INSTITUTIONAL BIOSAFETY COMMITTEE REVIEW

MEETING MINUTES

Meeting Date:Friday, June 27, 2025Time:11:00 am Central TimeLocation:Zoom Teleconference

Institution: Texas Oncology - Sammons Cancer Center, Dallas, TX

Principal Investigator: Charles L. Cowey, MD

Protocol: Moderna Therapeutics, Inc., mRNA-4157-P201

NCT Number: NCT03897881

Meeting Type: Continuing Review of Protocol and Site

Title: A Phase 2 Randomized Study of Adjuvant Immunotherapy With the Personalized

Cancer Vaccine mRNA-4157 and Pembrolizumab Versus Pembrolizumab Alone

After Complete Resection of High-Risk Melanoma

1. Call to order:

The Meeting was called to order at 11:00 am Central Time.

2. Introductions and orientation:

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

3. Declaration of quorum:

Four voting members were present, including one local member unaffiliated with the institution. Also present were two 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: 4 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 mRNA-4157 because the study agent consists of synthetic mRNA administered via intramuscular injection. 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 mRNA-4157 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:

Χ	APPROVED
	CONDITIONALLY APPROVED
	TABLED
	DISAPPROVED

DETERMINATION VOTE - YES: 4 NO: 0 ABSTAIN: 0

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10. Review of proposed facilities and practices:

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

Points of Discussion:

- The Committee recommended that the Institution confirm the effective disinfectants that are available where the study agent is handled, and update the Biosafety SOP and Site Inspection Checklist with these disinfectants.
- 2. An Institutional Representative confirmed that biohazardous waste containers are closed with a lid when not in use.

11. Site requirements:

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

12. Vote on the Site:

The Committee voted for the following determination on the Site:

Х	APPROVED
	CONDITIONALLY APPROVED
	TABLED
	DISAPPROVED

DETERMINATION VOTE - YES: 4

NO: 0

ABSTAIN: 0

13. Advice to the Institution: None.

14. Meeting adjourned: The meeting was adjourned at 11:10 am Central Time.