

Dermavant is committed to fostering unprecedented change and unparalleled impact in immuno-dermatology.

We're doing this by thinking differently, pushing the boundaries of science and partnering with providers in new ways—all with one goal: transforming the lives of millions of patients with skin diseases.

Dermavant at-a-glance

- Clinical-stage biopharmaceutical company
- Founded September 2015
- A subsidiary of Roivant Sciences
- U.S. operations in Long Beach, California and Raleigh-Durham, North Carolina

About Dermavant

A Passion for Science and a Commitment to Dermatology

Dermavant Sciences, a subsidiary of Roivant Sciences, is a clinical-stage bio-pharmaceutical company focused on developing products that represent new treatment categories and address some of the greatest challenges providers and patients encounter.

The company's robust medical dermatology pipeline includes both late-stage and early-development product candidates that target specific unmet needs in two of the largest growing immuno-dermatology markets, psoriasis and atopic dermatitis, as well as other large markets, including vitiligo, primary focal hyperhidrosis, and acne.

Dermavant is developing its lead product candidate, tapinarof (DMVT-505), as a novel therapeutic aryl hydrocarbon receptor modulating agent (TAMA) topical cream for the treatment of plaque psoriasis and atopic dermatitis, which affect approximately 8 million and 26 million people in the United States, respectively.

Management team

Todd Zavodnick

Chief Executive Officer

Philip M. Brown, MD, JD

Chief Medical Officer

Chris Chapman

Chief Commercial Officer

David Rubenstein, MD, PhD

Chief Scientific Officer

Michael Swartzburg

Chief Financial Officer

Christopher Van Tuyl, JD

General Counsel

Dermavant has four product candidates addressing:

Psoriasis · Atopic Dermatitis · Vitiligo · Hyperhidrosis · Acne

PRODUCT CANDIDATE	INDICATION	STAGE OF DEVELOPMENT				
		Preclinical	Phase 1	Phase 2	Phase 3	FDA Review
Tapinarof (DMVT-505) A topical, therapeutic AhR modulating agent, inhibiting several proinflammatory factors	Psoriasis	[Progress bar: Preclinical, Phase 1, Phase 2, Phase 3]				[Progress bar: FDA Review]
	Atopic Dermatitis	[Progress bar: Preclinical, Phase 1, Phase 2]			[Progress bar: Phase 3, FDA Review]	
Cerdulatinib (DMVT-502) A topical dual JAK/Syk inhibitor	Vitiligo	[Progress bar: Preclinical, Phase 1]		[Progress bar: Phase 2, Phase 3, FDA Review]		
	Atopic Dermatitis	[Progress bar: Preclinical, Phase 1]	[Progress bar: Phase 2, Phase 3, FDA Review]			
Oxybutynin/Pilocarpine (DMVT-504) An oral combination of an immediate-release muscarinic antagonist with a delayed-release muscarinic agonist	Primary Focal Hyperhidrosis	[Progress bar: Preclinical, Phase 1]	[Progress bar: Phase 2, Phase 3, FDA Review]			
DMVT-503 A novel mechanism of action for the topical treatment of acne vulgaris	Acne Vulgaris	[Progress bar: Preclinical]	[Progress bar: Phase 1, Phase 2, Phase 3, FDA Review]			

Excludes certain Asian territories

Key market data

Psoriasis: 8M patients in U.S.¹

- \$12.0B sales²
- \$19.3B by 2026²
- Topical treatments are ~58% of U.S. RxS³

Atopic dermatitis: 26M patients in U.S.⁴

- \$1.8B sales²
- \$5.7B by 2026²
- Topical treatments are ~97% of U.S. RxS³

Vitiligo: 0.5% to 2% of global population⁵

- No FDA-approved therapies

Hyperhidrosis: 15.3M patients in U.S.⁶

- No FDA-approved systemic treatments

References: 1. National Psoriasis Foundation. 2. Evaluate Pharma. 3. IQVIA National Prescription Audit (NPA), October 2020. 4. National Eczema Association. 5. Bergqvist C, Ezzedine K. Vitiligo: A Review. *Dermatology*. 2020;236:571-592. doi: 10.1159/000506103. 6. Doolittle, J. et al. *Arch Dermatol Res*. 2016;308:743-749; US Census.

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