

Application for independent verification/validation of environmental product declarations (EPD) that have successfully passed a verification/validation of an established ECO Platform Programme Operator

A prerequisite for the acceptance of an order is a fully filled-in application form.

Manufacturer company:		
Address:		
Plant(s) (with complete address(es) if different from company)		
Contact		
Function		
Telephone	E-mail	
<i>important:</i> For the plant, a distinction must be made between the <u>billing</u> address, <u>plant</u> address(es) and <u>correspondence</u> address if applicable. Please provide appropriate information. We will be happy to handle all correspondence and invoicing by e-mail. Please provide a central digital billing address if relevant.		
Declaration holder if different from the manufacturer, institution and address		
Author of the life cycle assessment Institution/address/tel./e-mail		
	Indication of whether internal or external assessment is used; in the case of external LCA practitioners, please indicate institution, persons (necessary for comparison with the team of verifiers and for checking impartiality)	
ECO Platform programme operator that verified the first version of EPD	Name, address, country, website	
Titel and unique identification number of EPD(s)	TEXT	

Product description:

The declared products must be described (please list separately if there are different products/product groups).

In addition to a general product description, the trade names of the products/product groups (including any product codes) to which the EPD applies must be stated.

If it is not possible to specify trade names, e.g. in the context of association EPDs, the product description must clearly distinguish the products/product groups.

Product definition (please select one of the following options and delete the alternatives that do not apply)

Product 1/Product group 1

Product description TEXT

Alternative 1a: Product according to CPR with hEN:

Regulation (EU) No. 305/2011 (CPR 'old') or (EU) No. 2024/3110 (CPR 'new') applies to the marketing of the product in the EU/EFTA (with the exception of Switzerland). The product requires a declaration of performance in accordance with the CPR, taking into account EN xyz: date, title and CE marking.
The respective national regulations apply to its use.

Alternative 1a: Product according to CPR with ETA:

Regulation (EU) No. 305/2011 (CPR 'old') or (EU) No. 2024/3110 (CPR 'new') applies to the marketing of the product in the EU/EFTA (with the exception of Switzerland). The product requires a declaration of performance taking into account ETA xyz: date, title and CE marking.
The respective national regulations apply to its use.

Alternative 2a: Product that is not harmonised under the CPR but under other regulations in the EU:

The following EU harmonisation regulation(s) apply(ies) to the placing of the product on the market in the EU/EFTA (with the exception of Switzerland):

- Directive No. xyz, date, title
 - Regulation No. xyz, date, title
- and their harmonised standard(s) based on this:

- EN xyz: date, title

The CE marking is affixed to the product in accordance with the proof of its conformity with the following harmonised standards based on the aforementioned harmonisation regulation(s):

- EN xyz: date, title

The respective national regulations apply to its use.

Alternative 2b: Product harmonised in the EU on the basis of both the CPR and other regulations:

For the product to be placed on the market in the EU/EFTA (with the exception of Switzerland), Regulation (EU) No. 305/2011 (CPR 'old') or (EU) No. 2024/3110 (CPR 'new') and the following EU harmonisation regulation(s) apply:

- Directive No. xyz, date, title
- Regulation No. xyz, date, title

The product requires a declaration of performance in accordance with the CPR, taking into account EN xyz: date, title or ETA No. xyz: date, title and the CE marking.

The CE marking is affixed to the product in accordance with the declaration of performance in accordance with the CPR and proof of its conformity with the following harmonised standards based on other harmonisation regulations:

- EN xyz: date, title

The respective national regulations apply to its use.

Alternative 3: Product not subject to EU harmonisation legislation:

The respective national regulations at the place of use apply to the use of the product, for example the building regulations of the federal states and the technical regulations based on these regulations.

Product 2

As above for product 1

Etc.

2 Number of EPD documents and project reports, number of data sets/average data sets

(Info: The LCA-practitioner can make useful suggestions here)

Text example (please adapt): X project report(s), X EPD documents with 1 data set each

3 Background database(s)

Specification of the background database used for upstream and downstream data sets, unless generic data is available (e.g. Ecoinvent or Sphera MLC (formerly GaBi) or others):

Name of the database:

Please send us an offer for the requested verification/validation of the EPD(s).

The applicant acknowledges that the programme operator Bau EPD GmbH performs independent verifications/validations of environmental product declarations (EPDs) in accordance with EN 15804 and ISO 21930 based on EN ISO 14025. At the same time, the applicant acknowledges the product category rules (PCR) that are published in this context on the website of Bau EPD GmbH and apply in addition to the standards, as well as the regulations of the programme operator Bau EPD GmbH for the verification/validations of EPDs (MS-HB and the pertinent applicable documents).

After the acceptance of the order has been confirmed by Bau EPD GmbH, the client agrees to provide all documents belonging to the EPD to be verified/validated (in particular the EPD document, the project report and the life cycle inventory analyses, etc.) to the programme operator Bau EPD GmbH if the applicant agrees with the offer of Bau EPD GmbH. For this purpose, a separate contractual document must be signed which also contains necessary confidentiality agreements (M-Doc 03).

The applicant takes note of the process flow of the conformity assessment programme according to M-Doc 26 and is aware of the associated obligations to cooperate.

Place/date/stamp

**Legally valid
signature(s)**

The following areas must be filled in by the programme operator:

Assessment of the completeness of documentation for an offer:

Complete _____

Incomplete, missing documentation:

Previous experience with the requested services _____

PCR available/to be prepared _____

Verifiers approved for the product group _____

Resources available in the planned project period _____

Restrictions for verification/validation if relevant _____

Impartiality of the certification body guaranteed _____

Impartiality of the verifier guaranteed _____

Decision by _____ *dated* _____:

The offer is made with M-Doc 03 _____

The offer cannot be made because _____