

Information and expression of consent to the processing of personal data (GDPR N 2016/679)

Data controllers and related purposes

The Promoter of the Study (the National Cancer Institute – “G.Pascale” Foundation of Naples) that has been described to you, in accordance with the responsibilities established by the rules of good clinical practice (DLgs 211/2003), will process your personal data, in particular health data and, only to the extent that they are indispensable in relation to the objective of the study, other data relating to your origin, your lifestyle and your sexual life (etc.), exclusively for the purpose of carrying out the study. study and for pharmacovigilance purposes. To this end, the data indicated will be collected by the testing center and transmitted to the Study Promoter (the National Cancer Institute – “G.Pascale” Foundation of Naples). The processing of personal data relating to your illness and the response to the proposed treatment is essential for carrying out the study: refusal to provide them will not allow you to participate.

Nature of the data

Your name will only appear on the consent forms that you will be asked to sign. These forms will be sent in electronic format through an encrypted system that guarantees your privacy to the National Cancer Institute of Naples. The doctor who will follow you in the study will identify you with a code: the data concerning you collected during the study, with the exception of your name, will be transmitted to the National Cancer Institute – “G.Pascale” Foundation of Naples, recorded, processed and stored together with this code, at your date of birth, your sex, your weight and height, information on your illness and the outcome of your treatment. Only your doctor and authorized parties will be able to link this code to your name.

Processing methods (GDPR N 2016/679)

The data, processed using tools including electronic ones according to the methods established by GDPR N 2016/679, will be disseminated only in a strictly anonymous form, for example through scientific publications, statistics and scientific conferences. Your participation in the study implies that, in compliance with the regulations on clinical trials of medicinal products, the staff of the National Cancer Institute – “G.Pascale” Foundation of

Naples, the Ethics Committee and the Italian health authorities will be able to know the data concerning you, also contained in your original clinical documentation , in ways that guarantee the confidentiality of your identity.

Exercise of rights

You may exercise the rights referred to in the art. 7 of the Code (e.g. access your personal data, integrate them, update them, rectify them, oppose their processing for legitimate reasons, etc.) by directly contacting the Sponsor of the Study, the Tumor Institute of Naples (in the person of Dr. Raimondo by Giacomo coordinator of the study). You can interrupt your participation in the study at any time and without providing any justification. Furthermore, no further data concerning you will be collected, without prejudice to the use of any data already collected to determine, without altering, the results of the research.

Consent

By signing this form I consent to the processing of my personal data for research purposes within the limits and in the manner indicated in the information provided to me with this document.

Name and Surname of study partecipant (*capital letters*)

Signature _____

Date ____/____/____