

## Extract from the Informed Consent Form

„According to the Order 904/2006 concerning the rules relating to the implementation of good practice in clinical trials, the basic principle is the protection of human rights and human dignity and the protection of personal data. (...)

Before being included in a study, you receive information about the objectives, risks and conditions which are conducting a clinical trial, as well as information regarding your rights to withdraw from the ongoing study. All this information can be found in the Informed Consent Form which is unique for each type of study, and signing and dating it at the beginning of each test, means your free agreement to attend the study. (...)

The personal data collected from you are related to: identification information, medical data and psychological data. (...)

According to the Law no. 677/2001 you have the right of access, intervention and the right not to be bound to submit an individual decision. Also, you have the right to request the change of your personal data, to oppose the processing of your personal data and to demand the withdrawal of the consent. To exercise these rights you can submit a written form, duly dated and signed, to the recruiting department.

Your collaboration with our company is conditioned by the existence of this agreement. This agreement may be withdrawn at any time without your justification only upon a written request addressed to the recruiting department. In this case your collected personal data will be stored to be accessed without being processed.

You may get information regarding your personal data whenever you want from the Recruitment Department.(...)

Your personal information will remain strictly confidential. (...)



**Recruitment Department**

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