Restoring Health, Transforming Lives Through Innovation



Variant is an emerging specialty pharmaceutical company leveraging advanced proprietary technologies to develop best-in-class drugs for patients with rare diseases.

Our lead drug candidate is hydroxypropyl beta cyclodextrin (HPBCD), which entraps and removes lipids that can cause injury to the kidneys and other organs. Our initial focus is on chronic treatment of two orphan kidney indications, Focal Segmental Glomerulosclerosis (FSGS) and Alport Syndrome (AS).

Corporate Highlights

- Established in 2014
- Focus: \$100b+ Orphan Drug Market
- Licensed HPβCD from L&F Research and University of Miami for Treatment of Kidney Diseases
- Leading Indications:
 - FSGS: Prevention of Kidney Disease Progression
 - FSGS: Prevention of Disease Recurrence in Patients with Kidney Transplants
 - AS: Prevention of Kidney Disease Progression
- Lead Indications: IND-ready in 9-12 Months
- Later Indication: Prevention of Diabetic Kidney
 Disease

Orphan Pipeline, With Potential For Additional Indications

Variant Pipeline						
Description	Pre-Clinical	IND Ready	Phase 1	Phase 2	Phase 3	NDA Filed
VAR 200: HPβCD for FSGS		\Rightarrow				
VAR 300: HPβCD for AS		\Rightarrow				
VAR 400: HPβCD for DKD						

Management Team

Stephen C. Glover
Co-founder & CEO

Nicholas Labella, Jr. MS, RPh Acting Chief Scientific Officer

Pablo A. Guzman, MD, FACC Acting Chief Medical Officer

Peter WolfeActing VP, Finance

Karen A. Cashmere
Acting VP, Marketing

Melda Uzbil

Acting VP, Corporate Development

Brian Piper, PMP

Acting VP, Project Management

Board of Directors

Shawn M. Titcomb - Chairman

Co-Founder and Managing Partner,
Allele Capital Partners

Stephen C. Glover

Variant Co-founder & CEO

Aaron Greenblatt, Pharm D

CEO, G&W Laboratories

Jules A. Müsing

Pharmaceutical Executive & Advisor

Nico P. Pronk

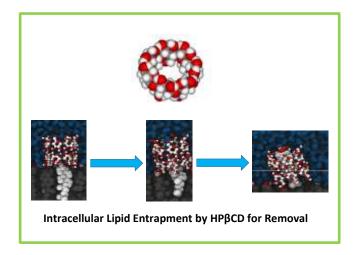
President & CEO
Noble Capital Markets

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Hydroxypropyl Beta Cyclodextrin (HPβCD)

HPβCD is comprised of seven sugar molecules bound together in a ring (cyclic oligosaccharide). The ring-shaped, three-dimensional structure has a hydrophobic cavity in its center, which is capable of entrapping and removing intracellular cholesterol and lipids demonstrated to cause kidney damage and impaired function in conditions such as FSGS and Alport Syndrome.



HPβCD: Potential to Fulfill Significant Unmet Needs in Orphan Kidney Diseases

Focal Segmental Glomerulosclerosis (FSGS) Market

- Affects up to 133,000 people in U.S.
- 5,400 new cases annually
- No disease-specific treatments
- 35% of patients in renal failure in 10 years
- 1,000 kidney transplants for FSGS/Year
- Recurs in 30% 40% of transplant patients

Alport Syndrome (AS) Market

- Affects up to 60,000 people in U.S.
- No disease-specific treatments
- 90% males, 12% females in renal failure by age
 40; 30% females by age 60
- Most deaf by age 40

HPβCD Opportunity

- Provide a first-in-class, FSGS disease modifying treatment
- Delay FSGS disease progression
- Prevent end-stage renal disease
- Prevent FSGS recurrence posttransplant
- Improve quality of life

HPβCD Opportunity

- Provide a first-in-class, AS disease modifying treatment
- Delay AS disease progression
- Prevent end-stage renal disease
- Prevent hearing loss
- Improve quality of life