



IEC System for Conformity Testing and Certification of Electrical Equipment CB Scheme

APPENDIX B (NCB only)

Legal differences

OD-CB 2007

MEMBER NATIONAL CERTIFICATION BODIES (NCBs) INFORMATION SHEETS			
<i>(Use additional forms as necessary.)</i>			
Product categories: CON, HOUS, INST, LITE, MEAS, MED, OFF, PROT SAFE, TOOL, TRON			
I	CB Scheme Procedure		
II	National Regulatory Requirements for Electrical Products within the Scope of IECEE CB Scheme	Yes	No
1	Is compliance with standards mandated by law?	✓	
	Comments: <i>Indirectly - through Provincial/Territorial legislation</i>		
2	Is product certification or approval mandatory?	✓	
	Comments: <i>Provincial/Territorial legislation</i>		
3	If approval is mandatory what aspects/characteristics of electrical products require mandatory certification or approval?		
	a) Electrical safety	✓	
	b) EMC		✓
	c) Hygiene – <i>N/A</i>		✓
	d) Energy Efficiency – <i>For some products (see list)</i>	✓	
	e) Ergonomics – <i>N/A</i>		✓
	f) Other (specify): - <i>Radiation – for medical radiation-emitting equipment</i>	✓	
	g) Telecom – Federal government certification for compatibility and network protection	✓	
4	Does the certification mark of the NCB provide regulatory recognition at National level?	✓	
	Comments: <i>Regulation is Provincial/Territorial matter - co-ordinated for uniform national implementation</i>		



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5	Does the NCB provide certification in all regulated areas listed in item 3 above?	✓ a),d) only	
6	If no, specify the body responsible for such certification or approval:		
	Electrical safety:		
	EMC:	Supplier's Declaration	
	Hygiene:		
	Energy Efficiency:		
	Other: Radiation	Bureau of Medical Devices, (Radiation Protection Branch) Health Canada (Fed. Gov't.)	
	Telecom	Industry Canada (Fed. Gov't.)	
7	Describe the conditions and the processes for obtaining each of the mandatory certifications or approvals in item 3 above, and include a flow chart for each process as necessary. Each description should address as a minimum the following:		
7a	Who can submit the application?	Foreign manufacturer directly/ foreign manufacturer via representative in a country/ commercial agent/ other NCB/ others	
7b	Language of the application:	English or French	
7c	Need to have a local representative to place a product on the country's market	Not required	
7d	Additional administrative requirements	samples for re-testing are taken from import lots/ requirements for product liability insurance 1. Registration of the manufacturer with the regulatory agency (<i>medical equipment only</i>) 2. <i>Mandatory periodic factory inspections.</i> 3. <i>Mandatory periodic re-testing requirements for some products only, e.g., thermal controls</i>	
7e	Can testing/ evaluation be carried out by NCBs in other countries?	Yes, in accordance with IECEE rules	
7f	Requirements for samples:	Samples are required with report	



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7g	Documentation required:	<i>CB Test Report, product description, photos, drawings</i>
7h	Language of markings, warnings and instructions	<i>English and French</i>
7i.	Time needed for processing (?)	<i>2 - 3 weeks</i>