Modul B

Type test ES

The EU-type examination test is that part of the conformity assessment which verifies and confirms that the design of the product meets the requirements of Directive 2014/28/EU.

The applicant shall send the technical documentation of the product to which he wishes to carry out the conformity assessment. The technical documentation must be in Slovak or English.

Notified Body shall check the technical documentation, if required requests additional information and send the applicant a price proposal for conformity assessment, including the determination of the place of where the tests will be performed, the number of test samples as well as the required activity and the date of receipt of the samples.

In case of approval of the price proposal, "Application for conformity assessment" form KOTA D-414 containing the conditions and the deadline for submitting the required samples, as well as the approval of the expected end date of the assessment, shall be send to the Notified Body. The form must be completed in full and legible. The veracity of the above data shall be confirmed by the applicant´s signature and a print stamp.

Subsequently, the Notified Body and the applicant shall agree on the location of the tests and the method of delivery of the samples.

The Notified Body shall draw up a draft contract incorporating all relevant and pre-discussed conditions for conformity assessment. Draft contract shall be send to the applicant by Notified Body for comments. Once the wording of the agreement is approved, both parties (the applicant's statutory representative and the statutory representative of the Notified Person) will sign the contract.

If on the basis of a signed contract the import of test samples is necessary from abroad, the Notified Body shall request the competent authority of the State Administration to issue an import license for the samples of explosives necessary to carry out the conformity assessment. NB shall also send the completed form 'Evidence on intra-Community transport of explosives' (Article 9 (5) and (6) of Directive 93/15 / EEC)to the competent authority.

Upon receiving the relevant license and transfer license, the Notified Body shall request from applicant to submit samples. Samples intended for conformity assessment need not be labeled according to the requirements of the relevant standards, but they must be clearly marked in a way that allows their unambiguous identification.

Upon recieving the samples, Notified Body shall perform the tests in accordance with the relevant harmonized standards. In the event of any inconsistency between test sample properties and the requirements, the testing is suspended. Information is sent to the applicant, including draft solutions to eliminate the nonconformity.

Check of the prescribed labeling is also part of the assessment. For this reason, the applicant must submit a labeling for each product under review before the end of the process.

If the product complies with the requirements of the Directive, the Notified Body will issue the final documents - the EU-type certificate and the final report. These documents, along with the invoice, will be sent to the applicant's address. The EU-type certificate is valid without any time limit.

If the product fails to meet the requirements of the Directive and the relevant harmonized standards, the Notified Body shall refuse to issue the certificate, giving the applicant detailed justification. At the same time, he will tell him about the possibilities and ways of appealing.

The applicant is required to inform the Notified Body about any modifications of the approved product which must be additionally approved if such changes may affect compliance with the essential requirements or the prescribed conditions of use of the product. This additional approval shall be issued in the form of an Annex to the original EU-type examination certificate.