

3P application for certification according to CPR

Table 1: Scope of desired certification:

Certification with Inspection by Notified Body of FCP and regular maintenance	Attestation of conformity system	Levels in fire classes	Desired certification level made by an X
	NA	A_{ca}	NA*
	system 1+	B₁**	NA
	system 1+	B₂	NA
	system 1+	C_{ca}	NA
Certification on received products only (without inspection and without regular maintenance)			
	system 3	D_{ca}	Yes, see table 2
	system 3	E_{ca}	Yes, see table 3

*Level A_{ca} is not applicable for cables

** By selecting B₁, it is not possible to down grade to B₂,C_{ca} or D_{ca} in case of non-compliance.

Sponsor, name and addresses:

<i>Company Name</i>	
<i>Address</i>	
<i>Address</i>	
<i>Address</i>	

Production site for products under certification, name and addresses:

<i>Company Name</i>	
<i>Address</i>	
<i>Address</i>	
<i>Address</i>	

Contact information:

<i>Contact person Name</i>	
<i>Telephone</i>	
<i>Email</i>	



Table 2: Definition of products to be covered by the class Eca certification:

<i>Data, Communication or Power Cable</i>	<i>Jacket type LSHF, LSRHF or PVC</i>	<i>Diameter of cable in mm</i>
.....mm
	mm
	mm
	mm
	mm
	mm
	mm
	mm
	mm
	mm
	mm
	mm
	mm

Table 3: Definition of products to be covered by the class (B2ca to Dca) certification:

<i>Data, Communication or Power Cable</i>	<i>Jacket type LSHF, LSRHF or PVC</i>	<i>Diameter of cable in mm</i>
.....mm
	mm
	mm
	mm
	mm
	mm
	mm
	mm
	mm
	mm
	mm
	mm
	mm

" Sponsor " agrees by signing below with the following terms and conditions for the certification activity. "Sponsor" is hereafter called manufacture:

- a) Manufacture will without delay implement any changes in requirements for cables or production control system if and when they are communicated by 3P.

Note from 3P: A timescale for changes will in this case also be specified by 3P and will depend on any issue in question. Changes will be needed if requirements are modified in EN ISO/IEC 50575 or by official EU Group of Notified Bodies guidance papers.

- b) Manufacture assures unchanged passing of requirements for all future continued production, unless changes have been approved by 3P or a new certification has been issued.
- c) Manufacture confirms that the 3P assignment is a CPR certification as specified in table 1 and table 2.
- d) Manufacture will not make any misleading or incorrect statements about the granted certifications.
- e) Manufacture will stop marketing of any granted 3P certification if the certification is suspended, withdrawn, or terminated.

Note from 3P: A timescale for stop of marketing will in this case be specified by 3P and will depend on any issue in question.

- f) Manufacture will not distribute 3P certification documentation to third parties unless it is in complete form or has been initially approved by 3P.
- g) Manufacture will not issue marketing material that does not comply with the point e) statement or has been initially approved by 3P.
- h) Manufacture will comply with requirements for marks of conformity and information specified in CENELEC standard EN 50575.
- i) Manufacture will keep a record of complaints related to the activities and components of the present CPR certifications, and make this record available for 3P inspections.
- j) Manufacture will inform 3P without delay about changes in the organisation, raw material supply, cable designs and dimensions, key production or testing equipment, quality management system, FPC system and contact information.



- k) Manufacture will make all necessary arrangements for the 3P inspection visits, including availability for 3P inspection of (1) documentation and records, (2) access to storage, production and testing locations, (3) files for complaints and (4) education files for staff (5) Observers (DANAK, National Danish accreditations body) will be allowed to join 3P at the inspection visit. If subcontracting is used this must be initially informed to 3P, and access to subcontractors must be assured.

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Place **Date**

Title and name

Signature