



## **MINIMUM QC REQUIREMENTS FOR RE-HOMOLOGATION PROCESS FOR FHR TETHER SYSTEMS ACCORDING TO FIA STANDARD 8858-2010**

### **1. Foreword**

According to the **Re-homologation process-clarification note** available on the FIA website <https://www.fia.com/regulation/category/762>, manufacturers choosing option 1 for re-homologating their products need to present to their ASN a declaration and explanation of their internal quality control system (QC). As stated in the aforementioned document, in order for the QC system to be acceptable for approval, it will need to comply with some minimum requirements. This document describes the minimum requirements of the QC system that the manufacturer will need to have in place, as well as the documentation that is necessary to provide to obtain the re-homologation.

For clarity purposes, it has been deemed useful to specify the meaning of several expressions that will be used in this document and during the assessment process:

To MAINTAIN OBJECTIVE EVIDENCE refers to the manufacturer being able to provide justification that what was planned has actually been done. It is not necessary to keep records of the actual values, but it must be possible to demonstrate that the controls have been carried out.

To RETAIN DOCUMENTED INFORMATION refers to the manufacturer keeping records of the data of the checks (with values).

To MAINTAIN DOCUMENTED INFORMATION refers to the manufacturer being able to provide justification of documented processes and controls. This could be in the form of explicative documents, but it could also be for example, videos of the processes or photographs.

### **2. Minimum requirements**

#### **2.1 Processes control**

In order for the QC system to be acceptable, the company must maintain objective evidence of the following:

- Procurement process control
- Client order review and control
- Production order review and control
- Staff training (including new staff)
- Internal audits

In addition, the company must maintain documented information of the following:

- Production processes, including drawing controls and process change records
- Non-conformities management

#### **2.2 Traceability of materials and components**

The QC system must ensure that key raw materials and components for the product can be traced for each item produced. Documented information on the traceability must be retained.



Key materials are those that could directly affect the outcome of any of the tests defined in section 2.4. In the case of FHR tether systems according to FIA standard 8858-2010, the following groups of materials as a minimum are considered key materials:

- Webbing of tethers straps,
- Metals of end fittings,

### **2.3 Random testing of components and/or final products**

In order to control the final product performance it is compulsory that the QC system includes a random checking and testing programme to confirm that the production still complies with the requirements of the standard.

For all FIA standard 8858-2010 FHR tether systems it is necessary to perform and retain documented information of at least the following tests:

- Visual inspection:
  - 2% of the tethers;
- Mechanical strength and forward elongation test as defined in Art. 7.2 of the FIA standard 8858-2010:
  - 1 tether system every 2.5 years for productions of less than 2000 tethers/year;
  - 0.2‰ of the tethers for productions equal or higher than 2000 tethers/year;
- Flame resistance tests as defined in Art. 7.3 of the standard FIA 8858-2010:
  - 1 tether system every 2.5 years for productions of less than 2000 tethers/year;
  - 0.2‰ of the tethers for productions equal or higher than 2000 tethers/year;
- Cutting tests as defined in Art. 7. of the standard FIA 8858-2010:
  - 1 tether system every 2.5 years for productions of less than 2000 tethers/year;
  - 0.2‰ of the tethers for productions equal or higher than 2000 tethers/year;

These tests can be done internally in the manufacturer's facilities or externally. It is not necessary to use an FIA-approved test house. Variations of the mechanical strength tests can be used but only with prior approval by the FIA.



### **3. Documentation to be provided for re-homologation**

When applying for re-homologation using option 1, the manufacturer must submit to its ASN the Re-homologation Application Template and, in order to explain and declare its QC system, it must also submit the following information, depending on whether or not the manufacturer is certified according to ISO 9001:2015.

#### **3.1 Manufacturers not certified according to ISO 9001:2015**

- Declaration, in a company letterheaded document, filled in and signed, in accordance with:
  - Appendix I Processes control;
  - Appendix II Traceability of the materials and components;
  - Appendix III Random testing programme.
- Flow chart indicating when the controls declared in Appendix III are done during the production process.

#### **3.2 Manufacturers certified according to ISO 9001:2015**

- Copy of a valid ISO 9001:2015 certificate
- Declaration, in a company letterheaded document, filled in and signed, in accordance with:
  - Appendix III Random testing programme.
- Flow chart indicating when the controls declared in Appendix III are done during the production process.

### **4. Review and audits**

During the process of assessing the re-homologation request, the FIA reserves the right to request examples of the evidence and documented information required in section 2 of this document.

In addition, and as provided for under Article 6 of the FIA Homologation Regulations for Safety Equipment, the FIA reserves the right to perform audits to confirm that the manufacturer follows the quality control, and during which the manufacturer may be requested to demonstrate the veracity of its declaration and provide justification and records of the controls requested.



### Appendix I Processes control

This declaration shall be supplied on letterhead paper of the applicant company and signed (full name and position within the company required).

Mr/Ms [redacted] as [redacted] at [redacted] (the company) declares that the management of the company ensures that quality objectives have been defined and communicated throughout the company. The company follows a Quality Management System in order to ensure that production and procurement are carried out under controlled conditions and to ensure that the final product conforms to the requirements of the FIA standard for which they are homologated.

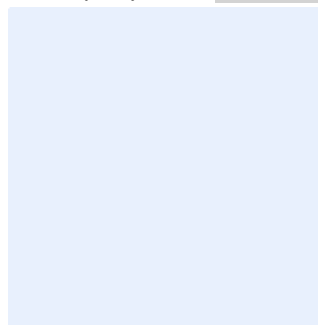
The company maintains objective evidence of the following:

- Procurement process control  
The company has processes in place to ensure that the products and services incorporated in the final product and supplied externally comply with the requirements and specification of the original homologated product.
- Client order review and control  
The company reviews the products that are going to be offered to customers in order to ensure that the requirements of FIA standard 8858-2010 are still complied with, and that no modification has been made with respect to the originally homologated product without authorisation by the FIA.
- Production order review and control
- Staff training (including new staff)
- Internal audits

In addition, the company maintains documented information of the following:

- Production processes, including drawing controls and process change records
- Non-conformities management

This Quality Management System has been in place in the company since [redacted]



Date: [redacted]



## Appendix II Traceability of materials and components

This declaration shall be supplied on letterhead paper of the applicant company and signed (full name and position within the company required).

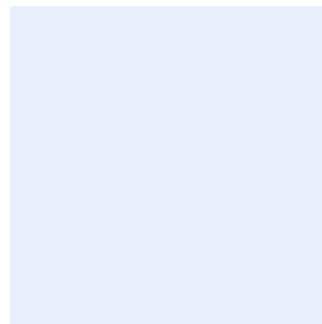
Mr/Ms [redacted] as [redacted] at [redacted] (the company) declares that the company retains documented information that allows all key materials of the products to be traced including information on the following:

- Supplier,
- Purchase date,
- Batch number,
- Controls or checks performed on arrival at the company.

The manufacturer is able to provide the above information on the following materials used in a specific tether system:

- Webbing of tethers straps;
- Metals of end fittings;

This traceability system has been in place in the company since [redacted]



Date: [redacted]



### Appendix III Random testing programme

This declaration shall be supplied on letterhead paper of the applicant company and signed (full name and position within the company required).

Mr/Ms [redacted] as [redacted] at [redacted] (the Company) declares that the information below is descriptive of the random tests done during the production of FHR tether systems according to FIA standard 8858-2010.

Tests	Test	How often?	Where are the tests done?
Visual inspection		tether system checked every	
Mechanical strength and forward elongation test as defined in Art. 7.2 of the FIA standard 8858-2010		tether system checked every	
Flame resistance tests as defined in Art. 7.3 of the standard FIA 8858-2010		tether system checked every	
Cutting tests as defined in Art. 7. of the standard FIA 8858-2010		tether system checked every	

Documented information of these controls is retained and can be provided if necessary.

These controls have been in place in the company since [redacted]

[redacted] Date: [redacted]