Content

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| Content： | Chapter 1 General Principles<br><br>Article 1<br>These Regulations are established in accordance with paragraph 2 of Article 11 of the Standards Act.<br><br>Article 2<br>The dedicated authority in charge of standards may announce a list of products to which the designated national<br>standards items mentioned in Article 10 of the Standards Act apply as products within the scope of application<br>of the CNS Mark (hereinafter referred to as “the CNS Mark Product List”). The same shall apply when the<br>dedicated authority in charge of standards intends to withdraw products from the list.<br><br>Article 3<br>Products included in the CNS Mark Product List as mentioned in the previous Article may be allowed to use<br>the CNS Mark if they meet the following requirements:<br>1.Where the quality management (hereinafter referred to as “QM”) of the factory has been registered in a<br>quality management system as specified by the dedicated authority in charge of standards.<br>2.Where the products meet relevant national standards after inspection and testing. |

The dedicated authority in charge of standards shall specify and announce the quality management system referred
to in subparagraph 1 of the preceding paragraph.

Article 4
The pattern of CNS Mark is:
The dimensions of the above pattern shall be prescribed and announced by the dedicated authority in charge of
standards.
Article 4.PDF
Article 4.doc

Article 5
The CNS Mark as mentioned in the preceding Article, together with the certificate number, shall be affixed
prominently to the body of the products that are granted the use of the CNS Mark (hereinafter referred to as
"the CNS Mark products"). Where it is not possible to affix the CNS Mark to the body of the product, the Mark
shall be affixed to the packaging or containers of the product. For products in loose packaging, the CNS Mark
shall be displayed on the delivery notes.
If the CNS Mark is not affixed to the product in accordance with the provisions of the preceding paragraph,
the drawback shall be rectified within one month;
beginning on the following day, from the date that the notification issued by the dedicated authority in charge
of standards or commissioned by juridical persons or organizations (hereinafter referred to as "the third-party
certification agencies") is served. The specified period may be extended for one more month upon approval for
a single time only.

Article 6
The dedicated authority in charge of standards may commission the third-party certification agencies to conduct
conforming evaluations, certificate issuance (or replacement), market monitoring, and relevant management
procedures for the commission of CNS Mark products.

Article 7
The dedicated authority in charge of standards may recognize QM certification bodies (hereinafter referred to as
"recognized QM certification bodies") or testing laboratories to conduct relevant activities, such as the QM
assessments, product sampling and inspection.

Chapter 2 Applications

Article 8
Manufacturers applying for the use of CNS Mark (hereinafter referred to as "applicants") shall select the
classification of CNS Mark applicable for their products, and different manufacturing plants shall file applications separately.
Each product is limited to one application. However, if the product is subject to further classifications,
applications shall be made based on such classifications. One application is allowed for each classification.
Separate applications shall be made for different products produced and manufactured by the same plant. Where
the same product is produced and manufactured by different plants of the same company, separate applications
shall be made in respect of different plants.

Article 9
Before applying for the CNS Mark, an applicant shall first apply for QM certification with the dedicated
authority in charge of standards or a recognized QM certification body.
The dedicated authority in charge of standards or recognized certification body mentioned in the preceding
paragraph shall dispatch auditors to the applicant to perform assessments and prepare the report, which shall
be delivered to the applicant.

Article 10
Before applying for the CNS Mark, an applicant shall first apply for product inspection with the dedicated
authority in charge of standards or a recognized testing laboratory.
The dedicated authority in charge of standards or recognized testing laboratory mentioned in the preceding
paragraph shall dispatch inspectors to the applicant to sample products, which shall be inspected or witness
tested with compliance with related national standards. A report shall be prepared after inspection or testing
and delivered to the applicant.
With respect to the product in acquisition of other certification mark and registration of sustainable
effectiveness, as specified in the announcement of the dedicated authority in charge of standards, may be
exempted from inspection conforming to national standards with the latest test report of such certification
mark; items not exempted from inspection shall still be inspected as required.

Article 11
An applicant shall prepare and submit an application form together with the following documents and the
application fee to the dedicated authority in charge of standards or the third-party certification

agencies:
1.A copy of company or business registration certificate and a copy of factory registration certificate or
other equivalent certificate. Except if the manufacturer has filled in the uniform serial number of the company or business and the factory registration number in accordance with the application form, then this
requirement is no longer needed;
2.Basic information of the manufacturer;
3.Copies of registration certificates recognized by designated QM system, issued by the dedicated authority
in charge of standards or recognized QM certification bodies (recognized registration scope shall include
product items in the application); and
4.Copies of compliance reports of product inspection issued by the dedicated authority in charge of standards
or recognized testing laboratories within six months prior to the application.
When making the application mentioned in the preceding paragraph, a foreign applicant may authorize an agent
who has a business place in the Republic of China to act on his/her behalf by executing an authorization letter.
Where any of the documents submitted are in a foreign language, their Chinese translations shall also be
provided at the same time.
Where an application does not comply with the requirements of the preceding 3 paragraphs of Article 11, the
applicant shall rectify within one month; beginning on the following day, from the date that the notification
issued by the dedicated authority in charge of standards or the third-party certification agencies is served.
The application shall not be accepted if the rectification is not made within the specified time limit or noncompliance still exists after rectification.

Article 12
Where an application made in accordance with the previous article complies with the relevant provisions after
review, the dedicated authority in charge of standards or the third-party certification agencies shall approve
for use of the CNS Mark and issue the CNS Mark certificates, and may issue an English certificate based upon
application. Where the application does not comply with the relevant provisions, the application shall be
rejected with reasons stated.

Article 13
Under any of the following circumstances, manufacturers may apply for witness testing with reasons stated to
the dedicated authority in charge of standards,in addition to situations where witness testing is already

specified:
1.Huge and heavy products that are not easy for delivery;
2.Delicate and fragile products that are easily damaged;
3.Dangerous products that are easy to cause dangers;
4.The testing laboratory of the manufacturer complies with the requirements of CNS 17025 and is accredited by
a laboratory accreditation body of our country or of the same country as the applicant, which has signed the
Mutual Recognition Agreement developed by the International Laboratory Accreditation Cooperation, with the
items and the accreditation scope covered by the applicable national standards for products.
5.Other special circumstances approved by the dedicated authority in charge of standards.
The dedicated authority in charge of standards after accepting the applications may conduct on-site evaluations
to the plants or the laboratories, which the applicant assigns. An approval of witness testing will be granted
if any of the circumstances mentioned in the preceding paragraph is met and the applicant is equipped with
adequate testing facilities and capable of performing testing items in compliance with national standards.

Chapter 3 Administration

Article 14
The dedicated authority in charge of standards or the third-party certification agencies may conduct non-periodical
surveillance visits to the manufacturers who are allowed to use the CNS Mark (hereinafter referred to as "CNS Mark
registered manufacturers") each year.
When non-compliance with the requirements of previous paragraph is found during the surveillance, the drawback
shall be rectified within one month; beginning on the following day, from the date that the notification issued by
the dedicated authority in charge of standards or the third-party certification agencies is served. Upon expiration
of the one-month period, the dedicated authority in charge of standards or the third-party certification agencies
may conduct the surveillance again.

Article 15
In accordance with the quality management system designed by the dedicated authority in charge of standards for
CNS Mark registered manufacturers who are recognized to be non-compliance with the requirements of subparagraph 1 of
paragraph 1 of Article 3 during the surveillance of QM and are notified for rectification; the CNS Mark shall not be
allowed on products manufactured during the rectification period.

Article 16
The dedicated authority in charge of standards or the third-party certification agencies may take samples of CNS Mark
product from the market, construction sites or the plants of CNS Mark registered manufacturers to conduct product
inspection on an irregular basis. The inspection reports shall be prepared and provided to the CNS Mark registered
manufacturers.
If the preceding inspection is not conforming to national standards, the CNS Mark registered manufacturers shall
complete the rectification within one month; beginning on the following day, from the date that the notification
issued by the dedicated authority in charge of standards or the third-party certification agencies is served; and
apply to the dedicated authority in charge of standards or the third-party certification agencies for re-inspection.
The re-inspection mentioned in the preceding paragraph shall be conducted on new samples further selected on random
basis. But if only the labeling is not conforming to national standards, then a partial inspection may be implemented
only for the labeling.
Such product shall not use the CNS Mark, starting from the following day on which the notice mentioned in paragraph
2 is served, until the service of compliance reports of product re-inspection is served.
Where the CNS Mark products are proved to be non-compliance with national standards and such non-compliance is
verified by the dedicated authority in charge of standards or the third-party certification agencies, it shall be
conducted in accordance with provisions in paragraph 2 to paragraph 4.

Article 17
The CNS Mark products that are identified to be in compliance with national standards through inspections of
recognized testing laboratories in accordance with paragraph 1 of the previous article may be exempted from the
inspections as stipulated in the first paragraph of the previous article.
If the preceding inspection is not conforming to national standards, the CNS Mark registered manufacturers shall
complete the rectification within one month; beginning on the following day, from the date that the notification
issued by the dedicated authority in charge of standards or the third-party certification agencies is served; and
apply to the recognized testing laboratories for re-inspection.
The re-inspection mentioned in the preceding paragraph shall be conducted on new samples further selected on random
basis. But if only the labeling is not conforming to national standards, then a partial inspection may be implemented
only for the labeling.

Such product shall not use the CNS Mark, starting from the following day on which the notice mentioned in paragraph
2 is served, until the service of compliance reports of product re-inspection is served.

Article 17-1
The CNS mark products in acquisition of other certification mark and registration of sustainable effectiveness,
as specified in the announcement, may be exempted from inspection conforming to national standards as stipulated
in paragraph 1 of Article 16 or paragraph 1 of the preceding article with the test report on product sampling of
such certification mark within one year; items not exempted from inspection shall still be inspected as required.

Article 18
The factory surveillance, QM surveillance, product sampling inspection or request for relevant information conducted
by the dedicated authority in charge of standards or the third-party certification agencies, can not be evaded,
impeded, or denied by the CNS Mark registered manufacturers without due reasons.

Article 19
The CNS Mark registered manufacturers that suspend manufacturing of CNS Mark products shall file the reasons and
period of suspension with the dedicated authority in charge of standards or the third-party certification agencies
within three months beginning on the following day of the date of suspension.
The period of suspension mentioned in the preceding paragraph shall not exceed one year. An extension of six months
can be granted if there are valid reasons and; an approval is given by the dedicated authority in charge of
standards or the third-party certification agencies.
The CNS Mark shall not be applied to products manufactured during the suspension period mentioned in the preceding
2 paragraphs. Where a filing is made according to the provisions of paragraph 1, the CNS Mark may continue to be
applied to products that comply with national standards and are manufactured prior to the suspension period.
Upon the expiration of the suspension period mentioned in the second paragraph or early resumption, the CNS Mark
registered manufacturers may apply the CNS Mark to their products only after they have filed with the dedicated
authority in charge of standards or the third-party certification agencies that manufacturing of CNS Mark products
has been resumed.

Article 20
Manufacturing of CNS Mark products shall be deemed suspended if any of the following situations

was found:
1.The factory has no record of manufacturing CNS Mark products and quality management for the most recent year;
2.No sufficient product quantity is produced required for inspection can be obtained from the factory after
the previous sampling date for two times within three months.

Article 21
Where national standards applicable to the CNS Mark are revised or rescinded and resulted in the change of other
applicable national standards, the affected contents are available for evaluation. The CNS Mark registered
manufacturers shall complete rectification within six months; beginning on the following day,from the date that
the notification issued by the dedicated authority in charge of standards or the third-party certification agencies is served. An extension of six months can be granted if there is a rectification plan approved by the
dedicated authority in charge of standards or the third-party certification agencies.
The CNS Mark may continue to be applied on the products that comply with national standards and are manufactured
prior to the revision or rescission of national standards described in the preceding paragraph.
However, the products manufactured and produced during the extension period as stipulated in the proviso of
paragraph 1 shall not apply the CNS mark.
The CNS Mark registered manufacturers that have completed rectification earlier and have filed with the dedicated
authority in charge of standards or the third-party certification agencies; or have completed rectification
within the rectification period prescribed in paragraph 1, and conducted production in accordance with the
revised or newly applicable national standards may continue to use the CNS Mark.
After filing of the CNS Mark registered manufacturer as stipulated in the preceding paragraph or upon the
expiration of rectification period as stipulated in paragraph 1, the dedicated authority in charge of standards,
third-party certification agencies or recognized testing laboratories shall conduct product sampling inspection
in accordance with the revised or newly applicable national standards as stipulated in Article 16 and Article 17.

Article 22
Where changes are made to the content specified in the CNS Mark certificates, the CNS Mark registered manufacturers
shall apply to the dedicated authority in charge of standards or the third-party certification agencies for
issuance of replacement certificates by submitting the original certificates and relevant documents.
Where the changes involve relocation of plants; after registering the new plant site, the CNS Mark registered

manufacturers shall enclose copies of recognized QM registration certificates, assessment or surveillance
compliance report within one year, and product inspection compliance report within six months to apply for issuance
of replacement certificates.
The CNS Mark registered manufacturers who do not apply for issuance of replacement certificates in accordance
with the provisions of the preceding 2 paragraphs, shall rectify within one month; beginning on the following day,
from the date that the notification issued by the dedicated authority in charge of standards or the third-party
certification agencies is served.

Article 23
If the CNS Mark certificate is lost, damaged or destroyed, the CNS Mark registered manufacturers may apply to the
dedicated authority in charge of standards or the third-party certification agencies for re-issuance of certificates.

Article 24
Where the scope of registration of a CNS Mark registered manufacturer is influenced by the revocation or rescission
of the recognition status or alteration of scope of recognition of a recognized QM certification body, the
manufacturer shall submit a copy of the QM certificate and the assessment or surveillance report of that year to
apply for a change of recognized QM certification body within three months; beginning on the following day, from
the date that the notification issued by the dedicated authority in charge of standards or the third-party
certification agencies is served. If the application for such changes is not made, the manufacturer shall rectify
within one month beginning on the following day, from the date that the notification issued by the dedicated
authority in charge of standards or the third-party certification agencies is served.
The application for a change of registered QM certification body mentioned in the preceding paragraph shall be
approved if it complies with relevant provisions after review of the dedicated authority in charge of standards
or the third-party certification agencies; otherwise, it shall be rejected with reasons stated. The CNS Mark
registered manufacturers may apply for a second review within one month beginning on the following day, from the
date that the rejection notice is served.

Article 25
Where a CNS Mark registered manufacturer takes the initiative to apply for a change of recognized QM certification

bodies, it shall submit a copy of the QM certificate and the assessment or surveillance report of that year
issued by such recognized QM certification bodies to the dedicated authority in charge of standards or the
third-party certification agencies. Application for such changes will not be accepted during the rectification
period prescribed in Article 15.
Where a CNS Mark registered manufacturer voluntarily applies for change of product inspection agency, it shall
submit a copy of product sampling inspection compliance report of that year issued by such inspection agency and
fill in the application form to apply with the dedicated authority in charge of standards or third-party
certification agencies. Application cannot be accepted during the rectification period as stipulated in Article 16
and Article 17.
The application mentioned in the preceding 2 paragraphs shall be approved if it complies with relevant provisions
after review of the dedicated authority in charge of standards or the third-party certification agencies;
otherwise, it shall be rejected with reasons stated.

Article 26
Where the CNS Mark certificates are obtained through fraudulent means, the dedicated authority in charge of
standards or the third-party certification agencies shall revoke such certificates and request the certificates
to be returned by the manufacturers.
Within two months, starting from the following day on which the revocation notice as stipulated in the preceding
paragraph is served; manufacturer shall revoke the CNS Mark products prior to manufacturing and production, and
the CNS Mark logo applied on their packages, containers, delivery notes or other relevant product information.
Within one year, starting from the following day on which the CNS Mark revocation notice as stipulated in paragraph
1 is served; manufacturer shall not make a new application for the CNS Mark for the same products subjected to
previous revocation.

Article 27
Under any of the following circumstances, the dedicated authority in charge of standards or the third-party
certification agencies shall rescind the CNS Mark certificates and request the registered manufacturers to return
the certificates:
1.Where the required rectification involving marking has not been made within the specified time limit in
accordance with paragraph 2 of Article 5;
2.Where the CNS Mark fees are not paid;

3.Where the required rectification has not been made within the specified period or no application of product
inspection in accordance with paragraph 2 of Article 14, paragraph 2 of Article 16, paragraph 2 of Article 17,
paragraph 3 of Article 22, or paragraph 1 of Article 24;
4.Where violations of Article 15, paragraph 4 of Article 16, paragraph 4 of Article 17, paragraph 3 of Article
19, or paragraph 3 of Article 21 are made,regarding prohibition of the use of CNS Mark;
5.Where violation of Article 18 is made, regarding prohibition of evasion, impediment or denial;
6.Where report was not made in accordance with the provisions of paragraph 1 of Article 19 and any of the
circumstances prescribed in Article 20 is found,or report was made but production not resumed within the
period prescribed in paragraph 2 of Article 19;
7.Where the use of CNS Mark is not reported to the dedicated authority in charge of standards or the third-party
certification agencies according to paragraph 4 of Article 19;
8.Where the registration of quality management systems has been rescinded, or the QM registration scope has been
reduced, not covering the CNS Mark of the factory or product which is unable to acquire other QM registrations;
9.Where the application for the second review is not made in accordance with the provisions of paragraph 2 of
Article 24 or noncompliance still exists after the second review;
10.The CNS Mark registered manufacturers use the CNS Mark in the products without approval of using CNS Mark and
fail to make rectification within the time limit specified by the dedicated authority in charge of standards; or
the same manufacturer makes the same violation again within five years;
11.The same products of CNS Mark registered manufacturers are verified by the dedicated authority in charge of
standards or the third-party certification agencies as non-compliance with national standards for two times
within one year; or
12.Other violations of mandatory rule of law and of serious circumstance as certified by the dedicated authority
in charge of standards.

Article 28
Within two months starting from the following day on which the rescission notice as stipulated in the preceding
article is served, the CNS Mark registered manufacturers shall revoke the CNS Mark products prior to manufacturing
and production, and the CNS Mark logo applied on their packages, containers, delivery orders or other relevant
product information.
Within six months, following the day after the CNS Mark rescission notice has been served as mentioned in the

preceding article; the CNS Mark registered manufacturers shall not make a new application for the CNS Mark for
the same products subjected to previous rescission.

Article 29
Under any of the following circumstances, the dedicated authority in charge of standards or the third-party
certification agencies shall rescind the CNS Mark certificates and request that the certificates be returned:
1.Where the registered manufacturers apply for cancellation of the CNS Mark certificates;
2.Where the applicable national standards are rescinded;
3.Where the product is announced to be removed from the CNS Mark Product List;
4.Where the company or business registration certificate or factory registration certificate of the registered
manufacturer is revoked, rescinded or cancelled by relevant authorities;
5.Where the registered manufacturers are dissolved or close their business.
The CNS Mark products that comply with national standards and are manufactured prior to the date of rescission
mentioned in the preceding paragraph may continue to bear the CNS Mark.

Chapter 4 Recognition

Article 30
Regarding the qualification, condition, application procedures, assessment, term of recognition, revocation,
rescission and related administrative matters involved in the recognition of testing laboratories, the Regulations Governing Recognition of Designated Testing Laboratories for Commodity Inspection shall apply.
The scope and specific requirements of recognition for testing laboratories shall be announced by the dedicated
authority in charge of standards.

Article 31
Once receiving application of product inspection from CNS Mark registered manufacturers, the recognized testing
laboratories shall take samples from the manufacturing plant of the CNS Mark registered manufacturers to conduct
product inspection in accordance with national standards. The inspection reports shall be prepared and provided
to the CNS Mark registered manufacturers, as well as the dedicated authority in charge of standards or the
third-party certification agencies for references. If necessary, the dedicated authority in charge of standards
or the third-party certification agencies may hold on-site checks in testing laboratories.

Article 32
A QM certification body (hereinafter referred to as the applicant) that applies for recognition shall be qualified for any of the following criteria:

1.Being accredited by Taiwan Accreditation Foundation (hereinafter referred to as the Accreditation Foundation); or
2.Being accredited by an accreditation body that is located in Taiwan or the same country as the applicant and
that is a member of the Multilateral Recognition Arrangements (MLA) of the International Accreditation Forum,
Inc. (IAF) or the Pacific Accreditation Cooperation (PAC).
If there is no accreditation body in an applicant's country, or if the accreditation body located in the applicant's country has not yet become a member of the IAF MLA or PAC MLA, the accreditation can be obtained
from other IAF MLA or PAC MLA member accreditation bodies located in other countries. When the accreditation
body located in an applicant's country becomes a member of the IAF MLA or PAC MLA, an applicant is required
to obtain accreditation from this accreditation body within one year in order to retain recognition.

Article 33
The range of recognition as mentioned in the preceding Article sought by an applicant shall be limited to those
covered in the CNS Mark Product List or in the accreditation scope by accreditation bodies mentioned in the
subparagraphs of paragraph 1 of the previous Article.

Article 34
An applicant shall apply for recognition to the dedicated authority in charge of standards by submitting a
completed application form and attaching a copy of an accreditation certificate.
A foreign applicant may authorize an agent who has a domicile or business place within the territory of Taiwan,
the Republic of China, to submit the application.
If the application form and copy of accreditation certificate is presented in a language other than Chinese,
their Chinese translation shall be submitted at the same time.

Article 35
The application for recognition mentioned in the preceding Article shall be granted if it is deemed complying
with relevant provisions after review; otherwise, the application shall be rejected. The applicant may apply
for the second review within two months following the day of receiving the rejection notice.
The term of validity of the recognition mentioned in the preceding paragraph is the same as the term of validity
of the applicant's accreditation certificate. An applicant may apply for extension of the term of validity by
submitting a new application form within a period of three months prior to the expiration of the recognition.
The extended term of validity of the recognition is also the same as the term of validity of the applicant's

accreditation certificate every time.

Article 36
The dedicated authority in charge of standards may request an applicant or recognized QM certification body to
provide related documents or may dispatch its staff to conduct inspection of an applicant or certification body,
where necessary. The applicant or certification body shall not evade, impede or deny such requests for inspection.

Article 37
If the recognized QM certification body's scope of accreditation has been reduced and the scope of recognition
is affected; the recognized QM certification body shall apply with the dedicated authority in charge of standards
for changes to the scope of recognition within three months.

Article 38
The recognized QM certification body will be informed when their registered manufacturers have obtained the CNS
Mark certificates issued by the dedicated authority in charge of standards or the third-party certification agencies.
When performing surveillance visits to these manufacturers, the recognized QM certification body shall audit the
quality management systems of these manufacturers during annual surveillance visits to:
1.Ensure that the quality management of these organizations continues to comply with the requirements of specified
quality management systems;
2.Ensure that the manufacturers maintain the most up-to-date versions of regulations and national standards
applicable to the CNS Mark;
3.Ensure that the organizations establish adequate inspection plans for the CNS Mark products in accordance with
related inspection standards and devote adequate resources for the continuous compliance of their quality
management systems with the requirements.
A recognized QM certification body shall prepare surveillance audit reports and checklists during surveillance
visits and send the documents to the dedicated authority in charge of standards or the third-party certification
agencies within three months of a surveillance visit. If necessary, the dedicated authority in charge of standards
or the third-party certification agencies may dispatch staff to jointly conduct the surveillance visits.

Article 39
A recognized QM certification body shall inform the dedicated authority in charge of standards or the third-party
certification agencies of changes in the contents of the registration certificates or the invalidation of

QM
recognized registration certificates issued to its registered manufacturers that have been approved by the dedicated
authority in charge of standards or the third-party certification agencies to use the CNS Mark.

Article 40
The dedicated authority in charge of standards shall revoke all or part of the recognition granted to a QM
certification body under any of the following circumstances:
1.If the recognition is obtained through fraudulent means; or
2.If the accreditation certificates have been revoked.

Article 41
The dedicated authority in charge of standards shall rescind all or part of the recognition granted to a QM
certification body under any of the following circumstances:
1.If a QM certification body applies for cancellation of recognition;
2.If the accreditation has been rescinded or the accreditation certificates have been cancelled; or
3.If violation of Articles 36-39 is found.

Chapter 5 The Third-Party Certification Agencies

Article 42
The third-party certification agencies (hereinafter referred to as "the applicant") that apply for certification shall be qualified for any of the following criteria:
1.Governmental institutions, public or registered private colleges or higher educational institutions, or juridical persons of charity purposes;
2.Complete with CNS17065 system and acquire certificates of related fields from the Accreditation Foundation;
3.The testing laboratory has been recognized by the dedicated authority in charge of standards; and
4.Other criteria announced by the dedicated authority in charge of standards.
In addition to the above qualifications, the applicant shall be equipped with certification facilities, work site, personnel, management systems, information and knowledge about related certification scope
requirements for the product certification process.

Article 43
Applicants qualified by the requirements as stipulated in the previous article, have to prepare the following
documents and apply with the dedicated authority in charge of standards:
1.Certificates and proof of documents for qualifications as stipulated in the previous article;
2.Organizational chart and list of functions of organization;
3.Institutional layout and geographical map of location;
4.QM brochure;
5.Framework and list of QM documentation system; and
6.Other documents as designated by the dedicated authority in charge of standards.
Where the above listed documents submitted by the applicant are not in conformance with related, the applicant

shall complement or rectify within one month; beginning on the following day, from the date the notification is
served. Failure to complete the documents or rectification within the limited time, or still not conforming to
regulations, the application will not be further reviewed.

Article 44
Since the documental review as stipulated in the previous article is completed for conformation, the dedicated
authority in charge of standards shall conduct on-site checks. The application will be rejected when the on-site
checks are confirmed as not conforming to the relevant regulations.
Where the above-mentioned documental review and on-site checks are confirmed to conform to Article 42, the
dedicated authority in charge of standards may process the price negotiation and the commission contract for CNS
Mark product certification, and issue the proof of document.

Article 45
Where the results of on-site checks stipulated in the preceding article are confirmed as not conforming to related
regulations, the applicant may apply for re-inspection within two months; beginning on the following day, from the
date the notification is served. Not applying for re-inspection or confirming as non-conforming to the relevant
regulations will result in the application being rejected. The applicant may re-apply after three months beginning
on the following day, from the date the rejection notice is served.

Chapter 6 Supplementary Provisions

Article 46
For CNS Mark products, the dedicated authority in charge of standards shall announce the names of the CNS Mark
registered manufacturers, their plants, their registered CNS Mark product names and other relevant information,
and publish the information on the Standards Gazette. The same shall apply when the CNS Mark products are subject
to a revocation or rescission decision.

Article 47
These Regulations shall take effect from the date of promulgation.

Attachments： Article 4.pdf
Article 4.doc