INSTITUTIONAL BIOSAFETY COMMITTEE REVIEW

MEETING MINUTES

Meeting Date:	**Thursday, August 7, 2025**
Time:	12:00 pm Central Time
Location:	Zoom Teleconference
Institution:	The University of Kansas Medical Center, Kansas City, KS
Principal Investigator:	**Pradeep Mammen, MD**
Protocol:	Sardocor Corp., **SRD-001-1004**
NCT Number:	NCT06224660
Meeting Type:	Continuing Review of Protocol and Site
Title:	A phase 1b, open-label, controlled trial evaluating the safety and efficacy of SRD-001 (AAV1/SERCA2a) in subjects with cardiomyopathy secondary to Duchenne Muscular Dystrophy.

1. Call to order:

The Meeting was called to order at 12:21 pm Central Time.

2. Introductions and orientation:

Introductions were made and the Chair oriented members to the meeting procedures.

3. Declaration of quorum:

Six voting members were present, including two local members unaffiliated with the institution and the Institution's Biosafety Officer. Also present were seven Institutional Representatives and IBC Services staff. The Chair declared that a quorum was present.

4. Conflict of Interest:

The Chair requested that voting members report any conflict of interest regarding this meeting. No conflicts of interest were reported.

5. Public posting:

An Institutional Representative confirmed that notice of the meeting was publicly posted. No public comments were received by the site or the Committee regarding this review.

6. Approval of previous meeting minutes:

Minutes Approved - YES: 6 NO: 0 ABSTAIN: 0

7. Review of proposed research:

The Chair provided an overview of the protocol and status of the study.

The Chair provided an overview of changes since the last review.

8. Determination for biosafety level and period of IBC oversight:

The Committee previously determined that **BSL-1 containment facilities and practices plus Standard Precautions** are required for SRD-001, since it consists of an AAV vector being administered by infusion in a clinical setting. The Committee reaffirmed this determination.

The Committee previously determined that IBC oversight will continue for **3 months after the last subject's last dose of SRD-001 locally**, provided that other biosafety criteria for study closure are also met. The Committee reaffirmed this determination.

9. Vote on the Protocol:

The Committee voted for the following determination on the Protocol:

X	APPROVED
	CONDITIONALLY APPROVED
	TABLED
	DISAPPROVED

DETERMINATION VOTE - YES: 6 NO: 0 ABSTAIN: 0

10. Review of proposed facilities and practices:

The Chair provided an overview of the arrangement for the facilities and practices.

Points of Discussion:

1. The Committee recommended that the Site Inspection Checklist, Item 21 be revised to indicate that there are no handwashing sinks inside the dosing rooms.
2. The Committee discussed that, per the Site Map, prefilled disposable eyewash bottles are available in the dosing rooms and recommended that the site confirms that this is accurate.
3. The Committee recommended that the mops stored next to the plumbed eyewash station in the [redacted] be moved to another area.

11. Site requirements:

The Chair reviewed training and communication requirements for maintaining IBC approval with the Biosafety Officer and the Institutional Representatives.

12. Vote on the Site:

The Committee voted for the following determination on the Site:

X	APPROVED
	CONDITIONALLY APPROVED
	TABLED
	DISAPPROVED

DETERMINATION VOTE - YES: 6 NO: 0 ABSTAIN: 0

13. Advice to the Institution: None.

14. Meeting adjourned: The meeting was adjourned at 12:25 pm Central Time.