

# Take the plunge

Visit the **Alhemo™ Product Theatre** to learn how Alhemo™ can support your haemophilia B patients with inhibitors<sup>1</sup>

Alhemo™ is the first subcutaneous prophylactic treatment for patients with haemophilia B with inhibitors, delivered with a first-in-class prefilled pen device.<sup>1</sup>

June 25<sup>th</sup> 2023 | 12:15–13:00 (EST)

## Presentation Theatre 4

### AGENDA

12:15	<b>Welcome and introduction</b>	Dr. Stephanie Seremetis
12:20	<b>What are the remaining unmet needs in haemophilia B with inhibitors?</b>	Dr. Manuel Carcao
12:30	<b>Bringing innovation in the treatment of haemophilia B with inhibitors</b>	Dr. Stephanie Seremetis
12:50	<b>Holistically addressing patient convenience with a new pen device</b>	Vanessa Bouskill
13:00	<b>Meeting closure</b>	All

### SPEAKERS



**Dr. Stephanie Seremetis**  
CVP and Chief Medical Officer Rare Disease and Advanced Therapies, Novo Nordisk, Bagsvaerd, Denmark



**Dr. Manuel Carcao**  
Hospital for Sick Children, Toronto, Ontario, Canada



**Vanessa Bouskill**  
Hospital for Sick Children, Toronto, Ontario, Canada

The image shown is a model and not a real patient.



The information presented is based on the Canadian label. Alhemo™ is indicated for routine prophylaxis to prevent or reduce the frequency of bleeding in patients  $\geq 12$  years of age with haemophilia B (congenital factor IX deficiency) with factor IX inhibitors.<sup>1</sup>



# Everyday protection for your haemophilia B patients with inhibitors (HBwI)<sup>1</sup>



## 1<sup>st</sup> subcutaneous prophylaxis for HBwI<sup>1</sup>

Mean ABR of 1.7 for patients with inhibitors in the explorer 7 trial<sup>1,2</sup>



## 1<sup>st</sup> in class pen<sup>1,2</sup>

Ready-to-use portable pen that can be stored at room temperature\*



## Established safety profile<sup>\*1,2</sup>

Well-tolerated and can be used concomitantly with both bypassing agents<sup>†1,2</sup>

\* Up to 30 °C for up to 4 weeks.<sup>2</sup>

† For both bypassing agents, the lowest approved dose and dosing interval in the approved label are recommended. For activated prothrombin complex concentrate (aPCC), a maximum dose of 100 international unit/kg body weight within 24 hours is recommended. For severe bleeds, it is recommended to follow the dosing interval provided in the approved label for each product, based on clinical judgement.<sup>1</sup>



### Indications:

Alhemo™ (concizumab injection) is indicated for the treatment of adolescent and adult patients (12 years of age or older) with haemophilia B (congenital factor IX [FIX] deficiency) who have FIX inhibitors and require routine prophylaxis to prevent or reduce the frequency of bleeding episodes.

There is limited clinical experience of Alhemo™ use in patients known to have mild or moderate haemophilia B (FIX activity > 2%).

Please visit <https://www.novonordisk.ca/content/dam/nncorp/ca/en/products/alhemo-en-product-monograph.pdf> for more information relating to adverse reactions, drug interactions, and dosing information, which have not been discussed in this advertisement. The Product Monograph is also available by calling us at 1-800-465-4334.

The information presented is based on the Canadian label. This material is intended to be used at the ISTH 2023 Congress only.