Հայաստանի Հանրապետության Գիտությունների Ազգային Ակադեմիա

Մոլեկուլային կենսաբանության Ինստիտուտ



National Academy of Sciences of the Republic of Armenia

Institute of Molecular Biology

<u>IRB/IEC: IRB00004079</u>

(IORG 0003427)

Project name

Date

Dr. *name*, as a Principal Investigator of the "*Project name*" has submitted all the required documents for IRB review to the Institutional Ethics Committee at the Institute of Molecular Biology, NAS RA:

No.	Document	Version No. & Date
1.	IRB General Application Form	#x – date
2.	Protocol Summary Form	#x – date
3.	Participant Consent Form	#x – date
4.	Questionnaire	#x – date

The following members of the Ethics committee were present at the meeting held on *date*, Yerevan.

No.	Name	Role in Ethics committee	Qualification	Gender	Affiliation to Institute (Yes/No)
1.	Lyudmila Nersesova	Chair	PhD, Biochemist	Female	Yes
2.	Zaruhi Khachatryan	Member	PhD, Molecular Biologist	Female	Yes
3.	Zaven Karalyan	Member	PhD, DrSc, Virologist	Male	Yes
4.	Hakob Devejyan	Member	PhD, Manager	Male	Yes
5.	Hrachik Vardapetyan	Member	PhD, DrSc, Prof, Biochemist	Male	No
6.	Ruben Hovhannisyan	Memeber	PhD, DrSc, Physician	Male	No
7.	Gohar Mkrtchyan	Member	PhD, Biochemist	Female	Yes
8.	Susanna Mkrtchyan	Member	MS, Public Health Officer	Female	No
9.	Rita Malakyan	Member	PhD, Biophysicist	Female	Yes
1.	Levon Yepiskoposyan	Alternative member	PhD, DrSc, Prof, Geneticist	Male	Yes
2.	Aren Khoyetsyan	Alternative member	PhD, Molecular Biologist	Male	No
3.	Elina Arakelova	Alternative member	PhD, Molecular Biologist	Female	Yes
4.	Lilit Hovhannisyan	Alternative member	PhD, Biochemist	Female	Yes

The IRB has conducted full review of all presented materials and concluded that the proposed study entitled "*Project name*" meets the human subjects protection requirements, therefore the project is approved in the presented form.

The IRB expects to be informed about the progress of the study, any changes occurring in the course of the study, any revision in the protocol and patient information/informed consent and ask to be provided a copy of the final report

This ethics committee is working accordance to the US FDA regulations at 21 CFR parts 50 and 56, International Ethical Guidelines for Biomedical Research Involving Human Subjects, ICH-GCP, Schedule Y and ICMR guidelines and other applicable regulation.

Chair of the IRB of the IMB NAS RA

Dr. L. Nersesova

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