



Documents required for the Dietary Supplements

Dietary Supplements Manufactured Inside the Eurasian Customs Union (EACU):

1. an application;
2. copies of documents in accordance with which the supplement is produced (standards, technical conditions, regulations, technological instructions, specifications, recipes, composition specifications), certified by the manufacturer;
3. a written notification from the manufacturer that the supplement meets the requirements of the documents in accordance with which it is made;
4. a manufacturer's document on the use (operation) of the supplement, (instruction, manual, datasheet, recommendations) or its copy certified by the applicant (if any);
5. copies of labels (packaging) or their layouts for applied goods, certified by the applicant;
6. copies of documents on the specific activity of the supplement (for products containing unknown components, unofficial recipes), certified by the applicant;
7. a sampling certificate;
8. a manufacturer's declarations on the presence of genetically modified (transgenic) organisms, nanomaterials, hormones, pesticides;
9. testing reports (hygienic certificates), scientific reports, expert opinions;

The applicant is responsible for the accuracy of the documents provided for issuing a document confirming the safety of products (goods).

