

Corporate Presentation

July 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

- 40-60% of patients on traditional irons often lead to discontinuation due to intolerable GI side effects or insufficient efficacy¹
- 15-20 million iron deficient individuals in the US with and without anemia



ACCRUFeR®/FeRACCRU® (ferric maltol) set to the 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



- 1. Cancelo-Hidalgo MJ, et al. Curr Med Res Opin. 2013;29(4):291-303
- 2. Shield management estimate

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**



















Capgemini







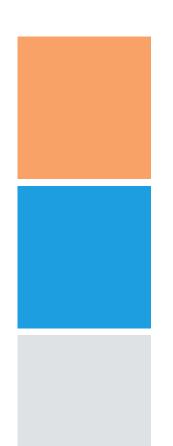






Iron Deficiency with & without Anemia (ID/IDA)

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

Patients comment: "Side effects of oral iron worse than the symptoms of IDA"

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

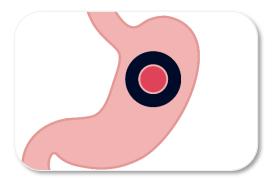
ACCRUFER® is formulated in a maltol complex vs. traditional oral irons, provided in ferrous-based formulations

Low iron dose

60 mg of elemental iron is delivered by ACCRUFeR® daily

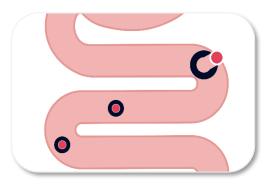
- 1. ACCRUFeR[™] is dosed at 30mg BID, MOA = mechanism of action
- ACCRUFeR® (ferric maltol) [Prescribing Information]. Austin, TX: Shield Therapeutics, 2019. Revised 02/22.

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

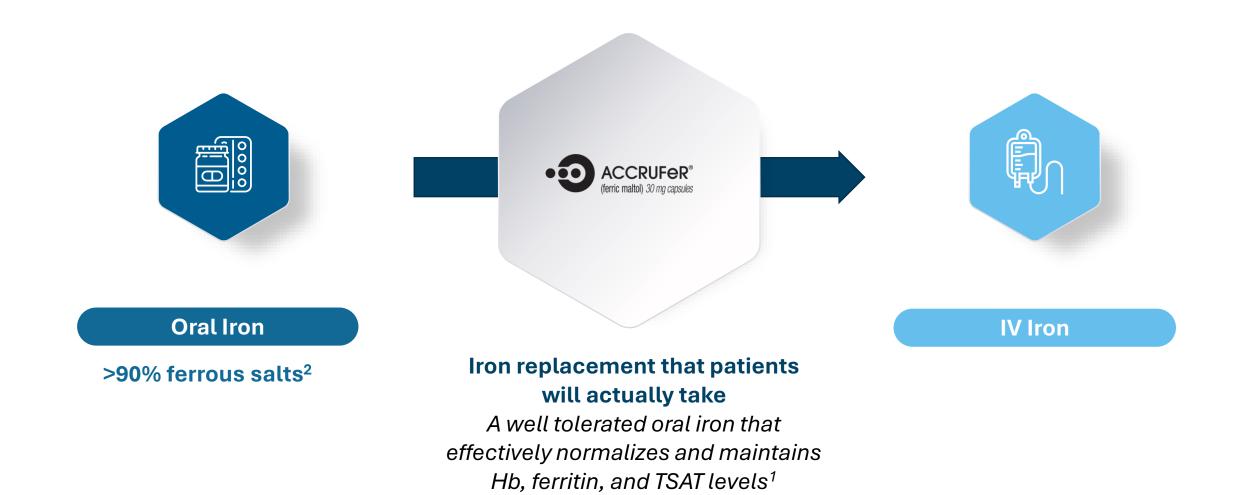


Iron remains bioavailable, chelated, and ready to replenish iron stores.

Excess iron is excreted in the stool



Significant window of opportunity exists for ACCRUFeR®





Global partnerships continue to progress

Deals include upfronts, milestones & double-digit royalties



United States

Co-Commercial Agreement

\$30m in available sales milestones



EU+1

Commercialized across Europe

€1m Pediatric EU Approval milestone

Double-digit royalties on net sales



Canada

Launched in Canada in Q1 2025

Revenue-based milestone payments

Double-digit royalties on net sales



Republic of Korea

Filed for approval; Pending successful review, approval anticipated in 2025

Revenue-based milestone payments

Double-digit royalties on net sales



China +2

Phase 3 Study completed Approval expected in H2 2026

Approval and
Revenue-based
milestone payments

Double-digit royalties on net sales



Japan

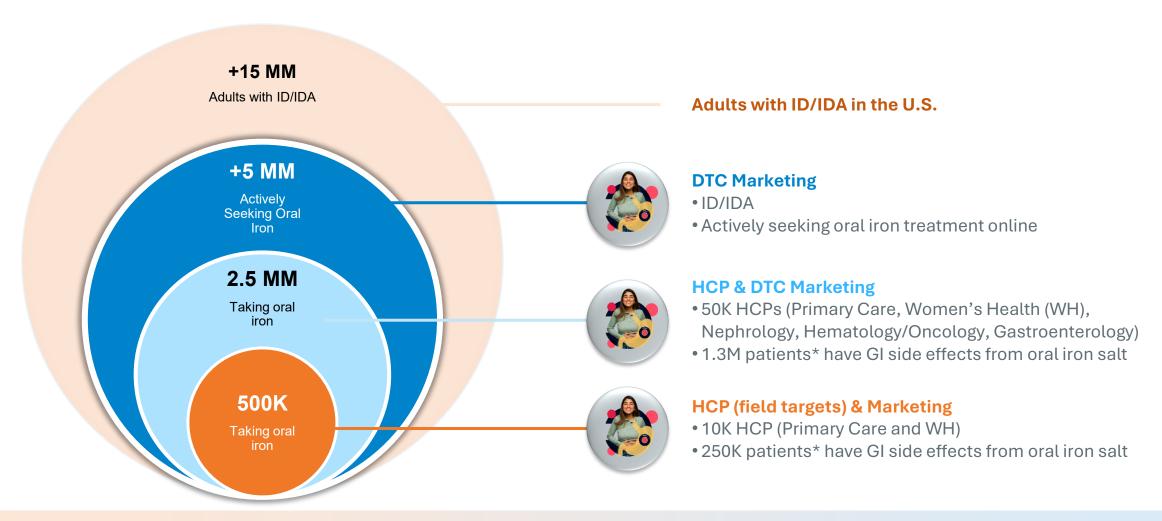
Deal signed in April 2025

Approval and Revenue-based milestone payments

Double-digit royalties on net sales



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





Key initiative mobilized to increase ACCRUFeR® awareness

Physician Focused Efforts





Physician Sampling Program:

- Initiated in June 2025
- ~100 sample orders in June



EHR Banner Marketing:

- Initiated in Jan 2025
- >9M impressions through June 2025







Digital Advertising:

- Ramp-up in March
- 14M impressions through June 2025

Patient Focused Efforts





Brittnev Fusilier



Sia Clyde

408K Followers 2.4M Followers 521K Followers



Ashlev Lemieux

Influencer Program:

- Initiated end of May 2025
- ~645K impression through June 2025











5 New Influencers





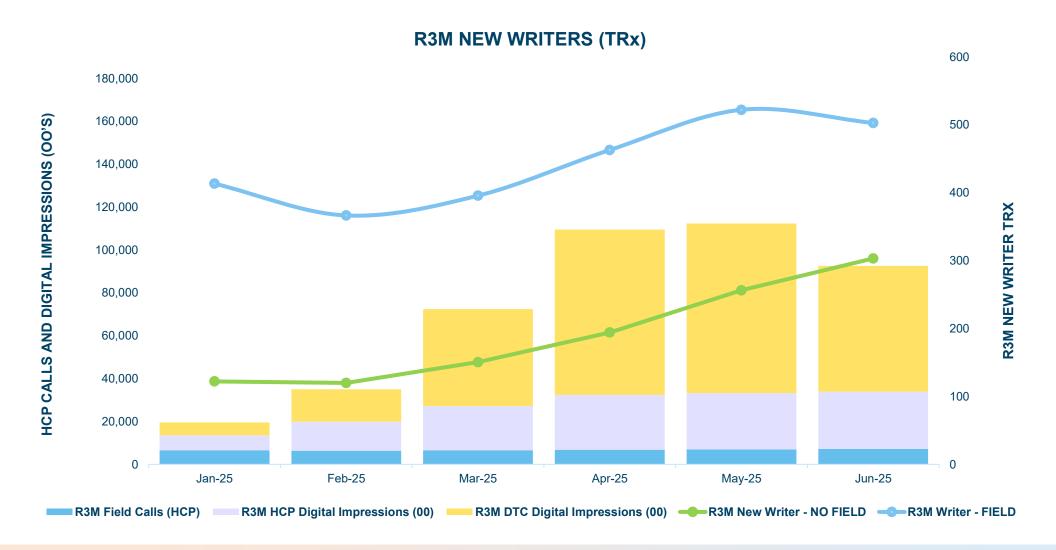


Social Media:

- Ramp-up in March
- 31M impressions though June 2025



Early indicators show positive impact of digital strategy on new writers





Key ACCRUFeR® initiatives for building momentum in Q2





Increase awareness of ACCRUFeR®

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



Enhance Sales Force Output

- Fully Staff all Territories and leverage Vacant Territory Sampling until rep start date
- Drive Demand in Top 6 States



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

- Contract with non-Blink pharmacies that have robust PA approval process for offices
- Focus on better pull-through of PA's from sales reps
- Leverage Case Managers to contact patients to pick up unfilled prescriptions



2025 Business Priorities



~\$12.8M ACCRUFeR® Net Revenues in Q2 25

Double Q1 25 revenues of \$6.4M

~47K TRx with ~23% consignment1

Highest quarter ever; 30% growth over Q1 25

~\$231 Net Selling Price in Q2 25

~25% growth over Q1 25 of \$187

Turn Cash Flow Positive by End of 2025

Q2 25

\$10.8M Q2 ending cash and cash equivalents

\$10.5M Q1 cash and cash equivalents

On track to turn cash flow positive by end of 2025

execute regulatory
process in Korea,
China, and the
Pediatric Population

Q2 25

Filed pediatric label expansion with EMA & FDA

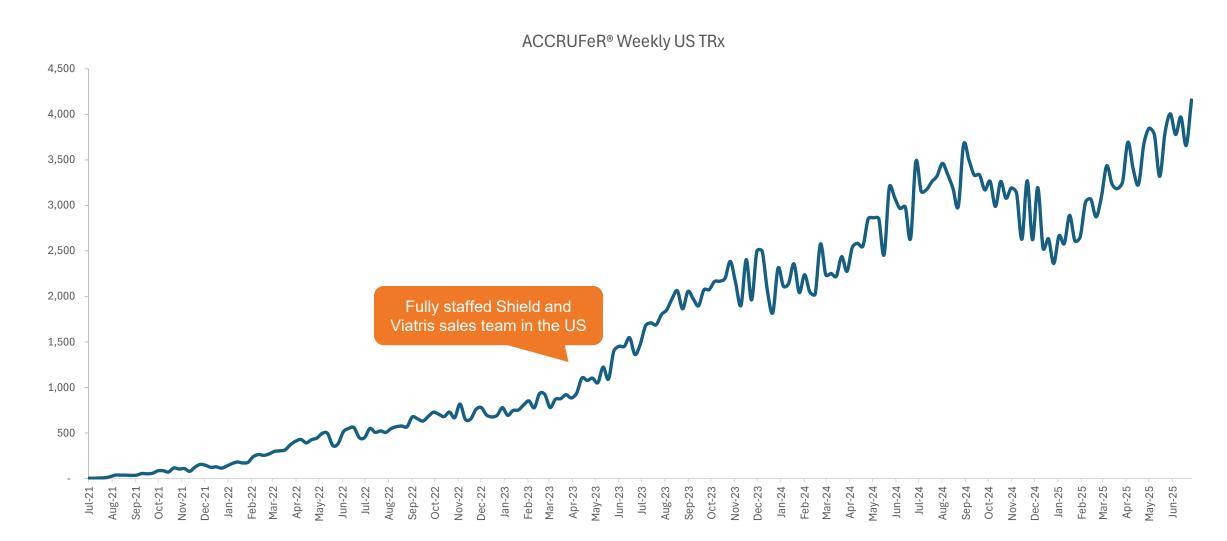
€0.5M in milestone payment from Norgine

Exclusive licence agreement in Japan with VITAL-NET, Inc.

Initial payment of ~\$665K plus regulatory and sales milestones, and double-digit royalties on net sales



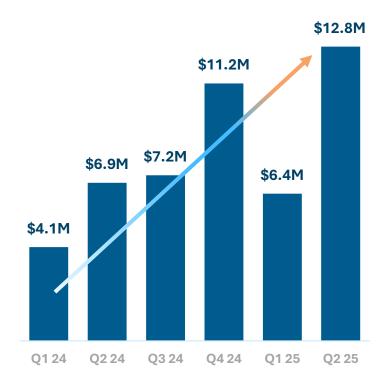
ACCRUFeR® prescriptions demonstrating significant uptake since launch



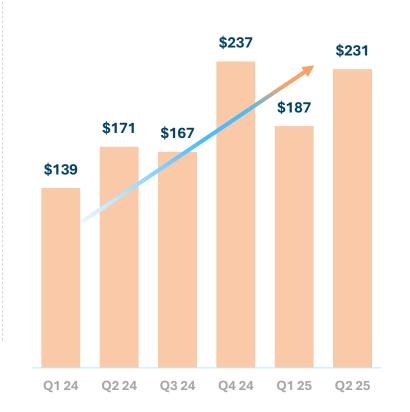


Strong growth in ACCRUFeR® in the US in Q2 2025

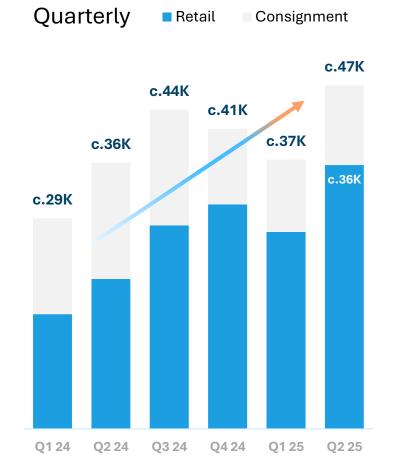
ACCRUFeR® Net Revenues Quarterly



Net Price Per ScriptQuarterly

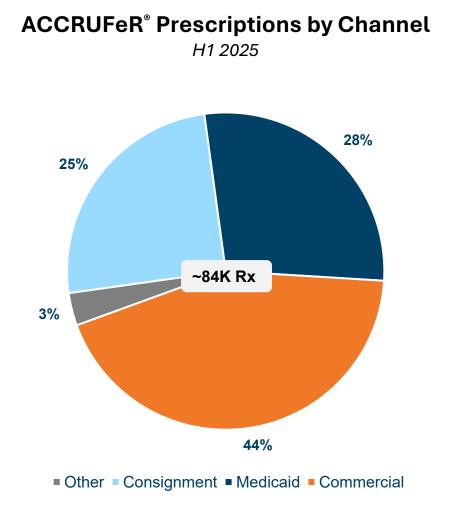


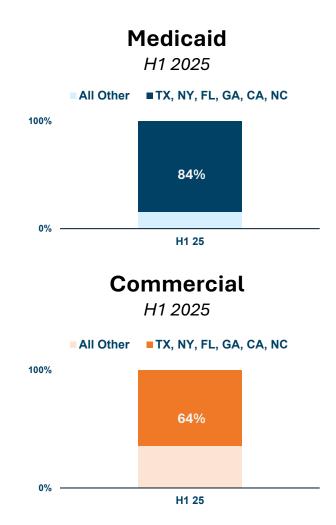
Total Prescriptions





TX, NY, FL, GA, CA, and NC are the key drivers of ACCRUFeR® TRx growth







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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



Dani - Patient using ACCRUFeR®

"When I had my first baby, my provider prescribed me a common, iron supplement. After a few weeks, my numbers weren't getting better, and the side effects were miserable"

"By the time I was pregnant with my third child, I knew I needed something different. I started taking ACCRUFeR right after my third baby was born."

"After five weeks, my hemoglobin was finally in the 12's—that was so exciting for me! I wish my doctor had known about ACCRUFeR sooner, especially when infusions and supplements didn't work for me."





Thank You!

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

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