Al-Farabi Kazakh National University

Faculty\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Department \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Full Name of Master student

Code and Name of Specialty :

Years of study:

**Research Protocol**

**«Title of Master Dissertation in Kazakh»**

**«Title of Master Dissertation in Russian»**

**«Title of Master Dissertation in English»**

Scientific Supervisors:

Full names, degrees and titles

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4. Abbreviations

**……………………………………………………………………………………..**

1. Researchers (name, names)

……………………………………………………………. CVs in Application 1

 **3.** **Introduction**

**3.1. Background**

**3.2. Research purpose**

* 1. Research Objectives

**3.4** Scientific Novelty and Practical Significance

During the study, it is planned to …………

For the first time will be …………..

Practical significance

1. Identified ………

2. Suggested………

**4. Methods**

**4.1.** Design

**4.2.** Research Objects

**4.3.** Data collection methods

 **4.4**. Data collection location

 **4.5.** Identification of sources

 **4.6.** Development of research tools

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 **4.9.** Calendar plan of research

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 **4.11.** *Exclusion criteria*

 **5.** Data Collection and Management

**5.1. Data collection**

**5.2. Data Storage and Archiving**

 **6.** Ethical Aspects

 **6.1.** Research ethics

 **6.2.** Interaction with Local Ethical Committee

The protocol, questionnaire, form, information leaflets and informed consent form are submitted for consideration to the Local Ethical Committee of Al-Farabi KazNU.

**6.3.** Information Leaflet

**6.4. Informed Consent**

 **6.5.** Signing of Form of Informed Consent

*The researcher must provide the participant with oral and written information about this study in a form that the participant can understand. Before the start of the survey and interview, the written consent of the participant must be obtained.*

*Before consent is obtained, the participant must be allowed sufficient time to:*

*- Obtaining interesting detailed information about the study;*

*- reflection and opportunities to ask clarifying questions;*

*- making a decision to participate in the study.*

*If the participant decides to participate in the study, then a member of the research team must sign the informed consent form with the participant and indicate the appropriate date of signing.*

*The original signed consent form should be kept at the study site or by the principal investigator. Another copy must be given to the study participant.*

**6.6.** Forms of Informational leaflet and Informed Consent (Applications 2,3)

**6.7.** Confidentiality, Privacy and Security of Personal Data

Participants' personal data will be stored and processed in accordance with the Law of the Republic of Kazakhstan "On Personal Data and their Protection".

**7.** List of Applications

7.1. CV of Master Student and Scientific Supervisors

7.2. Tools

7.3. Information Leaflet

* 1. Informed Consent