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Status: EU Directives are being published on this site to aid cross referencing from UK legislation. After IP completion day (31 December 2020 11pm) no further amendments will be applied to this version.

ANNEX VI

CONFORMITY TO TYPE BASED ON PRODUCT QUALITY ASSURANCE FOR SAFETY COMPONENTS FOR LIFTS(module E)

- 4. Surveillance under the responsibility of the notified body
- 4.1. The purpose of surveillance is to make sure that the manufacturer duly fulfils the obligations arising out of the approved quality system.
- The manufacturer shall for assessment purposes allow the notified body access to the 4.2. premises where final inspection, testing and storage are carried out and provide it with all necessary information, in particular:
- the quality system documentation; (a)
- (b) the technical documentation;
- the quality records, such as inspection reports and test data, calibration data, reports (c) on the qualifications of the personnel concerned.
- The notified body shall periodically carry out audits to ensure that the manufacturer 4.3. maintains and applies the quality system and shall provide the manufacturer with an audit report.
- 4.4. Additionally, the notified body may pay unexpected visits to the manufacturer's premises where final inspection and testing of safety components for lifts are carried out.

At the time of such visits, the notified body may, where necessary, carry out tests or have them carried out in order to check the proper functioning of the quality system. It shall provide the manufacturer, with a visit report and, if a test has been carried out, with a test report.