

Agent and Consultant for China Market Access

CCC (China Compulsory Certification) Approval

The government of China has introduced this new Safety/EMC licensing system. The system, beginning from April 30, 2002 and fully implemented on August 1, 2003, requires manufacturers in 19 product groups to obtain the CCC certificate and affix CCC Mark before exporting or selling into the Chinese market. Products do not meet CCC requirements may be held at the border and subjected to other penalties by Chinese Customs Officials.

The Catalogue for CCC Approval

1	Electrical Wires and Cables	11	Telecommunication Terminal Equipment
2	Switches for Circuits, Installation Protective and Connection Devices	12	Motor Vehicles and Safety Parts
3	Low-voltage Electrical Apparatus	13	Motor Vehicle Tires
4	Small Power Motors	14	Safety Glasses
5	Electric Tools	15	Agricultural Machinery
6	Welding Machines	16	Latex Products
7	Household and Similar Appliances	17	Medical Devices
8	Audio and Video Apparatus	18	Fire-fighting Equipment
9	Information Technology Equipment	19	Detectors for Intruder Alarm Systems
10	Lighting Apparatus		

Only contact us for subcategories defined for each product group.

Requirements for CCC Approval

Step One: Determine Whether Your Products Require CCC Mark

Step Two: Self-assessment Prior To Application For CCC Certification

Step Three: Apply to the Authorized Certification Body(ACB)

Step Four: Type Testing by an Accredited Testing Lab

Step Five: On-site Factory Inspection and Audit.

Step Six: Evaluation of Results and Approval

Step Seven: Purchase or Print the CCC Mark Labels

Step Eight: Supervision After Certification

Step Nine: Appeal, complaint and dispute

SGS Germany GmbH, Raboisen 28, 20095 Hamburg

Vertriebsbüro Dortmund: heiko.schmidt@sgs.com, phone: +49 (0)231 98 22 95 45, fax: +49 (0)231 98 22 95 47

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What Can SGS do for you?

- Determine whether your product requires CCC
- Plan testing and application processes
- Work out solution when failing in type test
- Entirely track the whole test process and feed back to applicant in time
- Pre-test is an option
- Priority of Initial Plant Inspection (IPI) owing to good cooperation with CQC and government labs
- Priority of type test owing to good cooperation with CQC and government labs
- If required, IPI training can be provided
- Translate all documents, if available
- Provide up-to-date regulations and standard
- Apply for Exemption of CCC Mark
- Transform CCIB, CCEE to CCC/CQC

Requirements for CCC

- Copy of the corporation's business license of the manufacturer
- Valid factory inspection report (if available)
- Holden other CCC certification number (if available)
- Factory introduction and organization structure (if available)
- the filled application forms
- the list of critical parts
- Product photos, the inner configuration and the circuit diagram
- description of the difference between the different model/type of products in the same application unit
- user's manual in Chinese
- nameplate in Chinese
- other necessary documents

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MII Approval for Telecom Equipment

The Ministry of Information Industry (MII) of PR China has formally issued a government document called “The Measures for Administration of Interconnection of Public Telecom Networks” on December 31, 1998. The state applies a network access permit system to telecom terminals connected to public telecom networks, radio communication equipment and interconnection-related telecom equipment. Telecom equipment subject to the network access permit system shall acquire a network access license issued by the MII. Telecom equipment that has no network access license shall not be connected to public networks and not be sold in China.

The catalogue for NAL Approval

Telecommunication Terminal Equipment	Radio Telecommunication Equipment	Network Equipment
Fixed Telephone Terminal	Radio Base Station	Fiber Optics Equipment
Cordless Telephone Terminals	Microwave Telecommunication Equipment	Softswitch
Group Telephones	Satellite Earth Station	SS7 Signaling Equipment
Facsimile Machines		Intelligent Network (IN)
Modems (Including Card)		Synchronization Equipment
Programmer Control Exchange User Switch Machines		Network Access Equipment
Mobile User Terminal		Telephone Exchange/Switch
Wireless Paging Terminal		ATM Switch
ISDN Terminal Equipment		Integrated Switching System
Data Terminal Equipment (including card)		Router Equipment
Multimedia Terminal Equipment		IP Switch and Network Security Equipment
Other Telecom Terminal Equipment		Data Communication Equipment
		Paging Center Equipment

Requirements for NAL Approval

1. Application Form
2. Copy of the corporation's business license of the enterprise (those enterprise outside China can provide valid business license of their Chinese Branch or Agency)
3. Government documents with the permission of marketing inside China (for China-foreign joint venture or wholly owned foreign enterprises)
4. Background information (company profile, production capability & capacity, instruments, quality assurance and measurements after sale services etc.)
5. Agent authorization certification (for the foreign manufacturing enterprises consign the local agencies to apply for the permission of entry)
6. Company Guarantee (with the signature of the corporation's delegates)
7. Equipment briefing (function, performance, general design and principium framework etc.)
8. Equipment photos and the inner configuration & schematic diagrams
9. User operation manual
10. Testing reports and examples of commercial application (for the enterprises outside China)

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SRRC Approval for Radio Transmission Equipment

On January 1, 1996, a provision is implemented in accordance with “The Radio Regulations of the People’s Republic of China” and “Provisional Measures on the Management of Import of Mechanical and Electrical Products” with a view to strengthening the management of import of radio transmission equipment. Based on the provision, for any radio transmission equipment that are exported to China, the foreign businessmen involved should possess a “**Radio Transmission Equipment Type Approval Certificate**” issued by the Office of State Radio Regulatory Commission of the People’s Republic of China (hereinafter referred to as SRRC Office) and a **Type Approval Identifier** (hereinafter referred to as SRRC ID) should be marked on the equipment.

Requirements for SRRC Approval

- 1) Application form for Type Approval of Radio Communication Equipment
- 2) Copy of the corporation’s business license of the manufacturer
- 3) Agent authorization certification (if the applicant is an agent)
- 4) Copy of the certificate of Type Approval recognized by the government of the Country of origin
- 5) Background information
- 6) Photos with nameplate and model clearly shown
- 7) Instruction, technique manual, technology specification, function and the later test report or data
- 8) 2~5 sets of prototype / sample
- 9) Other required data

The catalogue for SRRC Approval

All types of equipments that transmit radio wave, include:

- Radio communication equipments
- Navigation equipments,
- Location equipments ,
- direction-finding equipments,
- Radar equipments,
- Remote control equipments,
- Telemetry equipments,
- Broadcasting and television equipments,

Not include the following that radiate electromagnetic wave:

- Industrial, scientific and medical (ISM) equipments,
- Electric transport system,
- High-voltage power line
- Other electrical appliances.

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SFDA Approval for Medical Device

A new rule, Regulations for the Supervision and Administration of Medical Devices of PRC, had gotten fully effective as of April 1, 2000. It requires:

When importing medical devices into China for the first time, the agent of the imported device should submit the instruction for use, quality standards, testing methods, other relevant information, product samples, and marketing authorization certificates issued by the manufacturing countries (regions). They also required to be inspected and be approved by the drug regulatory authority under the State Council, and received an import product registration certificate before applying for customs formalities.

Requirements for SRRC Approval

1. The certificate of the manufacturer's legal production qualification
2. The certificate of the applicant's qualification
3. The certificate recognized or approved by the government of the Country (Region) of Origin to authorize the products as medical devices to enter into the market of the country
4. The Standards of the products to be registered shall apply the provisions for the management of the medical devices standards
5. Operation Manual of the Products
6. The Type testing Report presented by the medical devices quality test agency recognized by the State Drug Administration within the recent one year
7. The clinical trial report of medical devices
8. The Product Quality Guarantee presented by the Manufacturer, to promise that the quality of the products registered and sold in China are unanimously the same as that of the identical products put into market in the Country (Region) of origin.
9. The certificate of commission for the After-Sale Service Agency designated in China, the letter of commitment and business certificate of the commissioned agency
10. The Self-declaration on the authenticity of the materials submitted." The Self-declaration on the authenticity of the materials submitted " shall be presented by the manufacturer

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CPA Approval for Measurement Equipment

According to “Metrology Law of the People’s Republic of China” and “Measures of the People’s Republic of China for the Supervision and Administration of Imported Instruments of Measurements”, anyone who wants to import the instruments of measurement (listed in the Catalogue of Model Examination of the People’s Republic of China for Imported Instruments of Measurement) must apply to the competent department who is in charge of measurement under the State Council for model approval.

After receiving applications, the competent department shall arrange for an authorized technical agency to conduct design appraisalment, which shall be conducted in accordance with the requirements laid down in the appraisalment outline. Its includes inspection of the exterior, assessment of measurement functions, safety, adaptability to environment, reliability and service life.

Requirements for CPA Approval

1. 2~5 sets of prototype/ sample
2. Model approval application of instruments of measurement
3. Copy of the business license of the applicant
4. Photos of the sample instruments
5. The assembly diagrams, structural drawings and circuit diagrams of the instruments
6. The technical manuals of the instruments
7. The documents of technical standards and the methods for examination
8. The testing reports on the sample instruments
9. Explanation of safety of the instruments
10. The operation instructions



The Catalogue for CPA Approval

1. Weighing apparatus (including scales);
2. Sensors;
3. Sound meters;
4. Surveying machines with three coordinates;
5. Surveying instruments for roughness of surface;
6. Geodesic instruments;
7. Calorimeters;
8. Flowmeters (including water meters and gas meters);
9. Pressure gauges (including sphygmomanometers);
10. Thermometers;
11. Digital voltmeters;
12. Field intensity meters;
13. Electrocardiographs and electroencephalographs;
14. Monitors for toxic gases, dust and water pollution;
15. Ionospheric radiation protectors;
16. Spectrometers (including meters for ultraviolet rays, infrared rays and visible lights);
17. Gaseous phase and liquid phase chromatographs;
18. Instruments for measuring temperature and moisture content.

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Manufacturer Licenses for Import Boiler and Pressure Vessel

According to “Regulation on Safety Supervision of Special Equipment” and “Supervision Administration Regulation for Manufacture of Boiler and Pressure Vessel”, the boiler and pressure vessel, which are manufactured and/or used inside the territory of P. R. China, shall be subjected to the manufacture licensing system and mandatory supervision inspection for the safety performance of the product. The products produced by the manufacturers that have not obtained “manufacture licenses” are not allowed to sell and use inside the P. R. China.

Requirements for AQSIQ Approval

1. Original Application Form
2. Introduction on the manufacturer (background and manufacture capacity).
3. Copy of the legal business License
4. Copies of qualified license or certificates that the manufacturer has obtained
5. Typical products and their related parameters and specifications
6. Drawing and design document
7. Quality manual of manufacturer
8. Other supplementary documents if deemed necessary

The Catalogue for AQSIQ Approval

- Boilers
 - Pressure-resistant Steam Boilers
 - Pressure-resistant Hot Water Boilers
 - Organic Fluid Heaters
- Pressure Vessels
 - Various types of gas cylinder for containing gas, liquefied gas and liquid with maximum working temperature no less than its standard boiling point, and the maximum working pressure no less than 0.1MPa (gauge pressure), and the product of pressure and volume no less than 2.5 MPa&lm3
 - Various types of gas cylinder containing gas, liquefied gas and liquid with standard boiling point less than 60;æ; the maximum working pressure no less than 0.2MPa (gauge pressure) and the product of pressure and volume is not less than 1.0 MPa; &lm3
- Medical Oxygen Cabins

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