



NYU



Small Molecule Inhibitors of SPNS2 for the Treatment of Chronic Inflammatory Diseases

Technology ID

SCH14-01

Category

Life Sciences/Biochemicals &

Small Molecules

Autoimmune Disease

Life

Sciences/Therapeutics/Inflammat

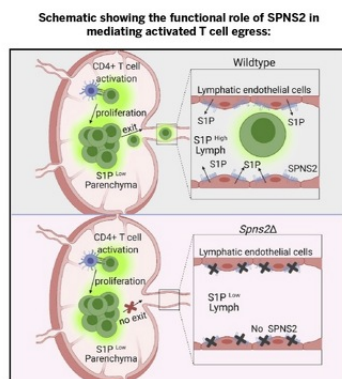
Disease

Doug Brawley

Authors

Susan Schwab, PhD

Learn more



Technology

The [Schwab Lab](#) at NYU Langone Health has elucidated a new therapeutic strategy for the treatment of chronic inflammatory diseases by trapping activated lymphocytes and achieving immunosuppression through the manipulation of sphingosine 1-phosphate (S1P) gradients rather than conventional S1P receptor 1 (S1PR1) targeting. In previous work (*Okuniewska et al. Cell Reports 2021*), they discovered that the transporter protein Spinster homolog 2 (SPNS2) is responsible for exporting S1P into the lymph guiding T cell egress from the lymph nodes (LN) during an immune response (see schematic). As discussed in the paper, this result immediately suggested that pharmacological inhibition of SPNS2 could downregulate S1P gradients resulting in the sequestration of activated T cells in the LN and prevention of subsequent inflammatory reactions and tissue damage. In target validation studies, they found that genetic deletion of *Spns2* is protective in a mouse model of multiple sclerosis. These proof-of-concept data strongly suggest that SPNS2 is a novel target for treating chronic inflammatory diseases and offers an innovative therapeutic avenue for achieving immunosuppression without common risks for adverse off-target effects.

As of present, the NYU inventors have identified and preliminarily characterized several small molecule inhibitors of SPNS2 as early therapeutic leads for the treatment of chronic inflammatory diseases. These compounds were initially identified from a 250,000-compound library and later selected for favorable activity in dose-dependent SPNS2 inhibition assays. The current lead candidate shows promising efficacy in mouse T cell sequestration experiments: lowers S1P levels (by ~25%) and reduces T cell counts (by ~60%) when administered daily (QD dosing).

Background

Chronic inflammatory diseases, which represent the most significant cause of death worldwide, are a result of aberrant immune system function and include a vast array of disorders/conditions characterized by long-term inflammation, such as heart disease, diabetes, inflammatory bowel disease (IBD), and multiple sclerosis (MS). New therapeutic strategies to treat chronic inflammatory diseases through suppression of lymphocyte-mediated inflammation and tissue damage are of critical importance to public health. In the immune system, the signaling lipid sphingosine 1-phosphate (S1P) directs the migration of lymphocytes from the lymph nodes (LN) into the lymph by binding and activating S1P receptor 1 (S1PR1) on the surface of lymphocytes. Several FDA-approved drugs target S1PR1 as a mechanism to sequester lymphocytes in the LN and cause immunosuppression; however, this therapeutic strategy suffers from adverse off-target effects due to the pleiotropic functions of S1PR1, specifically in the cardiovascular system.

Current Development Stage

Hit-to-lead optimization: Small molecule inhibitors of SPNS2 have been confirmed in primary and counter screens and profiled for dose-dependent activity. Several lead candidates have been selected for further optimization and characterization.

Applications

- Treatment of chronic inflammatory diseases:
 - Crohn's disease
 - Ulcerative colitis (UC)
 - Inflammatory bowel disease (IBD)
 - Autoimmune encephalitis
 - Relapsing remitting multiple sclerosis (RRMS)
 - Idiopathic Thrombocytopenic Purpura (ITP)
 - Fibrosis
 - Other autoimmune diseases

Advantages

- **Differentiated therapeutic strategy for modulating S1P signaling:** SPNS2 inhibition downregulates extracellular S1P gradients without directly manipulating the S1P receptor (S1PR1), the conventional target for existing FDA-approved therapeutics.
- **Low apparent risk for adverse off-target effects common with existing S1P modulating agents:** Unlike S1PR1, SPNS2 does not play critical functional roles in the cardiovascular system.
- **Spatially-specific mechanism of action:** SPNS2 inhibition targets S1P signaling gradients in the lymph and causes spatially-specific lymphocyte trapping within the lymph nodes.
- **Extensively validated target:** SPNS2 is well-characterized S1P transporter that exports S1P into the lymph guiding T cell egress from the lymph nodes during an immune response.
- **Target topology is favorable for pharmacological intervention:** SPNS2 is a membrane-embedded transporter with extracellular exposed surface (substrate binding pocket and interhelical loops) accessible to extracellular drug targeting.

Intellectual property

A PCT patent application has been filed covering the chemical compositions of the identified SPNS2 inhibitors and their methods of use.

Explore other available products at [NYU TOV Licensing](#)

References

1. Okuniewska M, Fang V, Baeyens A, Raghavan V, Lee JY, Littman DR, Schwab SR ,
<https://pubmed.ncbi.nlm.nih.gov/34260944/>
2. Mendoza A, Bréart B, Ramos-Perez WD, Pitt LA, Gobert M, Sunkara M, Lafaille JJ, Morris AJ,
Schwab SR. , <https://pubmed.ncbi.nlm.nih.gov/23103166/>