INFORMATION CLAUSE FOR PERSONS PARTICIPATING IN SCIENTIFIC RESEARCH CONDUCTED BY THE FOUNDATION

1. **Data Collector is Sano – Centre for Computational Personalised Medicine – International Research Foundation, Czarnowiejska 36 / C5, 30-054 Kraków, KRS: 0000797490, NIP: 6772446472, REGON: 384298430, e: info@sanoscience.org, t: +12 307 27 37, hereinafter referred to as "Sano".**
2. **Data Protection Officer Contact with the appointed Data Protection Officer: e-mail: iod@sanoscience.org**
3. **Purpose of processing**

We process personal data of participants in scientific research for the purpose of:

- conducting scientific research in accordance with specific research objectives;
- analysis and interpretation of data as part of research projects;
- monitoring, evaluation and validation of test results;
- comply with legal obligations arising from the regulations governing the conduct of scientific research;
- archiving of test results and documentation related to their conduct.

4. **Legal basis for processing**

- Article 6(1)(a) of the GDPR – consent of the study participant to the processing of their personal data;
- Article 6(1)(e) of the GDPR – processing necessary for the performance of a task carried out in the public interest (conducting scientific research);
- Article 9(2)(a) of the GDPR – the participant's consent to the processing of special category data (e.g. health data);
- Article 9(2)(j) of the GDPR – processing necessary for the purposes of scientific research, in accordance with appropriate safeguards for the rights and freedoms of the participant.

5. **Categories of relevant data**

Personal data processed as part of scientific research may include:

- identification data: name, surname, PESEL, date of birth;
- contact details: telephone number, e-mail address, correspondence address;
- health data, including medical, diagnostic and other physical or mental health information;
- data related to participation in the study, e.g. test results, measurements, answers to survey questions, biological samples;
- other data required for specific research purposes.

6. **Information about the recipients of personal dataThe recipients of personal data will be:**

- persons authorized by the personal data administrator (employees and associates of Sano carrying out scientific research);
- entities authorised by law (e.g. supervisory authorities, research funding funders);
- providers of services supporting the implementation of research, including providers of information systems, analytical systems, research laboratories and archiving services. Personal data will be transferred only within the limits permitted by law and with appropriate safeguards.

7. **Period of storage of personal data**

Personal data will be stored:

- for the duration of the research project;
- for the period specified in the law on archiving research documentation and research results (usually at least 10 years from the end of the study);
- for the period indicated in the consent, if applicable.

8. **Rights of the data subject**

Persons participating in the research have the following rights:

- access to your data;
- rectify data;
- delete data (within the limits resulting from the provisions of law or the principles of conducting scientific research);
- restriction of processing;
- withdraw consent at any time, if consent was the basis for processing;
- object to the processing of data if the basis for the processing is the public interest or the legitimate interest of the administrator.

9. **Right to lodge a complaint with a supervisory authority**

A survey participant has the right to lodge a complaint with the President of the Office for Personal Data Protection if they believe that the processing of their personal data violates the provisions of the GDPR.

10. **Requirement to provide personal data**

Providing personal data is voluntary, but necessary to participate in scientific research. Failure to provide data will prevent participation in the study.

11. **Source of personal data**

Personal data is collected directly from study participants during the recruitment process or during the implementation of the study.