



Corporate Presentation

October 2025

**Changing the Treatment
Paradigm for Patients with Iron
Deficiency Anemia**



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Shield Therapeutics

Fast Growing, Mission Driven, Specialty Pharmaceutical Company

Large global iron deficiency/iron deficiency anemia (ID/IDA) market ripe for disruption

- +15 million iron deficient individuals in the US with and without anemia
- 40-60% of patients on traditional irons often lead to discontinuation due to intolerable GI side effects or insufficient efficacy

ACCRUFER®/FeRACCRU® (ferric maltol) set to be the oral iron treatment of choice

- Approved by the FDA, EMA, TGA, Swiss Medic, and Health Canada as the only prescription oral iron indicated for the treatment of ID/IDA
- Highly tolerable proprietary oral formulation that offers a low side effect profile, distinguishing itself from conventional iron treatments

Poised to turn cash flow positive by end of 2025

-  • A strengthened balance sheet with enough cash to get to turning cash flow positive by the end of 2025
- Peak revenue potential of ACCRUFER® of ~\$450M², Strong IP through 2035



Experienced Executive Team with extensive US commercialization expertise



Anders Lundstrom
CEO



Santosh Shanbhag
CFO



Lucy Huntington-Bailey
General Counsel



Andy Hurley
Chief Commercial Officer



David Childs
VP, Manufacturing and
Strategic Alliance



Dr. Jackie Mitchell
VP, Quality, Clinical and
Regulatory Affairs



Iron Deficiency with & without Anemia (ID/IDA)

Universal problem: HCP's are struggling to treat IDA because patients can't tolerate the GI side effects of oral iron salts



Oral ferrous salts dissociate in the stomach. Unabsorbed iron (Fe⁺) generates reactive oxidative species (**ROS**), causing irritation and damage to the intestinal lining **and gastrointestinal (GI) side effects**

Up to 70% of patients can experience GI related side effects^{1,2} including bloating, dark stool, nausea, distention

Up to 60% of patients will discontinue treatment with ferrous (iron) salts primarily due to GI adverse events and lack of effectiveness³

ACCRUFeR® designed for efficacy and tolerability

Unique MOA (mechanism of action) Shields and Delivers Elemental Iron to the Small Intestine ^{1, 2}

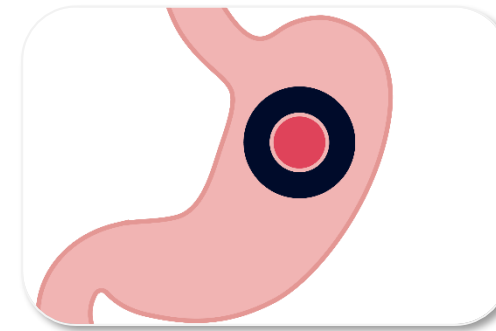
Proprietary formulation

ACCRUFeR® is formulated in a maltol complex vs. traditional oral irons, provided in ferrous-based formulations

Low iron dose

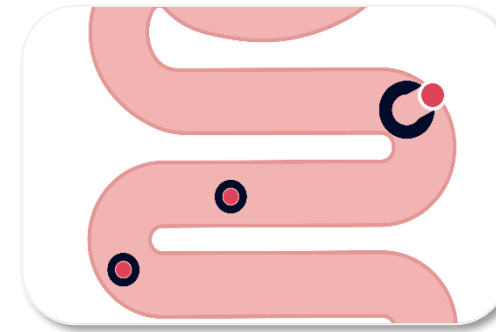
60 mg of elemental iron is delivered by ACCRUFeR® daily

ACCRUFeR® remains tightly bound in the stomach



The unique Maltol Shield™ protects iron from the stomach, remaining tightly bound as it passes through

Dissociates upon uptake in the duodenum



Iron remains bioavailable, chelated, and ready to replenish iron stores.

Excess iron is excreted in the stool

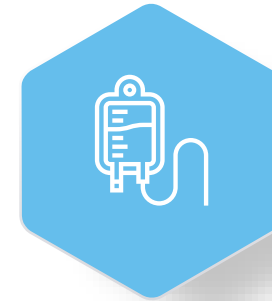
1. ACCRUFeR™ is dosed at 30mg BID, MOA = mechanism of action
2. ACCRUFeR® (ferric maltol) [Prescribing Information]. Austin, TX: Shield Therapeutics, 2019. Revised 02/22.

Significant window of opportunity exists for ACCRUFeR®



Oral Iron

>90% ferrous salts²



IV Iron

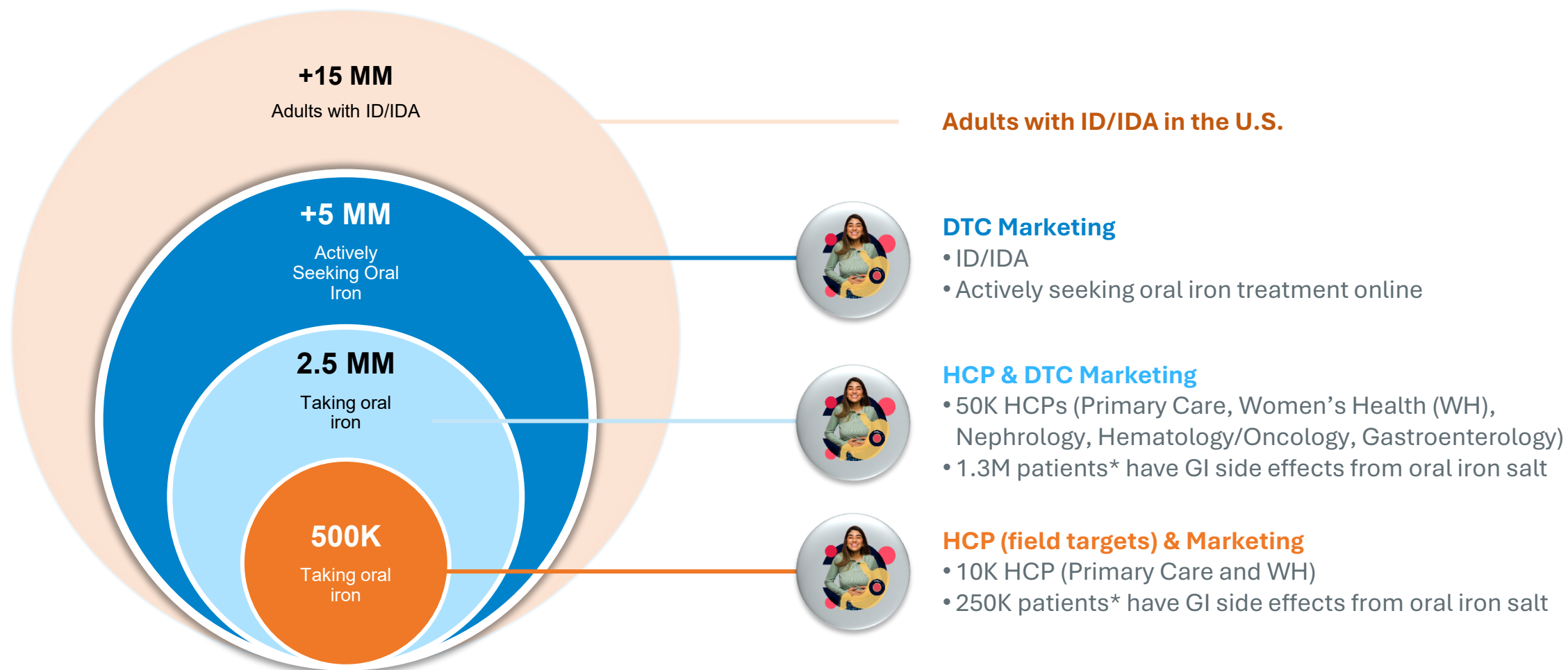
**Iron replacement that patients
will actually take**

*A well tolerated oral iron that
effectively normalizes and maintains
Hb, ferritin, and TSAT levels¹*

ACCRUFer®/FeRACCRU® (ferric maltol) portfolio – Global footprint

disease	phase 1	phase 2	phase 3	regulatory	marketed		market opportunity	Partner
ID/IDA (adults)							15M Adults with ID/IDA ³	 VIATRIS [™] co-promote
							6-7% of population have ID ⁴	 KYE Pharmaceuticals
							Prevalence in DE: ~3.3% of ID/IDA ⁵ , UK: ~8%/3% (women/men) IDA ⁶	 NORGINE
							5.2M people with ID/IDA ⁷	 한국파마 KOREA PHARMA
							15% prevalence of ID/IDA ⁸	 奥赛康
ID/IDA (pediatrics)							~1.0M ⁸ kids with ID/IDA ⁹	 VIATRIS [™] co-promote
							TBD	 NORGINE
Pulmonary Arterial Hypertension (PAH)							\$230M market in 2024 ¹⁰	 想いをつなぐ。明日へ進む。 バイタルネット

While ID/IDA is vast, reaching patients who have the urgency to treat is critical to the success of ACCRUFerR®



2024 HCP and Patient Quant studies, 2025 Digital Online analysis

*Patients taking Oral Iron Salts and have GI Side Effects estimated to be 40%-60% (Cancelo-Hidalgo MJ, et al. Curr Med Res Opin. 2013;29(4):291-303 and HCP and patient quant 2024

Consumer related marketing efforts rapidly increasing ACCRUFER® awareness

Social Media Influencers

Reached +1M patients



Sia Clyde
2.4M Followers



Krystal Nielson
565K Followers



Ashley Lemieux
535K Followers



Brittney Fusilier
409K Followers



Dr. Franziska Haydane
313K Followers



Louie Chan
125K Followers



Elisabeth Akinwale
160K Followers

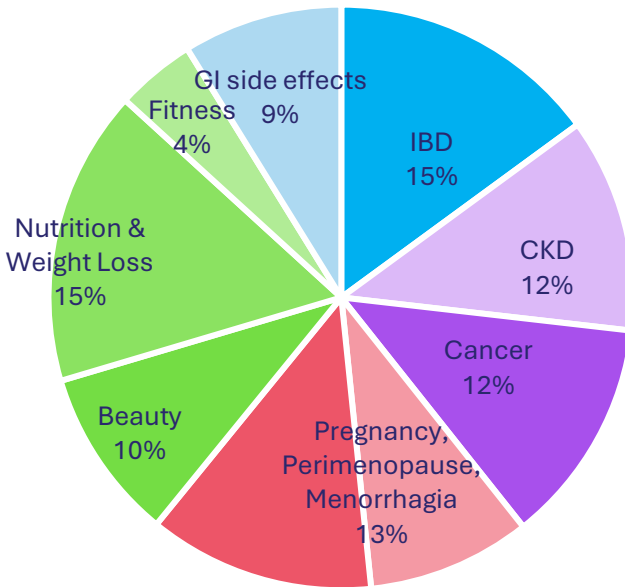


Onyi Azih
121K Followers

Top Social Media Channels and Patient Communities



Meta

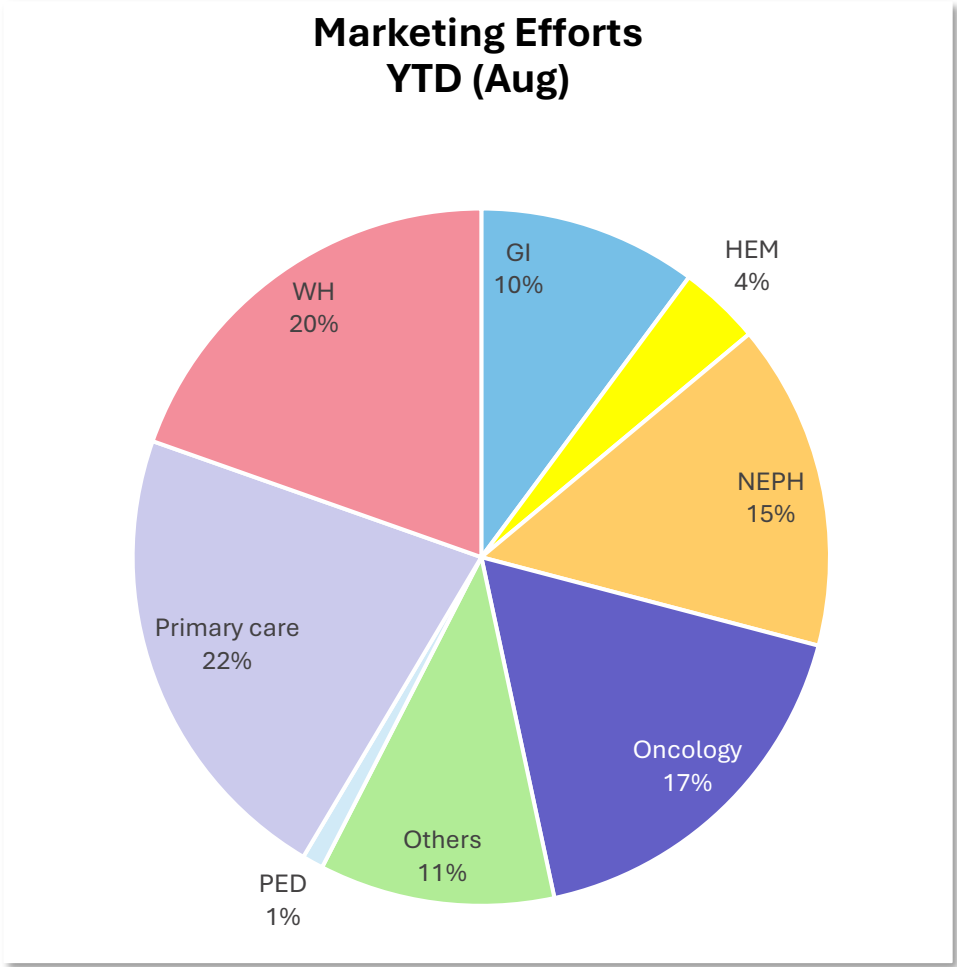
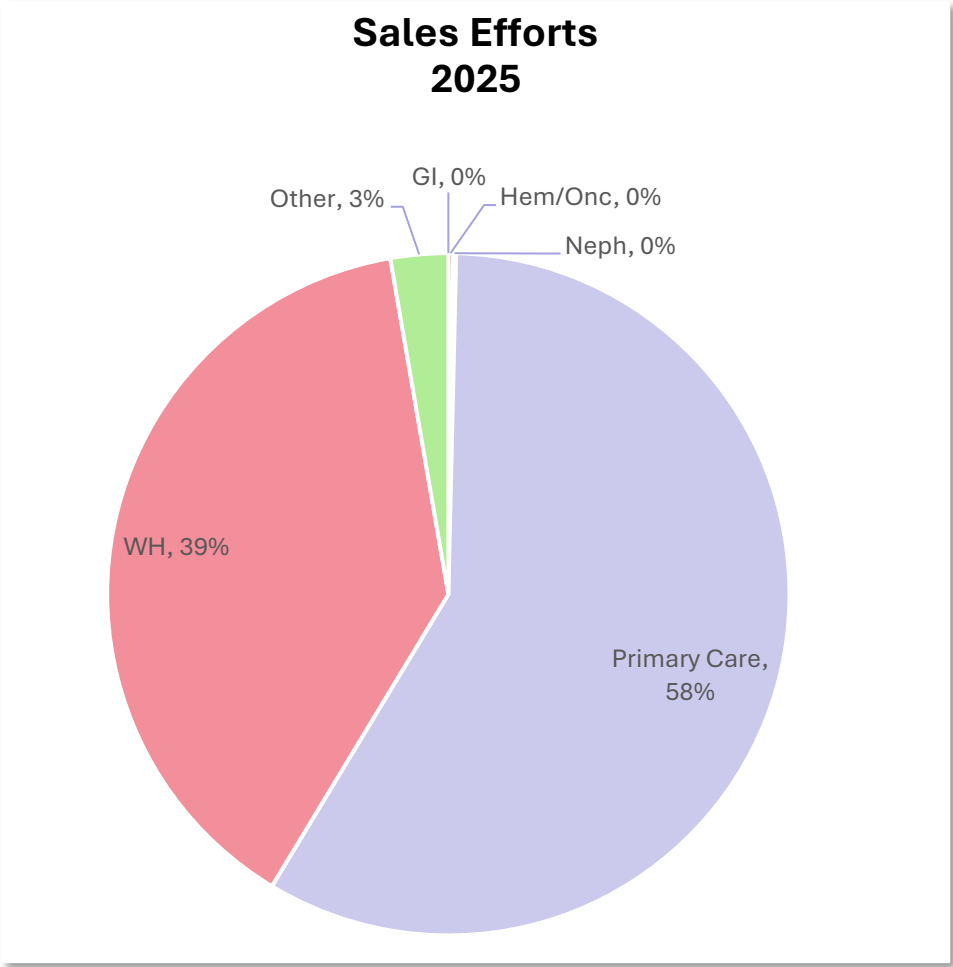


Digital Advertising

Reached +3.9M patients

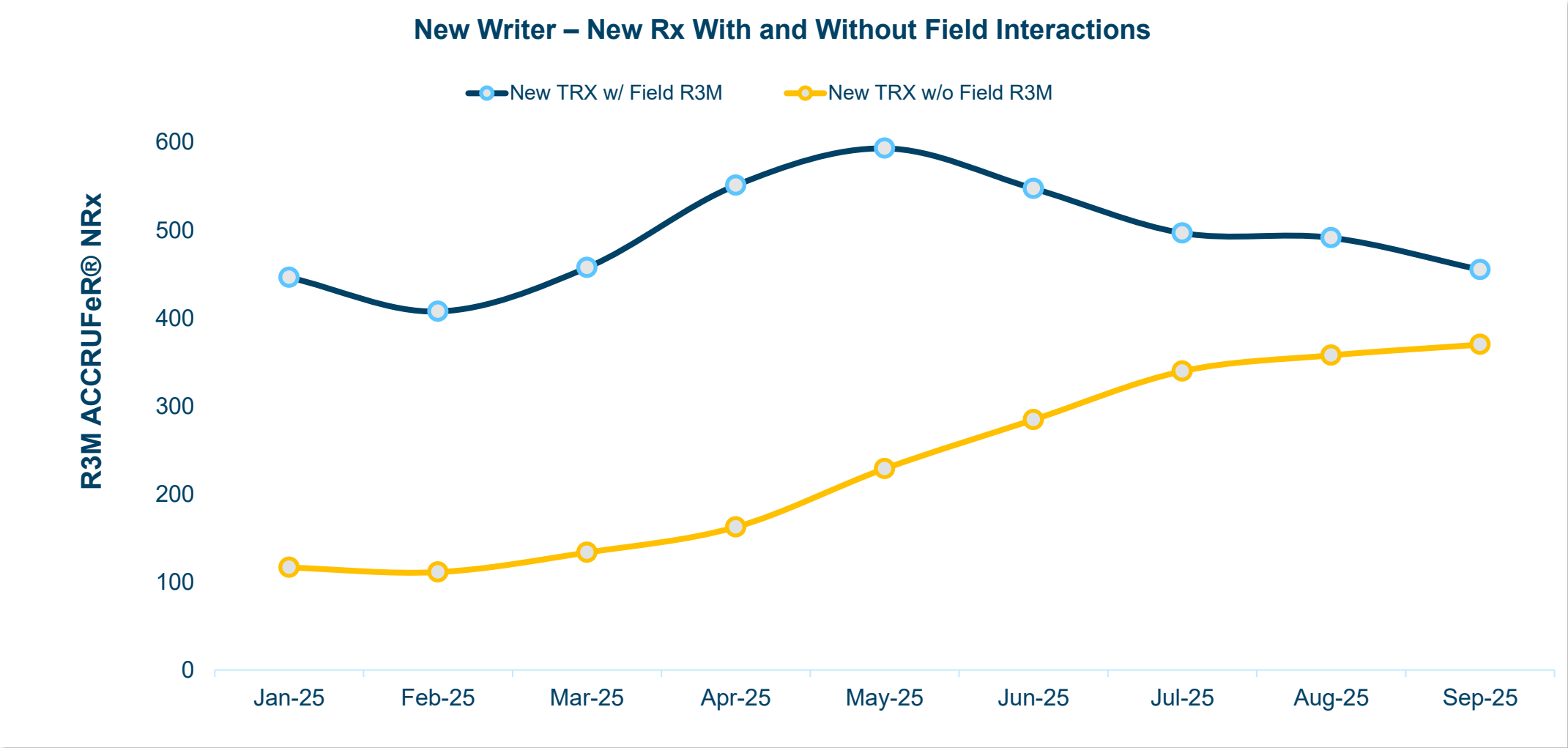


Physician focused marketing efforts has expanded reach into specialists



WH = Women’s Health; HEM = Hematology; GI = Gastroenterology; NEPH = Nephrology; ONC = Oncology

Early indicators show positive impact of digital strategy on Rx by new writers



Key ACCRUFeR® initiatives building ACCRUFeR® momentum



Increase awareness of ACCRUFeR®

- Patients and HCPs via digital marketing/initiatives
 - Endemic Ads and Rollout of Influencers within patient-focused social media channels
 - Significant increase in EMR banner marketing to providers



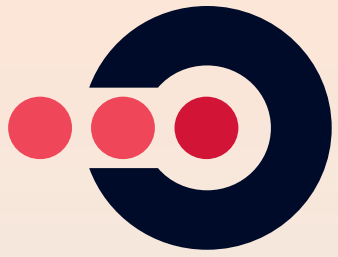
Enhance Sales Force Output

- Redesign of sales territories has resulted in strong prescription growth
- Continue to Drive Demand in Top 6 States (NY, CA, TX, FL, GA, NC)



Increase Prior Authorization (PA) submissions, PA approval rate & Covered Rx Fill rate

- Focus on more PA's submissions
- Contract with additional pharmacies that have robust PA approval process support for offices
- Leverage Case Managers to contact patients to pick up unfilled prescriptions



ACCRUFER[®]
(ferric maltol) *30 mg capsules*

#1 branded prescription oral iron
in the US ID/IDA* market

Source: IQVIA Xponent PlanTrak

*ID/IDA: Iron Deficiency with & without Anemia

2025 Business Priorities

**Grow ACCRUFer®
Net Revenues**

Q3 25

**#1 branded prescription oral iron
in the US ID/IDA market**

~\$13.1M ACCRUFer® Net Revenues
~54K TRx with ~22% consignment¹
~\$237 Net Selling Price in Q2 25

**Turn Cash Flow Positive
by End of 2025**

Q3 25

\$8.6M cash and cash equivalents
Net burn of \$3M excluding \$2.0M placing

~\$2.0M placing
@7.5p/share, 5% premium to 30D VWAP

On track to turn cash flow positive in Q4 2025

**Launch in Canada, and
execute regulatory
process in Korea,
China, and the
Pediatric Population**

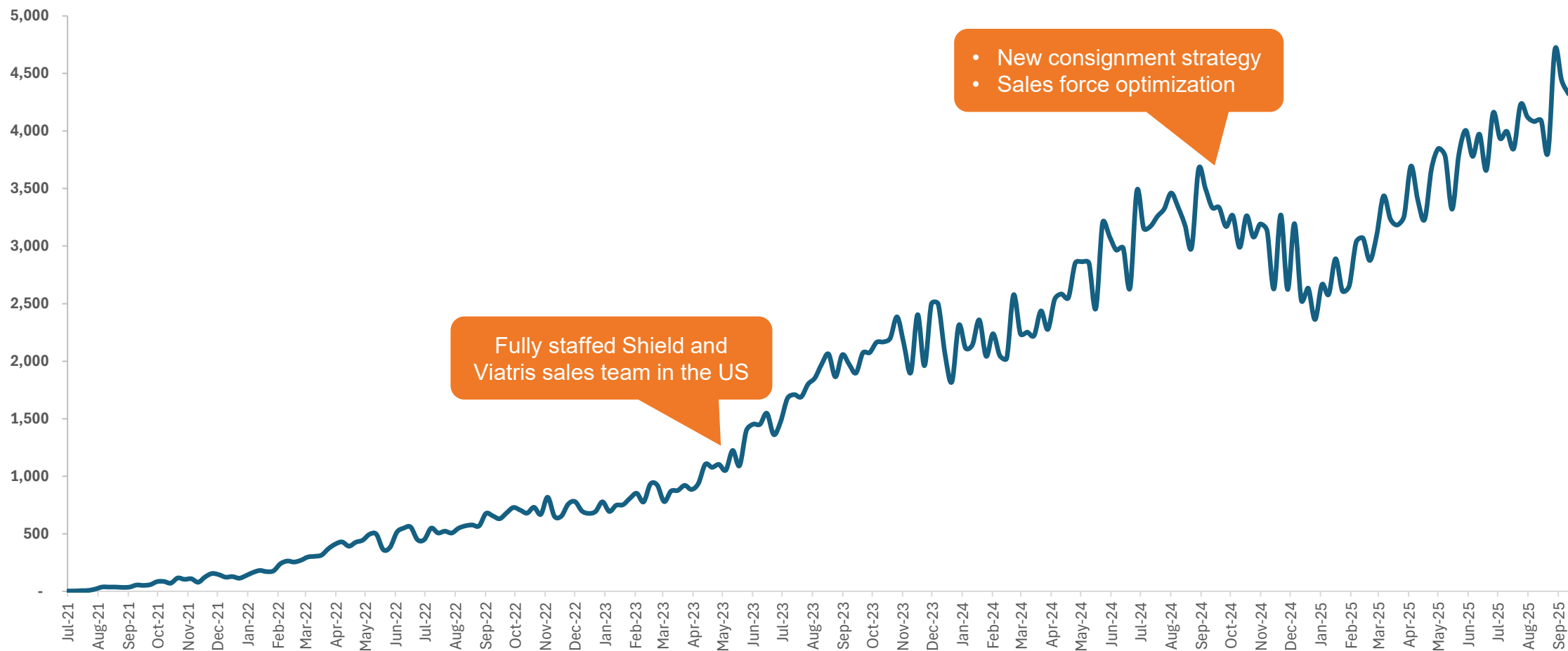
Q3 25

**Priority Review by FDA for ACCRUFer®
children with iron deficiency anemia (IDA)**

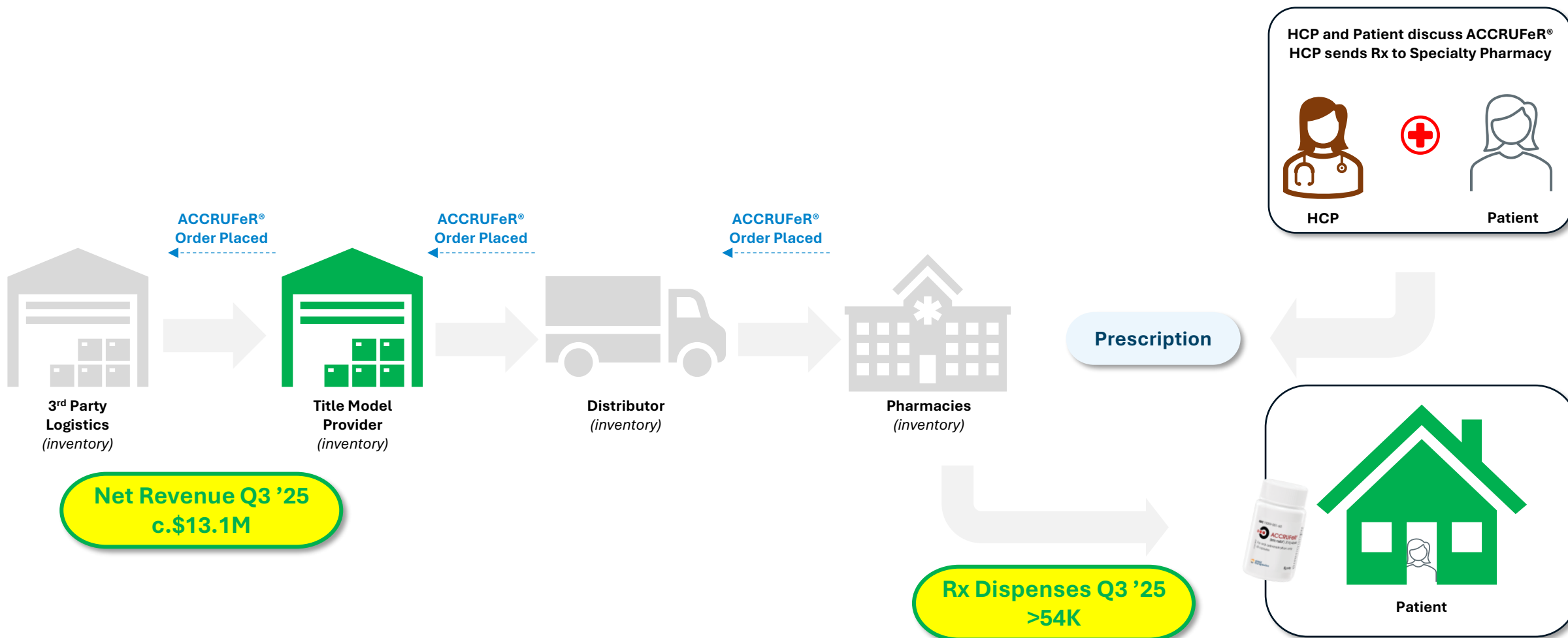
**Positive efficacy and safety results for
ACCRUFer® in Heart Failure (HF) and Iron
Deficiency Anemia (IDA)**

ACCRUFer® prescriptions demonstrating significant uptake since launch

ACCRUFer® Weekly US TRx

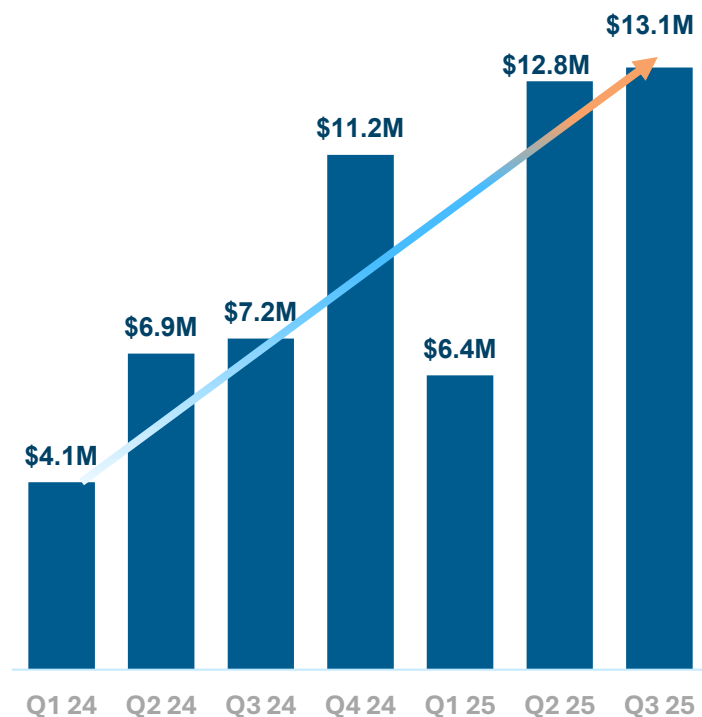


ACCRUFer® Distribution in the US

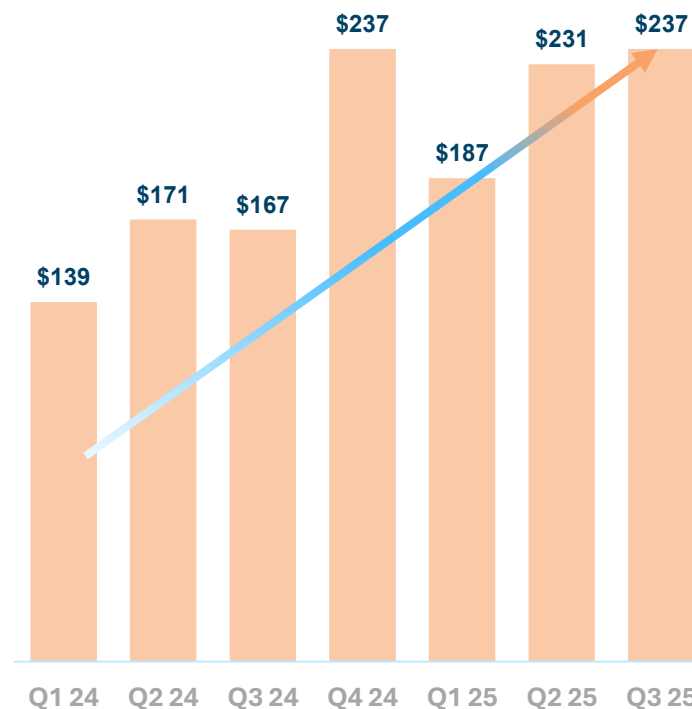


Continued growth in ACCRUFer® in the US in Q3 2025

ACCRUFer® Net Revenues
Quarterly

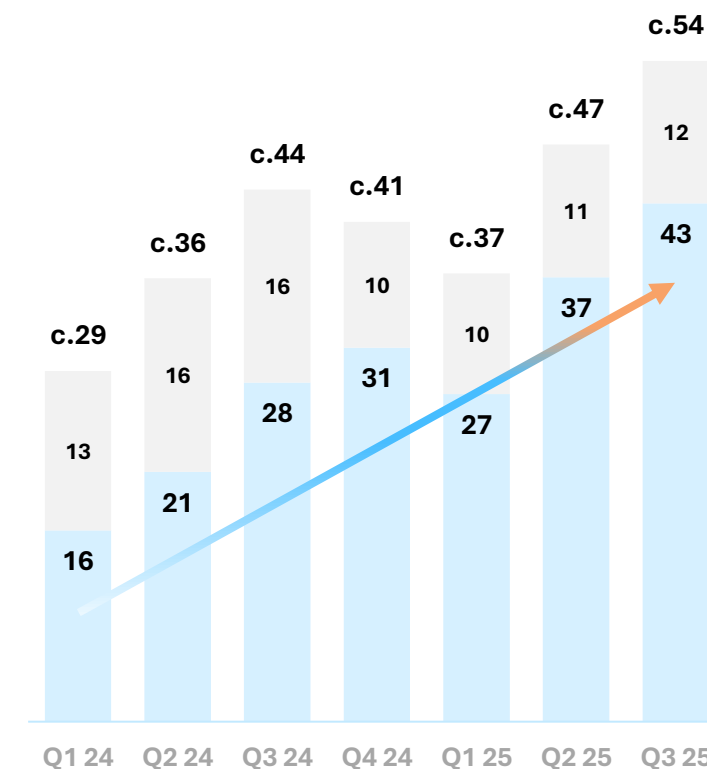


Net Price Per Script
Quarterly

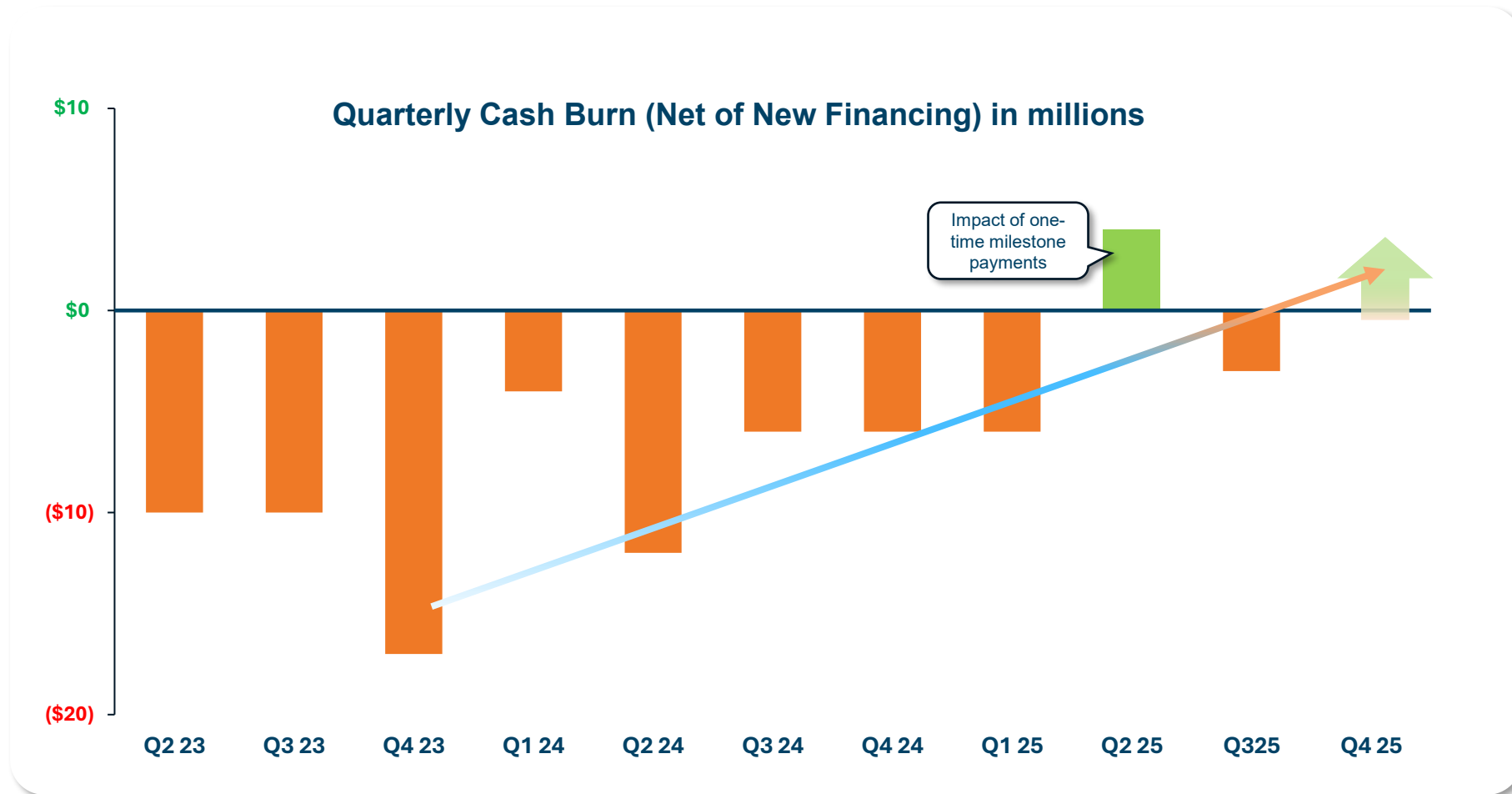


Total Prescriptions
Quarterly

■ Retail ■ Consignment



On track to turn cash flow positive in Q4 2025 driven by continued ACCRUFer[®] growth in the US and effective working capital management



Shield Therapeutics

Fast Growing, Mission Driven, Specialty Pharmaceutical Company



- **Vast market opportunity with significant revenue potential**
- **Shield-Viatris US partnership driving growth in ACCRUFER® prescriptions, net revenue and net selling price**
- **Global partnerships continue to progress at a steady pace with anticipated milestones and double-digit royalties**
- **Goal to be cash flow positive by end of 2025**
- **Looking for partnership opportunities to expand the portfolio**

Delivering on the brand promise



Thank You!

Anders Lundstrom – Chief Executive Officer
Santosh Shanbhag – Chief Financial Officer

www.shieldtherapeutics.com

