

Verkazia™ (cyclosporine 0.1%) for the Treatment of Severe Vernal Keratoconjunctivitis is Now Reimbursed by Non-Insured Health Benefits (NIHB) Program

August 17, 2021, TORONTO – Santen Canada Inc., a subsidiary of Santen Pharmaceutical Co., Ltd. (hereinafter, Santen), a global company focused exclusively on ophthalmology, today announced that Verkazia (cyclosporine topical ophthalmic emulsion 0.1% w/v) eye drops for the treatment of severe vernal keratoconjunctivitis (VKC) is now publicly reimbursed through the Non-Insured Health Benefits (NIHB) Program Limited Use Benefit for eligible patients who meet specific clinical criteria. The criteria and listing may be found [here](#).

Verkazia was approved for the treatment of severe vernal keratoconjunctivitis in children from 4 years of age through adolescence by Health Canada in 2018.¹

"Santen is pleased to see Verkazia reimbursed for patients living with severe VKC in Canada, where patients meet the specific criteria under the NIHB Program Limited Use Benefit," said Carol Stiff, Head of Santen Canada Inc. "As a company dedicated to Ophthalmology, we aim to reduce the loss of social and economic opportunities for people around the world due to eye conditions, and we hope this decision will help provide more access to the underserved population of Canadians." Santen is continuing to have discussions with other provinces, territories, and federal agencies regarding the listing of Verkazia under publicly funded drug programs.

About Vernal Keratoconjunctivitis (VKC)

VKC is a rare and recurrent allergic eye condition, most common in children and adolescents, that causes severe inflammation of the surface of the eye. The main symptoms of VKC – light sensitivity, tearing, itching, and mucous discharge – can prevent those affected from participating in everyday activities.^{2,3,4} Approximately one-third of VKC cases are considered severe, and without adequate treatment may result in corneal ulcers and even vision loss.⁵

About Verkazia

Verkazia cyclosporine 0.1% topical ophthalmic emulsion is indicated for treatment of severe vernal keratoconjunctivitis in children from 4 years of age through adolescence.¹ The recommended treatment regimen approved by Health Canada is one drop of Verkazia 4 times a day (morning, noon, afternoon, and evening) to be applied to each affected eye. The treatment should be discontinued if no improvement in signs and symptoms of vernal keratoconjunctivitis is observed. The treatment can be maintained at the recommended dose or decreased to one drop twice daily once adequate

control of signs and symptoms is achieved. Treatment should be discontinued after signs and symptoms are resolved and reinitiated upon their recurrence.

About Santen

As a global specialized company dedicated to ophthalmology, Santen brings a 130-year history of scientific knowledge and organizational capabilities to research, development, and commercialization of pharmaceuticals, surgical and medical devices, and OTC eye-care products. Santen is the market leader for prescription ophthalmic pharmaceuticals in Japan and its products now reach patients in more than 60 countries. Santen provides products and services to contribute to the well-being of patients, their loved ones and consequently to society. For more information, please visit Santen's websites www.santencanada.ca, www.santenusa.com and www.santen.com (Japan headquarters).

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¹ Verkazia™ (cyclosporine topical ophthalmic emulsion 0.1% w/v) Product Monograph. Santen Incorporated. December 21, 2018.

² Kumar S. Vernal keratoconjunctivitis: a major review. *Acta Ophthalmol* 2009;87:133-147

³ Leonardi A. Management of vernal keratoconjunctivitis. *Ophthalmol Ther.* 2013;2:73e88

⁴ Sacchetti M, et al. Development and testing of quality of life in children with vernal keratoconjunctivitis questionnaire. *Am J Ophthalmol* 2007;144:557-563

⁵ Bremond-Gignac D, et al. Prevalence of vernal keratoconjunctivitis: a rare disease? *Br J Ophthalmol* 2008;92:1097-1102

