

## Meeting Minutes

<b>Institution:</b>	Minnesota Urology P.A.		
<b>Meeting Date:</b>	September 10, 2025		
<b>Meeting Time</b>	10:30 AM Central Time		
<b>Meeting Type:</b>	Virtual Platform Teleconference (Remote) Open to the Public		
<b>Members in Attendance:</b>	<b>Member</b>	<b>Voting</b>	<b>Member Type</b>
	Bavaret, Tammy	Yes	Chair: Biosafety Expert/HGT Expert
	Rastein, Daniel	Yes	Core Member: Biosafety Expert/HGT Expert
	Campbell, Mark	Yes	Core Member: Biosafety Expert/HGT Expert
	Statz, Catherine	Yes	Local Unaffiliated Member
	Park, Gregory	Yes	Local Unaffiliated Member
	McKenzie, Carrie	No	Site Contact
<b>Invited Members Not in Attendance:</b>	<b>None</b>		
<b>Guests:</b>	Robb, Melissa Ogbu, Emmanuel Lavato, Katie (joining at 11:10am)		
<b>Staff:</b>	Payne, Kaylie		

**Call to Order:** The IBC Chair called the meeting to order at 10:32 AM. A quorum was present as defined in the Sabai IBC Charter.

**Conflicts of Interest:** The IBC Chair reminded all members present to identify any conflicts of interest (COI). No COI was declared by any voting member of the IBC for any of the items on the agenda.

**Public Comments:** No public comments were made prior to or at the meeting.

**Review of Prior Business:** None

**Previous Meeting Minutes:** Previous meeting minutes from **10/1/24** were reviewed and approved with no changes requested.

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### New Business:

<b>PI:</b>	Milbank, Aaron MD
<b>Sponsor:</b>	CG Oncology, Inc
<b>Protocol:</b>	CORE-008 A Phase 2, Multi-Arm, Multi-Cohort, Open-Label Study to Evaluate the Safety and Efficacy of Cretostimogene Grenadenorepvec in Participants with High-Risk Non-Muscle-Invasive Bladder Cancer (NMIBC)
<b>Review Type:</b>	Annual Review
<b>NIH Guidelines Section:</b>	III-C-1

**Trial Summary:** CORE-008 is a multi-arm, open-label Phase II trial sponsored by CG Oncology, Inc. and designed to assess the safety and efficacy of cretostimogene grenadenorepvec in participants with high-risk non-muscle invasive bladder cancer. The study agent cretostimogene grenadenorepvec consists of a recombinant, conditionally replicating oncolytic adenovirus. The investigational product (IP) is administered by intravesical instillation into the bladder.

**Biosafety Containment Level (BSL):** The study agent cretostimogene is based on a recombinant Risk Group 2 virus containing more than two-thirds of the native genome, requiring the use of BSL-2 containment under the NIH Guidelines.

### Risk Assessment and Discussion:

- The Committee reviewed the clinical trial Sponsor's study documents and the Sabai-generated comprehensive study-specific Risk Assessment which collectively provided a thorough description of the recombinant or synthetic nucleic acid molecules (investigational product/s) and the proposed clinical research activities involving the IP.
  - In summary, the primary risks in this clinical trial include potential occupational exposure from accidental spills, splashes, aerosols and needlesticks of the IP during preparation and/or administration procedures. These potential risks are mitigated through a combination of relevant staff training, safe clinical practices including Standard Precautions and sharps safety and use of appropriate PPE (as prescribed in the Risk Assessment and documented in the IBC submission package).
  - The Site confirmed that only study personnel who have been educated on the potential biohazards and the precautions to be taken when working with the IP will handle the IP or any materials contaminated by the IP.
  - The Site confirmed that study personnel are sufficiently trained in the practices and techniques required to safely work with the IP.
  - The Site confirmed that staff members receive Bloodborne Pathogens training.
  - Occupational Health Recommendations: The Sponsor notes that individuals who are at a potentially higher risk from working with or handling the study agent, such as pregnant or breastfeeding women and immunosuppressed or immunocompromised individuals,

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should not prepare, administer, or otherwise handle the study agent or materials contaminated with the study agent or provide direct care for treated participants presenting with any symptoms of illness attributed to cretostimogene for at least 1 week after treatment or until complete resolution of symptoms.

- The Committee had no additional significant comments or recommendations regarding the description of the potential risks and occupational exposure hazards associated with handling the IP in this clinical trial, or the proposed mitigation strategies, as detailed in the Risk Assessment.
- The Committee reviewed the Site's facility details, relevant study-specific procedures and practices, the Annual Review Report, and other applicable information provided by the Site for the purposes of the IBC review.
  - The Site verified that the information provided by the Chair was accurate.
  - In Response to a question from the Committee, the site confirmed the disinfectants described for spill cleanup are correct. The committee had no questions.
  - In response to a question from the Committee, the site confirmed that if the drainage bag is used to drain the urine, it will be wrapped in an absorbent pad then disposed of in the biowaste container.
  - In response to a question from the committee, the site confirmed that the Safety / OSHA Training Refresher does include Bloodborne Pathogens training.
  - The Committee stipulated that the Site contact their BSC Certifier to verify whether the report incorrectly identified the unit as having a direct exhaust connection instead of a canopy exhaust connection, as the indicated BSC model does not support a direct connection. If this was an error, the Site must confirm that the canopy was properly tested by the certifier and request an updated report reflecting the correct configuration and testing. The revised report must be provided to Sabai.

**Motion:** A motion of Full Approval for the study at BSL-2 was passed by unanimous vote. There were no votes against and no abstentions.

- Contingencies stated by the Committee: None
- Stipulations stated by the Committee:
  - Site to contact their BSC Certifier to verify whether the report incorrectly identified the unit as having a direct exhaust connection instead of a canopy exhaust connection, as the indicated BSC model does not support a direct connection. If this was an error, the Site must confirm that the canopy was properly tested by the certifier and request an updated report reflecting the correct configuration and testing. The revised report must be provided to Sabai by **10/9/2025**. The Committee agreed that resolution of this stipulation can be approved following review by the Chair

**Review of Incidents:** Nothing to report.

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**IBC Training:** Nothing to report.

**Reminder of IBC Approval Requirements.**

**Adjournment:** The IBC Chair adjourned the meeting at 11:11 AM

**Post-Meeting Pre-Approval Note:** None