



Res **Bio** ita

**A breakthrough in dermatology using
continuous topical protein therapy**

Corporate Overview
Q4 2024

Executive Summary

Platform

- ResVita Bio is a synthetic biology startup focused on high unmet need skin diseases
 - Platform produces continuous topical protein therapy using genetically engineered bacteria
 - Continuous protein therapy enables the function of biologics with the topical convenience of small molecules, while avoiding both modalities' systemic safety concerns
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Programs

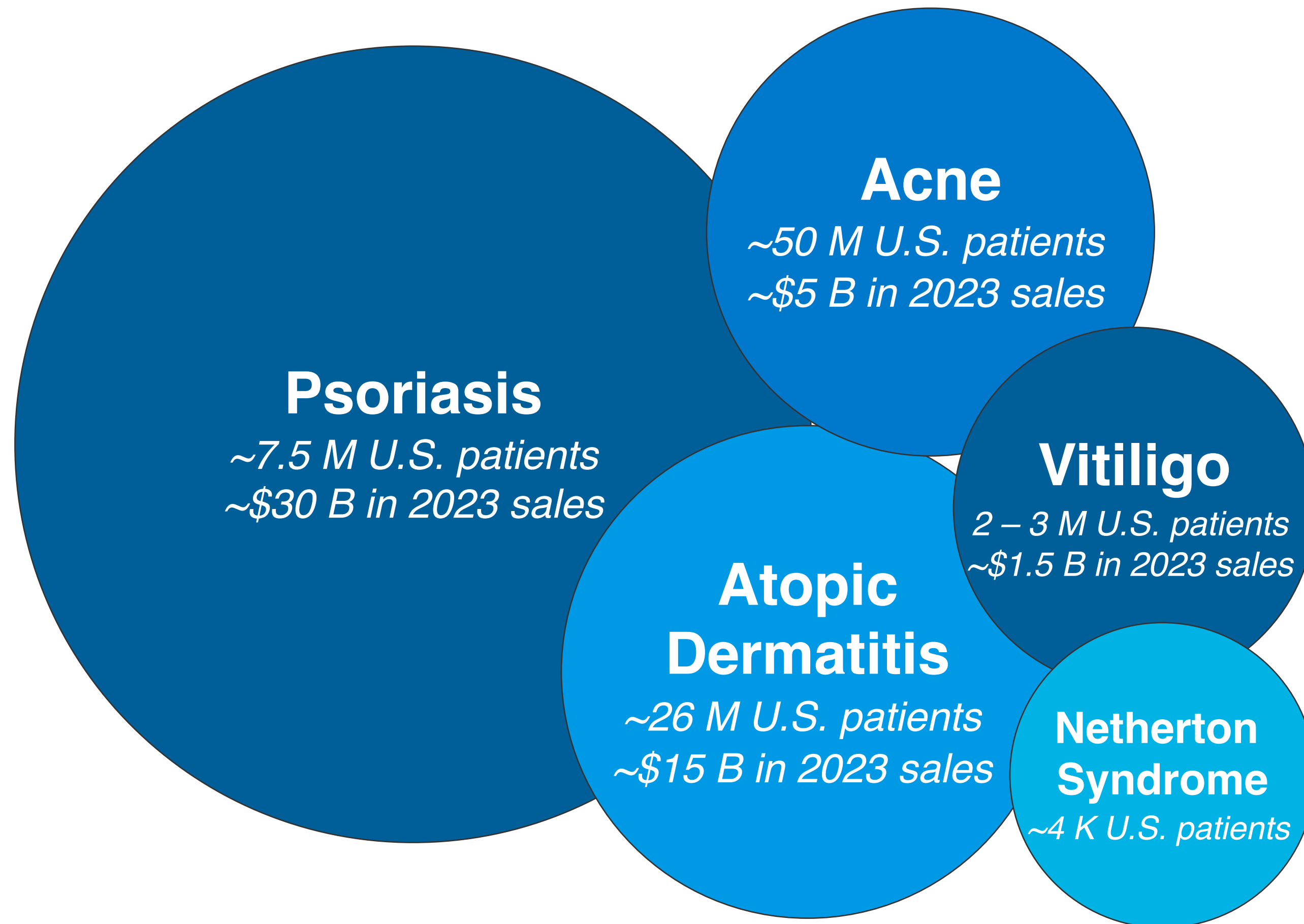
- Lead program starting IND-enabling studies in Q1 2025 to treat Netherton Syndrome, a serious rare pediatric disease affecting ~13 K patients in the U.S. and EU
 - Efficacy demonstrated in *ex vivo* human eczematous skin and *in vivo* Netherton mice
 - Rare pediatric voucher granted; FDA Interact aligned on biocontainment, CMC, preclinical
 - Additional programs in atopic dermatitis, acne, and others in discovery
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Corporate Context

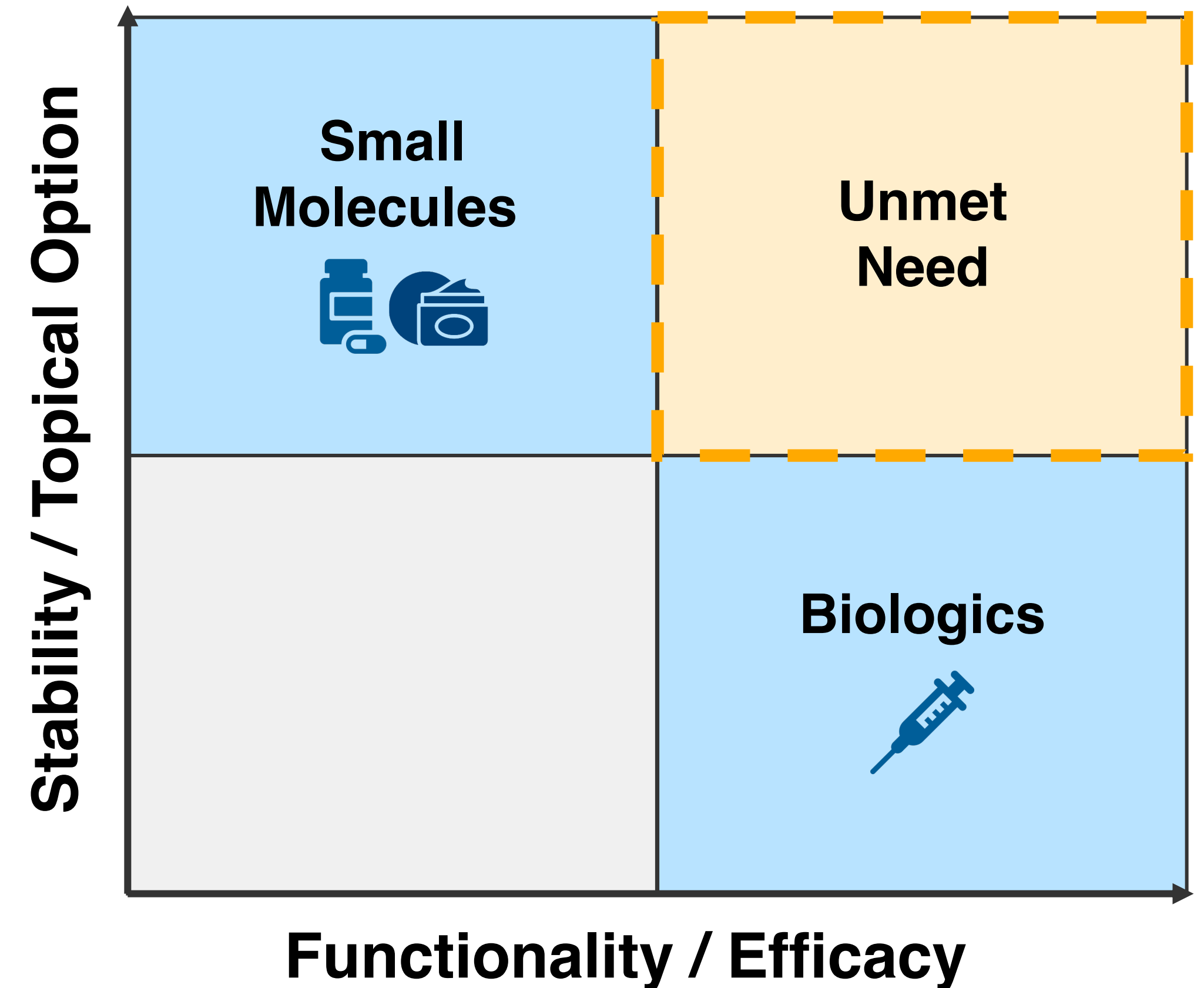
- Developed platform and nominated lead development candidate
- Seeking to deliver Phase I/II results for Netherton Syndrome, IND for Atopic Dermatitis, and platform buildout to partner on other diseases
- Advisory board consists of leaders in drug development, inflammatory dermatology diseases, and Netherton Syndrome

Large derm market in need of efficacious, safe topicals

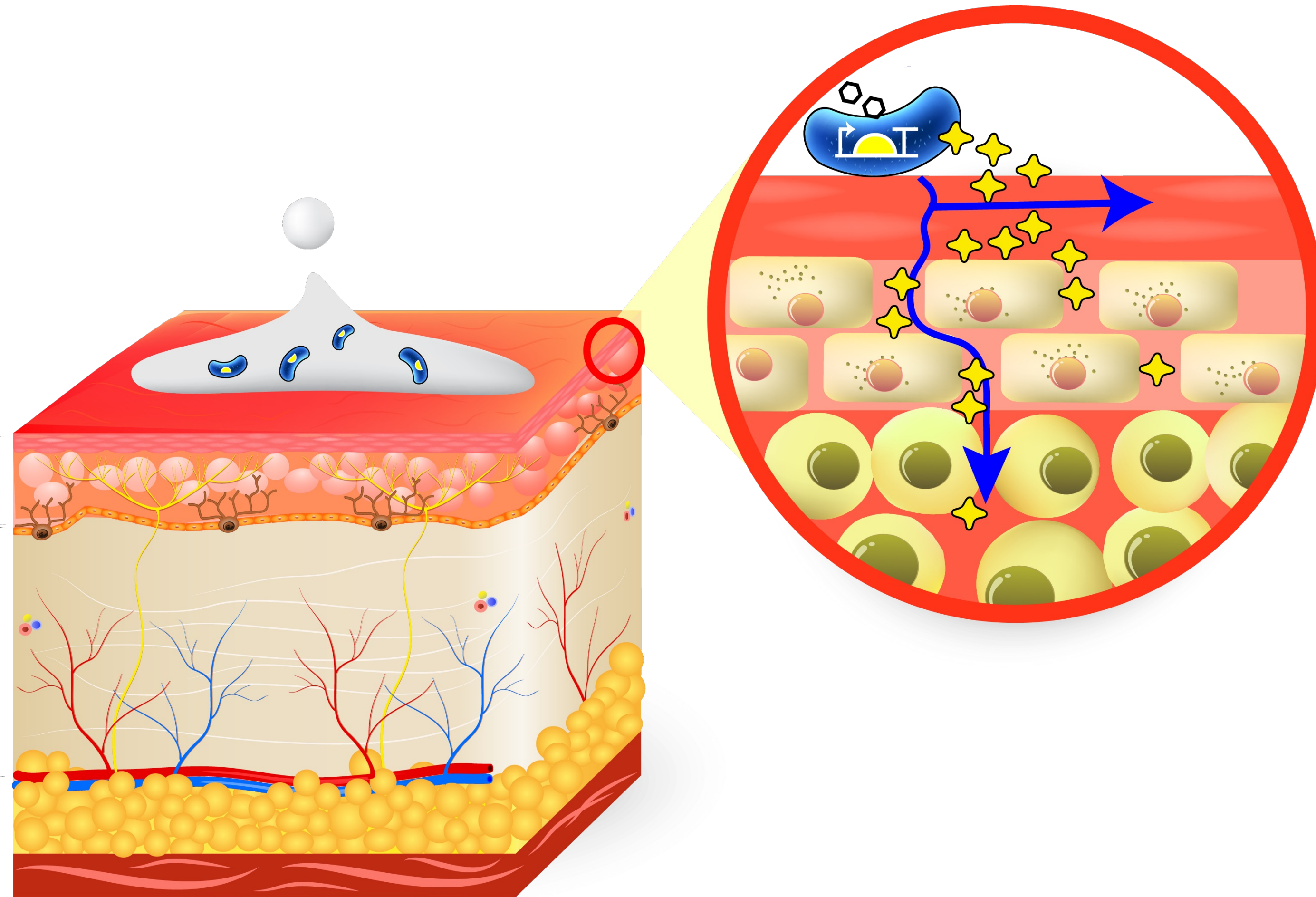
Select Dermatology Diseases



Limitation of Current Modalities







The ideal topical: continuous protein therapy via RVB Cells



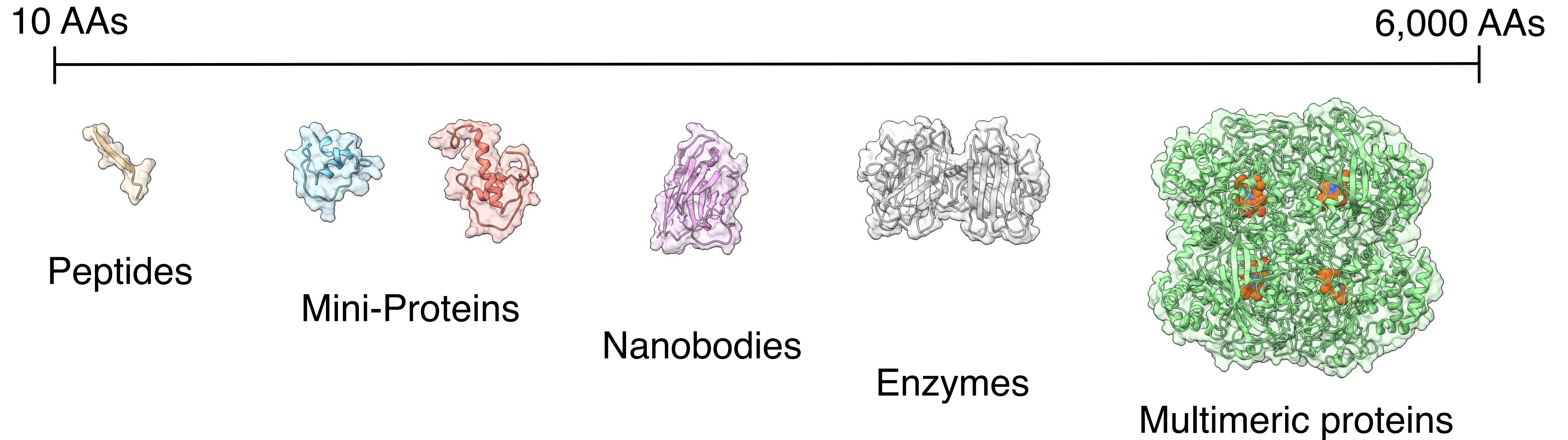
- Our RVB Cell platform enables **continuous topical protein therapy**
- RVB Cells convert nutrients from moisturizer to therapies, delivering **continuous** therapy on skin for **24 hours**
- Continuous protein therapy **increases efficacy & safety while reducing costs**

RVB Cells offer a differentiated function/convenience profile

	 Biologics	 Small Molecules	 Gene Therapy	 Continuous Protein Tx.
Efficacy (Functionality / Specificity)	High	Low-to-moderate	High	High
Safety	Moderate (Global immunosuppression)	Low-to-moderate (JAK black box)	Low (Immunogenicity)	High
Topical Optionality	No	Yes	Yes	Yes
Dosing Regimen	Bi-weekly	Twice daily	Weekly	Daily
Costs	High	Low	Exorbitant	Moderate

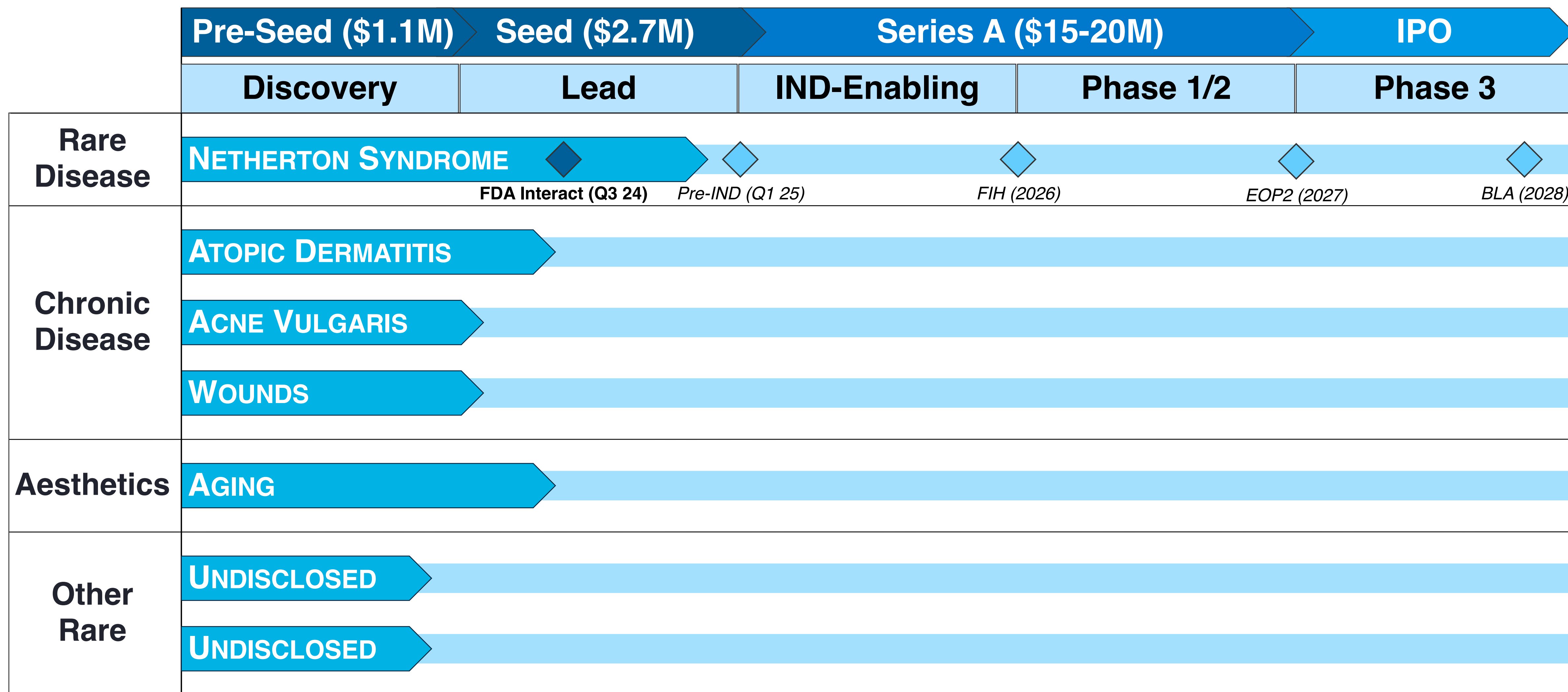
RVB cells deliver **specificity & functionality, high skin concentration levels & minimal off-site toxicity**

Broad applicability to deliver peptide/protein therapeutics



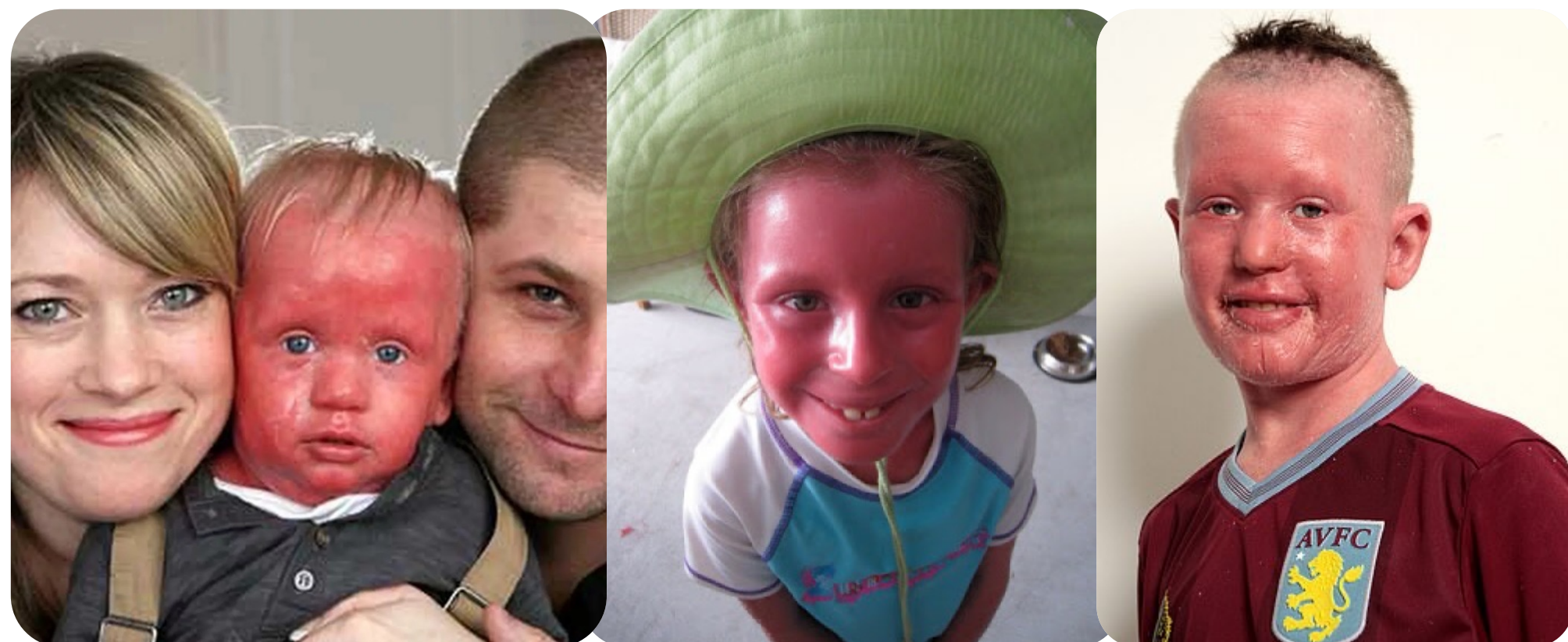
RVB Cells can be engineered to deliver a breadth of signaling, enzymatic, and cellular targets, including cytokines, proteases, and immune cells

Developing a pipeline in high unmet need skin diseases



RVB-003: disease modifying therapy for Netherton Syndrome

Netherton Syndrome



- Caused by a loss-of-function mutation in the *SPINK5* gene, encoding the **LEKTI protease inhibitor**
- LEKTI deficiency results in excess protease activity, leading to **skin barrier disruption** and **inflammatory immune response**

Epidemiology

- **Rare pediatric and orphan** condition
- Estimated prevalence of **~3.5 K** patients in the **U.S.**, **~11 K** in the **EU**

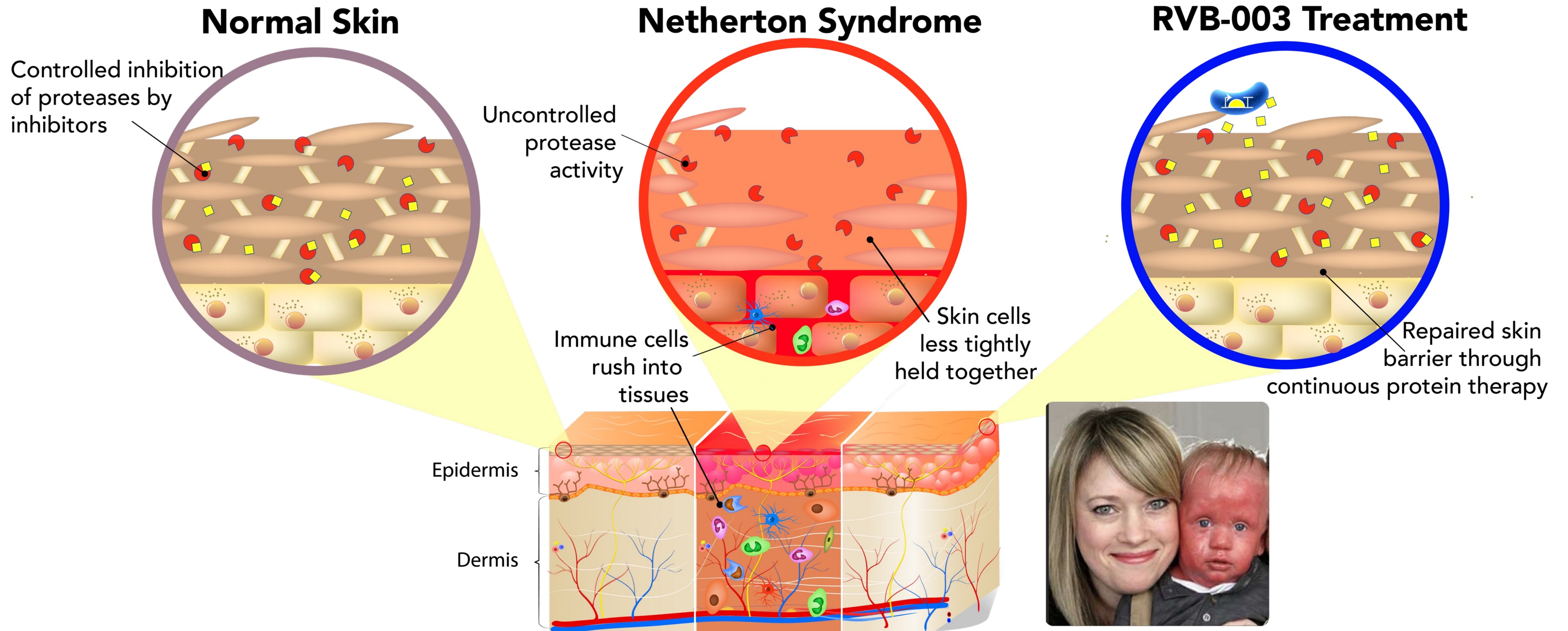
Unmet Need

- Currently **no approved therapies** with treatment limited to moisturizers
- High **burden** of disease
- **Fatal in ~20% of children**

RVB-003 Overview

- RVB-003 targets the **primary driver of disease** by inhibiting the protease KLK5
- **Rare pediatric voucher** granted, orphan designation filed, FDA Interact complete

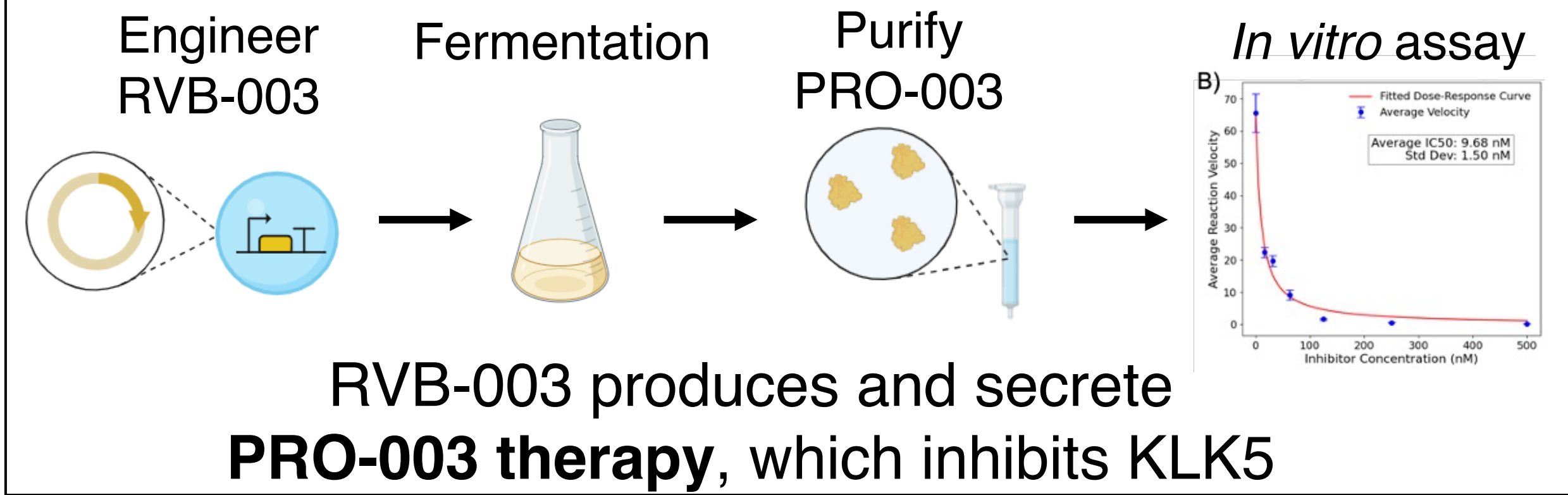
RVB-003 inhibits the KLK5 protease to restore the skin barrier



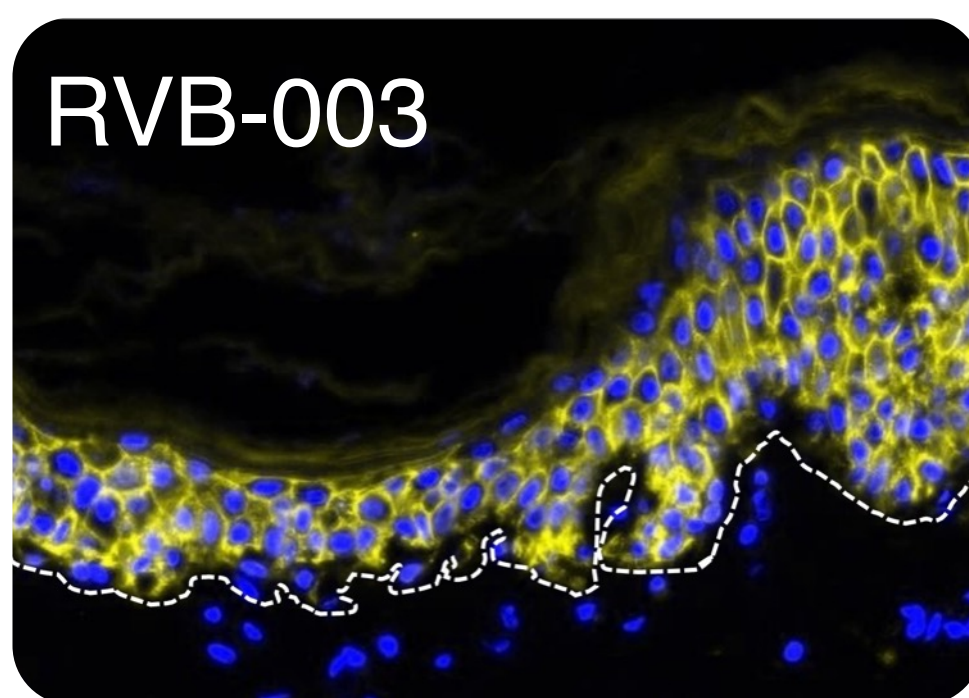
RVB Cells are genetically engineered to produce protease inhibitors to treat the cause of Netherton Syndrome

RVB-003 development status: *ex vivo* and *in vivo* efficacy

In Vitro Proof of Concept



Ex Vivo Proof of Concept

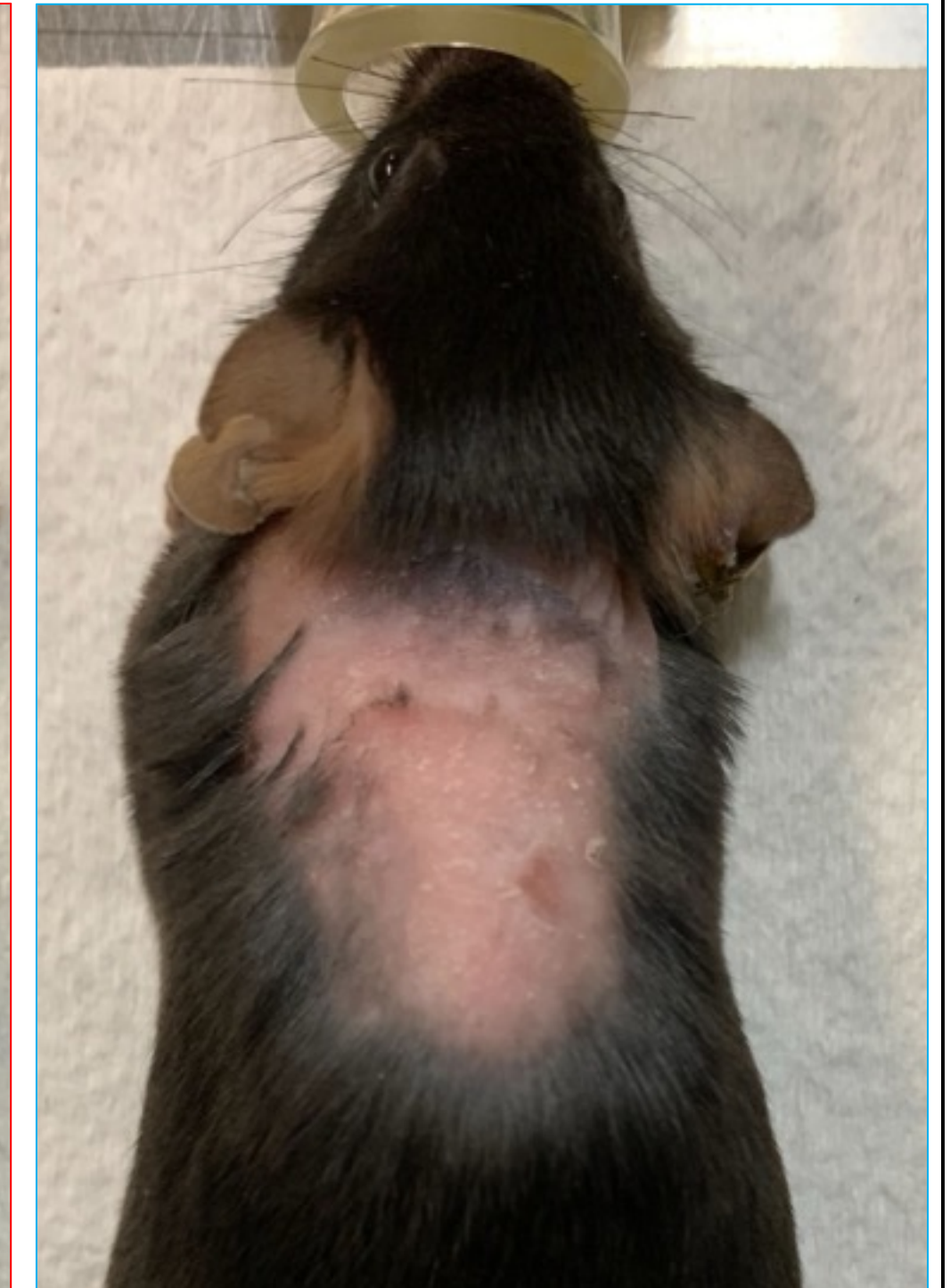


RVB-003 produces PRO-003 therapy (yellow) which **permeates** throughout human atopic epidermis and **reduces inflammation**

In Vivo Efficacy



Standard of Care



RVB-003

Skin barrier **rebuilt** within 4 days

RVB-101 in AD: building upon our NS program

Atopic Dermatitis (AD)



- Heterogeneous disease characterized by immune dysfunction and skin barrier damage.
- Hyperactive protease activity disrupts skin barrier function, contributing to AD and itch.
- Emerging therapies target protease activity to restore barrier integrity and reduce symptoms.

Epidemiology

- AD is a highly prevalent skin condition impacting over **7% of adults** and **15% children** in the U.S.
- U.S. prevalence estimated to be **>26 M**

Unmet Need

- **~60% of patients** treated with Dupixent **fail** to achieve a near-complete response
- Topical JAK inhibitors pose **black box safety concerns**

RVB-101 Overview

- Building upon validation in NS, our AD program **disease-underlying targets**
- Targets available **under CDA**

ResVita Bio Leadership

Founders



DR. AMIN ZARGAR

CEO & Co-Founder

- UC Berkeley Scientist
- NIH Awardee
- Bakar Innovation Fellow



DR. JAY KEASLING

Co-Founder

- Discover Magazine “Scientist of the Year”
- Nat. Academy of Eng.



Advisory Board

Drug Development



Steve Lo

CEO, Vaxart



Greg Went

CEO, Dextera

Inflammatory Skin



Lisa Beck

Derm, Rochester



Eric Simpson

Derm, OHSU

Rare Disease



Keith Choate

Derm, Yale



Vinzenz Oji

Derm, U of Munster



Kira Süßmuth

Derm, U of Berlin