Table of Contents

0. Introduction ...........................................................................................................2
1. Language .............................................................................................................2
2. Numbering .........................................................................................................2
3. Naming ................................................................................................................2
4. Origin ..................................................................................................................2
5. Scope of PMA .....................................................................................................3
6. Recognized safe use (applies only where a history of safe use is claimed) ...........3
7. Contents of PMA ................................................................................................3
8. Structure of document .......................................................................................3
9. Verification testing requirements ......................................................................4
10. Certification and marking ................................................................................4
11. Information for further processing .................................................................4
12. Availability ........................................................................................................4
Appendix 1 ..............................................................................................................5
Appendix 2 ..............................................................................................................6
0. Introduction

A Particular Material Appraisal (PMA) is the process by which the pressure equipment manufacturer ensures that each proposed material that is not in a harmonized standard or covered by a European Approval for Materials (EAM) conforms to the applicable Essential Safety Requirements (ESR) for materials. Particular material appraisals are part of the “records and correspondence relating to conformity assessment” referred to in PED Article 10 (4) and therefore subject to the requirements, which include the preparation of appropriate documentation.

As required by the Pressure Equipment Directive 97/23/EC (PED) Annex I section 4.2, the pressure equipment manufacturer is responsible for preparing documentation for materials to be used for pressure equipment in categories I to IV.

For pressure equipment in categories III and IV this documentation is submitted to the Notified Body in charge of the conformity assessment for the equipment so they can perform an appraisal of the particular material with regard to the pressure equipment, the intended use and applicable design rules.

This documentation shall describe the material properties in a manner that is concise, complete and correct. In general, rules for calculation, design and manufacture relating to the specific material should be considered in the PMA.

Decisions made concerning a specific material for use in a specific sets of conditions, do not need to be repeated. However, manufacturers and where applicable notified bodies must ensure the PMA is suitable for its application and that this is recorded in the technical documentation.

Pressure equipment manufacturers may utilize PMAs prepared and supplied by another manufacturer, e.g. the manufacturer of an assembly, on condition that the PMA is reviewed and adopted by the PE manufacturer and where applicable the NB in charge of the conformity assessment procedure, to ensure accuracy of the document and suitability for the application.

1. Language

No specific requirements, however generally in the same language as the technical documentation.

2. Numbering

No specific requirements however shall be traceable to the application and material specification.

3. Naming

It is not mandatory to provide a material name, however the identification of the material shall be given.
4. Origin

The material specification shall be indicated with its date. Where this is a publicly available specification the actual data need not be included in the text of the PMA. However, supplements to or deviations from these specifications shall be stated. PMAs related to publicly available specifications are only valid for the particular dated version. If the basic specification is not publicly available, it shall be attached to the PMA.

5. Scope of PMA

The document shall describe the proposed application and range of conditions that are applicable; these must be supported by assured properties in the material specification.

6. Recognized safe use (applies only where a history of safe use is claimed)

In the case of materials recognized as being safe to use before 29 November 1999, the manufacturer and/or the notified body shall take account of the existing data when certifying such conformity.

7. Contents of PMA

This will comprise qualitative and quantitative data.

Qualitative data includes delivery conditions, and where applicable, heat treatment requirements, and information on further processing and weldability. Quantitative data will comprise mechanical data, creep resistance etc. where this differs from or supplements that in the specification adopted.

The material grouping in accordance with CR ISO 15608:2000 is relevant where welding is undertaken.

Evidence shall be provided that the relevant ESRs have been met.

Any restrictions applicable to the use of the material shall be stated, e.g. extent of cold work, special controls for permanent joining, etc.

Note: Although the directive does not require PMAs for welding consumables (See WGP Guideline 7/12), however, it may be appropriate to utilize this form of documentation to demonstrate conformity with the ESR 4.2(a) - (see WGP Guideline 7/10).

8. Structure of document

There is no format specified in the directive, however an example of a suitable layout is given in Appendix 1. The information required may also be given elsewhere in the technical documentation of the pressure equipment.

The equipment manufacturer must also address the ESRs relating to material selection. This format could be extended to include documentation of ESRs relating to selection, e.g. PED Annex I – 4.1(b), (d) and (e).
9. Verification testing requirements

In general, testing by the material manufacturer (sampling, frequency of testing, and test method) is to be performed as foreseen in the original specification. However, the limitations in Appendix 2 apply.

10. Certification and marking

The pressure equipment manufacturer is required to obtain certification from the material manufacturer that affirm compliance with "a specification and comply with PED Annex I, 4.3 and relevant guidelines.

11. Information for further processing

See Section 7

Verification testing may be required in cases where the properties of the material have been altered by processing operations performed subsequent to the material manufacturing process. Where required testing shall be performed as specified in the original specification.

For example, where plate material is purchased from a material manufacturer and is subsequently cold formed into a dished end, the mechanical properties will be altered. In such cases it may be necessary to “verify” that appropriate properties are still available in the material and continue to meet the applicable ESRs.

12. Availability

PMAs shall be available for national authorities upon request for a period of 10 years after the last pressure equipment has been manufactured. Contrary to the requirements for EAMs, there is no requirement to make PMAs publicly available. Documentation to demonstrate conformity with the ESR 4.2(a) - (see WGP Guideline 7/10).
## Appendix 1

<table>
<thead>
<tr>
<th>Property</th>
<th>Requirement</th>
<th>Compliance</th>
</tr>
</thead>
<tbody>
<tr>
<td>Appropriate properties</td>
<td>e.g. Yield (design temp): Re/t&lt;br&gt;Ultimate: Rm/20 if required Ultimate (design Temp) Rm/t</td>
<td>Suitable properties for design and testing with average safety factors of XYZ</td>
</tr>
<tr>
<td>Sufficiently ductile</td>
<td>In steel; 14 %</td>
<td></td>
</tr>
<tr>
<td>Sufficiently tough</td>
<td>e.g. in steel; 27 Joule at 20 0C or lowest operating temp.</td>
<td>required with PO, min allowable value 27 J @ -20 F</td>
</tr>
<tr>
<td>Not significantly affected by ageing</td>
<td>At intended operating temperatures</td>
<td>N/A for this CS</td>
</tr>
<tr>
<td>Suitable for intended processing</td>
<td>e.g. welding, forming, etc</td>
<td>Yes</td>
</tr>
<tr>
<td>Material Marking</td>
<td>Identifiable to Certificate</td>
<td>Material traceable to specific MTR</td>
</tr>
<tr>
<td>Type of Certificate</td>
<td>Cat I; Certificate of Conformance</td>
<td>Nozzles: EN 10204-2.1 / -2.2 (generic MTR)</td>
</tr>
<tr>
<td></td>
<td>Cat II, III, IV; – e.g EN 10204 3.1.B</td>
<td>EN 10204-3.1B / 3.2 (specific MTR)</td>
</tr>
<tr>
<td>Material Manufacturers Approval</td>
<td>ISO 9000 by Euro Accredited Body</td>
<td>Nozzles: No / Piping</td>
</tr>
<tr>
<td></td>
<td>Independent test e.g EN 10204 3.1.C or 3.2.</td>
<td>See others 2</td>
</tr>
</tbody>
</table>

The named material specification may be used for the construction of pressure equipment intended for use within the stated limits and subject to the following restrictions:

**Restrictions:**

**Additional Testing:**

**Notes:**

______________________________  ___________________________
Signature                            City / Date
Pressure Equipment Manufacturer

______________________________  ___________________________
Signature                            City / Date
Notified Body (Cats III & IV only)

**Others:**

1) Materials used for equipment in category II, III or IV shall be supplied with a certificate of specific product control. If a certificate 3.1.B or 3.1.C according to EN 10204:1991 is used an affirmation of compliance with the specification must be given in the certificate, or in a separate document.

2) If material certificates issued by the material manufacturer shall be accepted he must hold at least an ISO 9002-certificate issued by a competent body established within the Community and having undergone a specific assessment for materials.
Appendix 2

Supplementary Testing Requirements

For new materials with limited data or for situations where supplementary requirements have been included in the PMA, the following applies:

A test program will be developed to produce results, which are of sufficient reliability for the intended use. It is recommended that a notified body is involved with agreeing the test program and witnessing the resulting testing. EN 764-4 provides guidance on the preparation of test plans. Consideration will be given to the number of samples, the variability of the materials, known facts, which could invalidate the results. E.g. impact properties on high sulphur casts and/or with an adverse Mn:C ratio.

The results obtained shall not be used to justify increases in allowable stress on the basis of individual tensile test values.

In general, testing (sampling, frequency of testing, and test method) is to be performed as foreseen in the original specification.

However,

- the testing methods have to be compatible with the PED requirements. In particular, the test results have to be consistently comparable;
- if a European testing standard (EN or EN ISO) is available, the testing method employed shall be replaced by the tests contained in the respective EN (or EN ISO) unless the testing foreseen in the original specification better reflects the state of the art than the corresponding EN (EN ISO). Examples of such standards are:

  - EN 10002-1 Tensile testing at room temperature
  - EN 10002-5 Tensile testing at elevated temperatures
  - EN 10045-1 Charpy impact test
  - EN ISO 3651-2 Intergranular corrosion resistance

Depending on product form and type of material, the sampling and frequency of testing shall as a minimum be based on the analogous European standard (EN). e.g.: EN 10028-1 and –7 for flat products, EN 10213-1 for castings, and EN 10222-1 for forgings.