CERTIFICATION IN THE EUROPEAN AUTHORITY IN ACCORDANCE WITH EN ISO

European certification and, as a result, the CE marking is recognized in Europe as the only marking that guarantees compliance of products with the requirements of European directives and harmonized standards. CE marking allows you to sell your product in all European countries.

The manufacturer, by placing the CE mark on his products, makes it clear to the consumer and the supervisory authority that the product complies with European standards and directives and is safe and he is responsible for it.

The European certification center "MEDSTANDART" competently performs European certification work together with its European partners.

The main stages of certification for the manufacturer / importer:

1) definition of applicable directives and standards;

2) verification of product requirements;

3) Preparation of the necessary technical documentation;

4) determining the need for conformity assessment (notified body);

5) product testing and conformity testing;

6) marking and drawing up the Declaration of Conformity.

To determine the certification procedure with the right to mark products with the European CE mark, deadlines for the procedure, cost (price of the conformity assessment procedure), you need to send the following information to our e-mail:

- company name and contact details;
- name and purpose of products;
- technical documentation (available).

After receiving this information, we will respond to you within one business day.

Call or write to get a free consultation:
+38 (073) 4192745, +48 (791) 063674