As required in the Medical Research Decree, the actual Informed Consent Form must include the following:

1. Name, personal identity code or date of birth, and address the research subject
2. A mention that the information has been given to the research subject and data about the giver of the information
3. A mention of other sources where information concerning the research subject will be gathered from
4. Whom the information gathered in the context of the research can be delivered to and how the confidentiality of the information is protected
5. Research subject's voluntary consent
6. A mention of the right to withdraw consent without it affecting the research subject's right to receive the care needed
7. The consent document must be dated and signed both by the person who gives consent and by the person receiving the consent
8. If the study subjects consent is given verbally, a witness independent of the research shall sign the consent document.
9. A copy of the document is given to the giver of the consent