and strong commercial capabilities

Develop

a more differentiated pipeline

Expand

into adjacent businesses and geographies

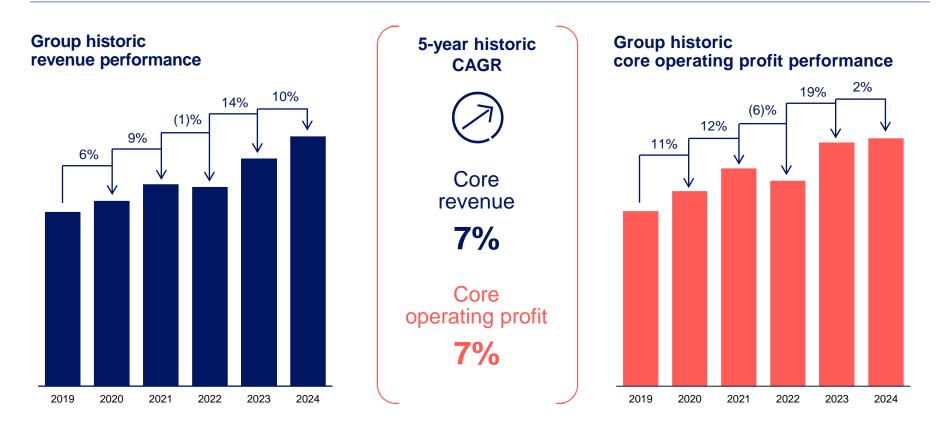
Empower

our people and cultivate a unified culture

Act

responsibly across our local markets and communities

Our growth over the last 5 years has been supported by our strong foundation



Our clear strategy supports our future growth ambitions



Efficiency without compromise on quality

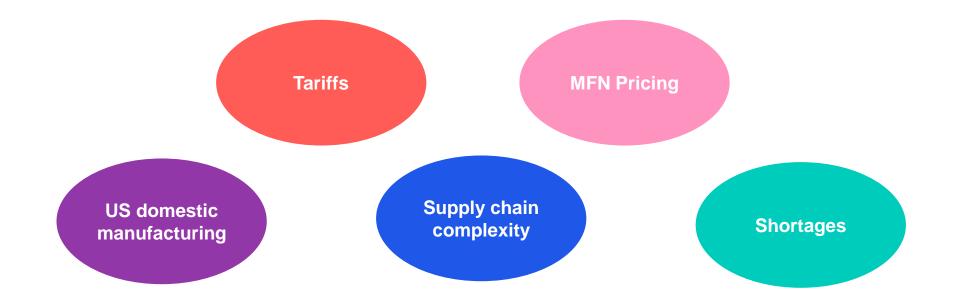


Unlocking value through synergies across the organisation

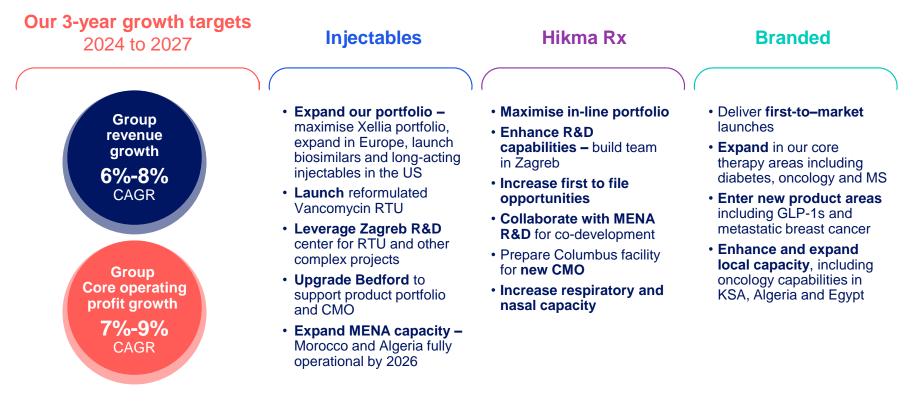
Operating sustainably to ensure we can deliver our purpose

Engaging in strategic partnerships to accelerate growth

Navigating uncertainty and balancing opportunities and risks



Our focus on execution will deliver our medium-term targets



Our strategic plans, underpinned by significant investment, will enable us to achieve our Group revenue target of \$5 billion in 2030

- Invest 5%-6% of Group revenue on R&D
- Extract synergies across our global R&D sites
- Add complexity including RTU, nasal, inhalation
- Leverage Zagreb to accelerate EU submissions
 - **Investing in R&D**

- Focusing on developing CMO as a core competency across sites and business segments
- Leveraging Group manufacturing expertise

Growing CMO

Group revenue of **\$5 billion** in 2030

- Building compounding
- Strong contribution from 503B
 business
- Leveraging our position as vertically integrated compounder with sterile manufacturing expertise

 Build integrated commercial team to support specialty launches - reformulated Vanco RTU, 505(b)(2)s, Epi Nasal Spray

Launching complex products and specialty brands

Partnering

 Benefitting from licensing partnerships for innovative products and healthcare solutions, e.g. SK Bio, Guardant, Rakuten

Expanding capacity

- Investing >\$1.5 billion in CAPEX to enhance and expand capacity
- Significant US domestic capacity, including complex aseptic bag filling and extensive LYO capacity

Expanding geographically

- Increased scale in EU, covering over 80% of the market by volume
- Expanding Hikma Rx portfolio into Canada

