The list of documents required for the provision of state service when the Applicant (or his/her representative by power of attorney) applies in accordance with the Section II of the Unified list of goods subject to sanitary and epidemiological supervision (control) at the customs border and customs territory of the Customs Union, approved by the decision of the Commission of the Customs Union dated May 28, 2010 No. 299:

1) to obtain a certificate of state registration:

1. Certificate of the state registration of a legal entity

2.copies of documents in accordance with which the products are manufactured: standards, technical conditions, regulations, certified by the manufacturer (producer);

3. technological instruction
4. specification
5. formulation, information on the composition

6.written notification of the manufacturer (producer) stating that the products manufactured by it meet the requirements of the documents in accordance with which they are made;

copies of the quality certificate, safety (quality) data sheet, quality qualification certified by the manufacturer (producer) or letter of the manufacturer of products manufactured in the Eurasian Economic Union (one of the listed documents is provided);

7.copies of the quality certificate, safety (quality) data sheet, certificate of analysis, quality qualification, certificate of free sale or letter of the manufacturer (with the translation into Russian certified in the prescribed manner) of products manufactured outside the territory of the Eurasian Economic Union (one of the listed documents is provided);

8. document of the manufacturer (producer) on the application (operation, use) of controlled goods (instructions, guidelines, regulations, recommendations) or its copy certified by the applicant (if any);

9. copies of labels (packing) or their layouts for controlled goods, certified by the applicant;

10. copies of documents on the specific activity of biologically active food supplements (for preparations containing unknown components, unofficial formulations), certified by the applicant;

11. accredited laboratory's act of sample collection or copy of the document confirming the import of samples for testing (invoice);

Documents are submitted on two media (1 hard copy, 1 copy on electronic media).