

Fire Detection and Fire Alarm Products

Please complete the form electronically where possible. If completing by hand, please use BLOCK CAPITALS and BLACK INK

Please submit this application form with the completed Technical Appendices as appliable to the product type(s). See Section 6.

SECTION 1 – General information

1.1 Type of product(s) requiring certification Please state below the type of product(s) to be certified (e.g., smoke detector, sounder, etc). If there are multiple product types, please list them here. Please select the type of application to be submitted:

New product application

Modification(s) to a certified product \Box

Change due to component shortage (Fast Track required)

and UKNI Marking are delivered by BRE Global (UK) Ltd.

1.2 Required Certification Marks

Please select which certification marks are required. Please note: To use the multiple certification process at least two certification marks must be selected.

See also Section 7 - Contractual information. Upon selection, the contractual statements of the related certification bodies will be labelled as APPLICABLE in Section 7

Please tick required certification marks



Please select the certification body you want to manage the project.

(Note: The chosen certification body (CB) will become the primary certification body for the project. In the case of modifications or updates to an existing product the primary certification body is likely to have issued the Constancy of Performance certificate. See below).

AFNOR Certification	BRE Global	=	DBI fication	VdS			
1.3 Constancy	of Performance ce	rtificate					
Do you require a 'Co	onstancy of Performance	e' certificate to sup	port the following?	a)	CE marking		The Construction Product Regulation
					b) UKCA marking		(CPR) requires fire detection equipment placed on the European market to have a
				c)	UKNI marking		Constancy of Performance certificate and be suitably marked.
Which certification b	oody do you want to carr	y out the assessme	ent and verification of	of constancy o	f performance in	accordar	nce with the CPR?
AFNOR	BRE	CNPP	DBI			~	of Performance certificates to support CE delivered by BRE Global (Ireland) Ltd.
Certification	Global ¹⁾	Cert.	Certification	VdS		-	f Performance certificates to support UKCA

1.4a Details of the Applicant – This will be the company that is authorised to apply for multiple certification

Registered company name:

Please provide evidence of company registration (e.g. Register of Commerce)

VAT No.

Address:

Please state the name that should appear on the certificate:

Nominated contact within the Applicant company:

Title:

Name:

Contact details Email:

Position:

Telephone:





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1.4b Details of the Applicant – Continued

There are circumstances where there is a requirement to include a second organisation associated with the applicant that handles specific functions, such as reviewing/accepting quotations, processing invoices or answering technical/administrative questions.

An example where this circumstance might be applicable, is a parent company that owns a group of companies, one of which applies for multiple certification and some of the functions normally attributed to the applicant are performed by the parent company. If this situation or a similar applies please complete the details of the second organisation associated with the applicant. Note, the name of second organisation will not appear of the certificate.

Registered company name:

Please provide evidence of company registration (e.g. Register of Commerce)

	VAT No.					
	Address:					
Nominated contac	Nominated contact within the Applicant company:					
Title:	Name:	Position:				
Contact details E	imail:	Telephone:				
Please state the relationship with the Applicant company declared in section 1.4a						

1.5 Details of the European Agent(s)

Please enter the details of the legal entities (European Agents) acting on behalf of the Manufacturer located outside of the European Economic Area (EEA) and/or the United Kingdom.

	Not Applicable	If No European Agents are required, please select Not Applicable		
Company name:				
Address:			CE Marking	
Company name:				
Address:			UKCA Marking	

I hereby authorize the companies named above to represent me throughout the EEA and the UK on all matters relating to the use of the CE/UKCA marking (*tick boxes as applicable*). I commit myself to immediately inform the primary CB whenever a new authorized agent, as referred to above, is appointed.

Nominated contact

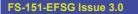
Title:	Name:	Position:
Contact details	Email:	Telephone:

1.6 Details of existing or previous certification(s)

Please select the certification bodies that currently audit or have previously audited the designated manufacturing site, and enter the certification reference (i.e., the certificate number).

If the designated manufacturing site(s) have no products currently certified select N/A

AFNOR Certification	DBI	
BRE Global (LPCB)	VdS	
CNPP Cert.	N/A	







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SECTION 2 – Manufacturing sites

Please enter the details of each company (if more than one) involved with the manufacture of the product(s) for which certification is required

2.1 Details of n	nanufactu	ring company (Sit	e1)			
Company name:						
Address:						
Indicate the activities	s performed	at site 1 Select all the	at app	ly		If other please provide details
Design		Manufacture		Assembly		Final test
		Marking		Management of modifications		Other
Nominated contact	within the Ma	nufacturing company 1				
Title:	Name:				Pos	osition:
Contact details	Email:				Telepl	phone:
Legal representative	e within the M	lanufacturing company	1			
		the person who overse actual declarations of S			mpany. I	. IMPORTANT - By signing this Application form the person
Title:	Name:				Pos	osition:
Contact details	Email:				Telepl	phone:
Się	gnature					Date:
Is the Applicant and	d the Manufa	acturing Site the same e	entity?	Yes No		

2.2 Manufacturing company (Site 2) Please complete this section if there is more than one manufacturing site involved in the production of the product(s) to be certified Not Applicable: If there is No additional Manufacturing site 2, please select Not Applicable Company name: Address: Indicate the activities performed at site 2 Select all that apply If other please provide details Design Manufacture Assembly Final test

Management of modifications

Nominated contact within the Manufacturing company 2

Marking

Title: Name:

□ Inspection

Contact details Email:

Position:

Other

Telephone:





Date:

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Legal representative within the Manufacturing company 2

This section must be signed by the person who oversees the legal affairs of the manufacturing company. **IMPORTANT -** By signing this Application form the person named below accepts the contractual declarations of Section 7.

Title:	Name:			_		Pos	sition:			
Contact details	Email:				Telephone:					
Sig	gnature			Date:						
Is the Applicant and	the Manufac	turing Site the same en	tity?	Yes	ю					
2.3 Manufacturi	ng compa	ny (Site 3)								
Please complete this	s section if th	nere are more than two i	nanuf	acturing sites involve	d in the productio	on of the	e product(s) to be certified			
	Not Applicab	ble:		If there is No a	additional Manufa	acturing	site 3, please select Not Applicable			
Company name:										
Address:										
Indicate the activities	s performed a	at site 3 Select all tha	t apply	Ý			If other please provide details			
□ Design		Manufacture		Assembly			Final test			
		Marking		Management of mo	odifications		Other			
Nominated contact	within the Ma	anufacturing company	3							
Title:	Name:					Pos	sition:			
Contact details	s Email: Telephone:									
Legal representative	e within the I	Manufacturing compan	y 3							
		the person who oversed actual declarations of S			anufacturing corr	npany. II	MPORTANT - By signing this Application form the person			
Title:	Name:					Pos	sition:			
Contact details	Email:					Teleph	hone:			

Is the Applicant and the Manufacturing Site the same entity? Yes No

Signature





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SECTION 3 – Financial

3.1 Billing information

Please enter the details of the legal entity responsible for paying the test and certification fees

Same as Applicant	(Section 1.4)	If the legal entity responsible for paying the test and certification fees is the same as the Applicant, please check box and jump to Section 4
Registered cor	mpany name:	
Please provide evid	dence of company reg VAT No. Address:	gistration (e.g. Register of Commerce)
For France: SIRET No.		APE Code:
Nominated contact	The nominated	l contact must be the person responsible for authorising payment of the test and certification fees
Title:	Name:	Position:
Contact details	Email:	Telephone:

SECTION 4 - Quality Management Systems & Factory Production Control of manufacturing

4.1 Contacts responsible for the Quality Management System

Please enter the contact details of the primary contact within the manufacturing site(s) responsible for the Quality Management System.

Contact person name:	Site 1	Telephone (Direct dial number	email
	Site 2	preferred)	
	Site 3		
Please enter the details	of existing ISO 9001 registrations		

4.2 ISO 9001 registered companies				
Are all the manufacturing sites ISO 9001 registered?	⊖ YES	O No	but wish to apply for registration	<i>Please give details below and complete Sections 4.3 and 4.4 as applicable</i>
		O No	and do not wish to apply for registration	Jump to Section 5

Name the organisation(s) from which the ISO 9001 registration was obtained:

Site 2

Site 3

Please confirm the respective certificate numbers:

Expiry Date of the certificate(s):

Site 1

Please provide copies of the ISO 9001 certificates for our records





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4.3 Companies that have no ISO 9001 registration but have completed an application for registration

Please answer the following questions if the manufacturing site(s) have no current ISO 9001 registration but an application for registration made has been submitted

To which accredited organisation has the application for registration been made?		Site 2	Site 3
What is the expected date the certificate(s) for registration is(are) to be granted?			
4.4 Companies that have no ISO 9001 registration but wis	sh to	apply for registration	
If applying for ISO 9001 registration please check the following boxes as	applica	able, to confirm the information supplied	
Current quality manual submitted with this application form		Technical documentation - Installation manuals	s, drawings, part lists, schematics, etc
Please provide an English language copy of the quality manual		Critical component list, including the supplier ic	lentification (number and name)
Product data sheets/Product specifications		Other relevant documentation submitted with the	nis application form
SECTION 5 – Additional information			
Country/territory specific requirements			
Please indicate to which other authorities the product certificates will be s	ubmitte	ed:	
Dubai Civil Defence (For further information see separate Dub	ai Civi	I Defence Submission Guidance)	
If applicable, please list any other authorities			

Other information

If you wish to provide further information or specific instructions, please use the space below





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SECTION 6 – Technical information

Complete one or more of the Appendices as applicable to the type of product(s) for which certification is required.

Please submit the completed Appendices with your application.

Please check all boxes that apply

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	APPENDIX 1	Included standards SOUNDERS (EN 54-3:2001 + A1:2002 + A2:2006) SOUNDERS (EN 54-3:2014 + A1:2019) VAD (EN 54-23: 2010) SHORT-CIRCUIT ISOLATOR (EN 54-17:2005 + AC:2007) INPUT/OUTPUT DEVICE (EN 54-18:2005 + AC:2007) COMPONENTS USING RADIO LINKS (EN 54-25:2008 + AC:2010 + AC:2012)
	APPENDIX 2	SMOKE (EN 54-7:2018) HEAT (EN 54 5:2017+A1:2018) SHORT-CIRCUIT ISOLATOR (EN 54-17:2005 + AC:2007) INPUT/OUTPUT DEVICE (EN 54-18:2005 + AC:2007) COMPONENTS USING RADIO LINKS (EN 54-25:2008 + AC:2010 + AC:2012) COMBINATION OF SMOKE AND HEAT SENSORS (EN 54-29:2015) COMBINATION OF CO AND OPTIONALLY HEAT DETECTORS (EN 54-30:2015) COMBINATION OF SMOKE, CO AND OPTIONALLY HEAT SENSORS (EN 54-31:2014 +A1:2016)
	APPENDIX 3	LINE DETECTORS USING AN OPTICAL BEAM (EN 54-12:2015) SHORT-CIRCUIT ISOLATORS (EN 54-17:2005 + AC:2007) INPUT/OUTPUT DEVICES (EN 54-18: 2005 + AC:2007) COMPONENTS USING RADIO LINKS (EN 54-25:2008 + AC:2010 + AC:2012)
	APPENDIX 4	MANUAL CALL POINT (EN 54-11:2001 + A1:2005) SHORT-CIRCUIT ISOLATORS (EN 54-17: 2005 + AC:2007) INPUT/OUTPUT DEVICES (EN 54-18: 2005 + AC:2007) COMPONENTS USING RADIO LINKS (EN 54-25 :2008 + AC:2010 + AC:2012)
	APPENDIX 5	FLAME (EN 54-10:2002 + A1:2005) SHORT-CIRCUIT ISOLATORS (EN 54-17:2005 + AC:2007) INPUT/OUTPUT DEVICES (EN 54-18:2005 + AC:2007) COMPONENTS USING RADIO LINKS (EN 54-25:2008 + AC:2010 + AC:2012)
	APPENDIX 6	POWER SUPPLY EQUIPMENT (EN 54-4:1997 + A1:2002 + A2:2006) POWER SUPPLIES (EN 12101-10:2005 + AC:2007)
	APPENDIX 7	CONTROL & INDICATING EQUIPMENT (EN 54-2:1997 + A1:2006) POWER SUPPLY EQUIPMENT (EN 54-4:1997 + A1:2002 + A2:2006) POWER SUPPLIES (EN 12101-10:2005 + AC:2007)
	APPENDIX 8	SMOKE ALARM DEVICES (EN 14604:2005)

When complete, each Appendix provides the technical information necessary to generate the quotations.

Applicants should expect a quotation from each Certification body selected in Section 1. These quotations cover the activities that each certification body (and where

applicable, the associated testing laboratories) must undertake to complete the multiple certification process.

To comply with Competition Law each certification body (and where applicable, the associated testing laboratory) must issue its own (separate) quotation to the Applicant.





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SECTION 7– Contractual information

In accordance with the certification guidelines of ISO-IEC 17067 a contract shall exist between the certification body and the Applicant. Each EFSG member certification body has its own contractual requirements. To permit the use of a common application form, the Applicant and legal representatives of the associated Manufacturing companies must agree with following declarations as applicable to the certification marks required. The contractual statements that apply based upon the selection of the required certification marks (Section 1.2), are labelled "APPLICABLE".

7.1 AFNOR Certification

HOLDER'S COMMITMENT TO THE NF MARK:

For this purpose, I declare that I know and accept the General Rules of the NF brand, the certification rules NF508 – fire safety system (NF-SSI) its appendices and I undertake to respect them throughout the lifetime of use of the NF brand. I also undertake not to use the NF mark before obtaining the right of use. In addition, I note that the NF mark is assigned to a central component OR hardware in a certified NF (SSI) system and in this case only. I certify that the product(s) meets the regulatory requirements applicable to them and undertake not to present for certification counterfeit products.

I undertake to communicate to my subcontractor(s) the general rules of the NF mark as well as the NF-SSI certification rules.

Commercial reference:

Commercial Brand:

In case the product is a component in according to the NF terms please indicate the SSI number:

In case of mutual request (sub-contractor, nominated representative or distributor) please tick one of the boxes and complete the address and the legal name, this document should be signed by the both (nominated representative + subcontractor or distributor):

OPTION 1: The central equipment OR component is designed by the subcontractor OR supplier, making a joint application, in the following design unit: "legal name + full address"

OPTION 2: The central equipment OR component is manufactured by the manufacturer OR subcontractor OR the supplier, making a joint application, in the following manufacturing unit(s): "full address name".

7.2 BRE Global (UK) Ltd and BRE Global Ireland Ltd

The contractual declarations required by BRE Global (UK) Ltd and BRE Global Ireland Ltd are incorporated into their corresponding quotation templates and the associate terms and conditions. Agreement with the BRE contractual arrangements is required upon acceptance of their quotation(s). The quotation(s) will be generated in response to the receipt of this application form.

7.3 CNPP Cert

COMMITMENT

I would like to request a certificate of constancy of performance issued by CNPP Cert allowing the producer to affix the CE mark on the abovementioned product(s), in compliance with the enclosed technical file.

I make a commitment not to present for this, product(s) where other simultaneous requests for a certificate of constancy of performance have been made. Nor to maintain for the same product, another certificate of constancy of performance issued by another certification body (please attach a copy of a valid certificate of constancy of performance for any products already certified by another notified body).

I declare that I have been informed about, and understand and agree with; the reference document A201 Procedure, including its appendices and application circular letters and the applicable fees.

Within the frame of the obligations associated with this application request, I commit myself to abide by the rules prescribed by these documents now and in the future, for the duration of the use of the certification, and especially to:

Is from ONDE O at immediately about an immediately and to the number (1) and information of an immediately and the

- Inform CNPP Cert, immediately, about any important change to the product(s), organization, personnel and associated equipment, etc...;
- Facilitate the mission of every person mandated by CNPP Cert. in order to perform the requested inspection(s) as required by the reference document;
- Not to abuse the certification;
- Make all the requested payments according to the reference document, no matter which decision is taken, i.e., agreement or refusal of the certification. In this regard, I validate that this present agreement serves as a placement of the order for the audit service prescribed in the reference document, without sending an individual purchase order for every occurrence.
- I am aware this letter and the reference document referred to above, addresses the obligations, rights and requirement of each party and constitutes together a certification contract.
- I hereby authorize CNPP Cert. to go ahead with any and all exchanges necessary with the CNPP laboratories.





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7.4 DBI Certification

DECLARATION

I, the applicant, have completed this application and established the facts presented in this application. I declare that the information provided in this application is true and correct and that the same application has not been lodged simultaneously with any other certification body.

I agree that the DBI Certification A/S and the DBI Certification UK Ltd Agreement and Service Terms shall govern the evaluation pursuant to this application, and to provide information required to perform the evaluation.

I declare that the samples placed at the disposal will be fully representative of the production envisaged, and agree to provide further specimens if needed for carrying out the eventual test programme.

Where historical data is submitted as part of the evaluation, I declare that the product tested to obtain the historical data is fully representative of the current production or I have identified any modifications to the design, production processes or materials.

I declare that should any significant changes in the product(s) or in my organisation or any other activity that could affect conformity with the standards that the product(s) is(are) evaluated against occurs, DBIC and DBIC-UK will be informed.

Were DBI has been chosen to provide the assessment and verification of constancy of performance in accordance with the CPR (See Section 1.2), I understand that DBI Certification A/S has the role of notified body in accordance with the product certified and all obligations regarding the CE-CPR and CE-MED certification are towards DBI Certification A/S. I understand that DBI Certification UK Ltd has the role of approved body in accordance with the product certified and all obligations regarding the UKCA-CPR certification are towards DBI Certification UK Ltd. DBI Certification A/S will be the main point of contact, and provide the quotation, testing arrangements, and Factory assessments.

I, the applicant, authorize DBIC and DBIC-UK to proceed with this as an active application evaluation against the Standards required by the CPR/MED/UKCA Program(s).

7.5 VdS Certification

DECLARATION

Data protection

As the responsible body, VdS Schadenverhütung GmbH collects and processes personal data in the course of the procedures described here only to the extent necessary for the performance of the contract (chapter II, art. 6, no. 1 lit. b), GDPR) or on the basis of a specific declaration of consent by the person concerned (chapter II, Art. 6, no. 1 lit. a), GDPR) which can be revoked informally at any time.

For further information on data protection, please refer to the General Terms and Conditions (VdS 3177) or the information on our website (https://vds.de/de/unternehmen/datenschutz/).

Declaration and consent

I (we) agree that,

- the General Terms and Conditions of VdS certification body, VdS 3177,
- the guidelines Procedure for Testing, Approval and Certification of Products and Systems for Fire Protection and Security Technologies, VdS 2344 as well as the guidelines Procedure for the Performance of Product Surveillance, VdS 2841,
- the associated price list or the offer,

form the basis of this order and accept them in the respective valid version as an integral part of the contract.

Furthermore, I (we) consent to,

- VdS Schadenverhütung GmbH collecting, processing and using personal data or other data within the context of delivering the contract,
- VdS Schadenverhütung GmbH making the certification accessible to third parties (i.e., EFSG member companies) by publishing it in freely accessible directories,
- Documents (e.g., order confirmations, test reports, draft certificates) being transmitted by e-mail. This shall be done exclusively to the e-mail address of the contact person stated under "applicant". Transmission to another e-mail address requires the written consent of this contact person.





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By signing this application form the Applicant and the legal representatives of the manufacturing companies/sites (Section 2) confirm their acceptance of the following;					
Select as appropriate					
The applicable declarations of Section 7 based on the selection of the required certification marks of Section 1.2 of this application form					
Consent to permit EFSG and all its certification body members named in Section 1.2 (and their associated testing laboratories) to exchange information related to this project that includes but is not limited to; product samples, documentation and test results.					
By agreeing to give consent the Applicant absolves the above from of their confidentiality obligations associated with this project.					
Nominated Representative This section must be signed by the person who oversees the legal affairs of the Applicant company or a nominated representative					

Title:

Name:

Signature

Position:

Date:

Please complete if Section 1.5 is applicable		
European Agent	This section mus	be signed by the person who oversees the legal affairs of the European Agent or a nominated representative
Title:	Name:	Position:
	Signature	Deter
		Date:

Next steps

Upon completion of the application form please return it to the primary certification body of your choice, i.e., the certification body you want to manage the project (*Refer to the selection in Section 1.2*). You will be contacted to acknowledge receipt.

The primary certification body will coordinate with the other certification bodies involved to create a common test programme (referred to as a 'test schedule') upon which, corresponding quotations are based.

You will receive quotations from each certification body (and where applicable associated testing laboratory) involved.

Each quotation covers a specific aspect of the testing/multiple certification that is to be performed by each certification body (and where applicable associated testing laboratory).

The testing and certification programme can begin when the Applicant confirms acceptance of each quotation.

