

Certification Section	Muscat Quality Services SPC Conformity Certification Services Exhibit – Certification scheme	No.	MQS-CSC1a
		Rev. No.	00
		Date	27-10-2024

MQS Certification Scheme for Cosmetic and Personal Care Products

Issued by	Certification Manager		
Revision details			
Rev. No.	Issued on	Details of amendment / revision made	Reason for amendment
00	27/10/2024	First Issue	Due to implementation of ISO/IEC 17065
01	02/11/2025	Second Issue	Due to the external audit from GAC-NC

Prepared by	Approved By	Signature
Marwa AL-Ghafri	Quality Manager	

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1 Introduction

MUSCAT QUALITY SERVICES is leading Omani company specialized in providing product certification services which is authorized by Directorate General of Standardization and Metrology (DGSM). We have practical and scientific expertise and have extensive experiences and deep understanding of local, regional, and international standards related to quality and risk management. DGSM give the responsibilities to issue the Certificate of Conformity for several product including **cosmetics and personal care** in order to clear the product through Invest Easy and Bayan platform.

2 Scope of the Scheme, including the Type of Products covered and Type of Product Certification Scheme

This document describes a system for the Evaluation and Certification of **Cosmetics and personal care products**.

This Scheme is operated under the ownership of **Muscat Quality Services Conformity Certification** and is bound to be followed by **MUSCAT QUALITY SERVICES** and its Client. The Scheme is prepared based on the guidelines given in ISO/IEC 17067, Type 1a, to comply with the requirements of ISO/IEC 17065. The detail Scope of Accreditation is as below: -

Product / Process / System covered under the Scope	Requirement of DGSM	Normative Document(s) [Standard(s)/ Regulation(s)]
Cosmetic and personal care	Ref: DGSM-CPS-2	-GSO 1943:2024 -Guidelines for Product Classification (MOH-GD)

3 External References (Including the Reference Standards used in the Product Certification)

Management System Standards	
ISO/IEC 17000	Conformity Assessment – Vocabulary and General Principles
ISO/IEC 17065	Conformity Assessment – Requirements for bodies Certifying Products, Processes and Services
ISO/IEC 17067	Conformity Assessment – Fundamentals of Product Certification and Guidelines for Product Certification Schemes

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4 Terms and Definitions

For the purposes of this document, the definition is as below for better understanding:

1. **Client** – Organization or Person responsible to a Certification Body for ensuring that Certification Requirements, including Product Requirements, are fulfilled.
2. **Consultancy** – participation in the designing, manufacturing, installing, maintaining, or distributing of a Certified Product or a Product to be Certified, or the designing, implementing, operating, or maintaining of a Certified process or a process to be Certified, or the designing, implementing, providing, or maintaining of a Certified Service or a Service to be Certified.
3. **Evaluation** – combination of the Selection and Determination functions of Conformity Assessment Activities.
4. **Product** – result of a Process.
5. **Process** – set of Interrelated or Interacting Activities which transforms inputs into outputs.
6. **Service** – result of at least one activity necessarily performed at the interface between the Supplier and the Customer, which is generally intangible.
7. **Procedure** – specified way to carry out an activity or a process.
8. **Certification Requirement** – specified requirement, including Product requirements, that is fulfilled by the Client as a condition of establishing or maintaining Certification.
9. **Product Requirement** – requirement that relates directly to a Product, specified in standards or in other Normative Documents identified by the Certification Scheme.
10. **Certification Scheme** – Certification system related to specified Products, to which the same specified requirements, specific rules and Procedures apply.
11. **Scope of Certification** – identification of the Product(s), process(es) or service(s) for which the Certification is granted, the applicable Certification Scheme, and the Standard(s) and other Normative Document(s), including their date of publication, to which it is judged that the Product(s), Process(es) or Service(s) comply.
12. **Scheme Owner** – Person or Organization responsible for developing and maintaining a specific Certification Scheme.
13. **Certification Body** – third-party Conformity Assessment Body operating Certification Schemes.
14. **Surveillance** – systematic iteration of conformity assessment activities as a basis for maintaining the validity of the statement of conformity.
15. **Testing** – determination of one or more characteristics of an object of conformity assessment according to the procedure.
16. **Inspection** – examination of an object of conformity assessment and determination of its conformity with detailed requirements or, on the basis of professional judgement, with general requirements.
17. **Identification Test** – A simple check or measurement used to confirm that the current production batch matches the certified reference sample.
18. **Review** – consideration of the suitability, adequacy and effectiveness of selection and

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determination activities, and the results of these activities, with regard to fulfilment of specified requirements by an object of conformity assessment.

19. **Decision** – conclusion, based on the results of review that fulfilment of specified requirements has or has not been demonstrated.
20. **Attestation** – issue of a statement (Certificate) based on a decision that fulfilment of specified requirements has been demonstrated.
21. **Suspension** – temporary restriction of the Certificate of conformity by the body that issued the statement, for all or part of the specified scope of attestation.
22. **Withdrawal** – cancellation/revocation of the Certificate of conformity by the body that issued the statement.
23. **Conformity** – fulfilment of a requirement.
24. **Non-conformity** – non-fulfilment of a requirement.
25. **Impartiality** – presence of objectivity.
26. **Appeal** – request by the person or organization that provides, or that is, the object of conformity assessment to a conformity assessment body or an accreditation body for reconsideration by that body of a decision it has made relating to that object.
27. **Complaint**– expression of dissatisfaction, other than appeal, by any person or organization to a conformity assessment body or an accreditation body, relating to the activities of that body, where a response is expected.
28. **Corrective Action** – action to eliminate the cause of a nonconformity and to prevent recurrence.

5 Abbreviations

Abbreviation	Full Name
OS	Omani Standard
GSO	GCC Standardization Organization
ASTM	American Society for Testing and Materials
BIS	Bureau of Indian Standards
BS	British Standards
CA	The Concession Agreement entered into between the Concessionaire and the Government
CC	Central Computer
CCH	Central Clearing House
CENELEC	European Committee for Electro technical Standardization
IS	Indian Standard
ISA	Independent Safety Assessor
ISO	International Standards Organization
IT	Information Technology
JIS	Japanese Industrial Standards
DGSM	Directorate General of Standardization and Metrology
GAC	GCC Accreditation Center

6 Duties and Responsibilities

MUSCAT QUALITY SERVICES fulfills the requirements of ISO/IEC 17065 and will ensure that the Certification Scheme(s) for Product Certification, are controlled and operated so as to ensure, amongst other things, that they are impartial, and that decisions taken and implemented at all levels, including Management and Committees, are free from commercial or other pressures that may prevent the objective provision of Certification Services. The Certification Scheme is designed based on ISO/IEC 17067.

Ensure that information obtained during the Certification Process, from sources other than the Client, is not disclosed to anyone without the written consent of the individual concerned, except where the law of the land requires such information to be disclosed.

7 Objectives of Product Certification

The fundamental Objectives of Product Certification are: -

- to address the needs of Consumers, Users, and more generally, all Interested Parties by giving confidence regarding fulfillment of specified requirements.
- to demonstrate to the market that their Product has been attested to fulfill specified requirements by an impartial third-party body.

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Product Certification provides the following:

- Confidence for those with an interest in fulfilment of requirements.
- Sufficient value so that suppliers can effectively market Products.

8 Certification Scheme

The Certification Scheme is designed based on Type 1a as per the ISO/IEC 17067. The Scheme covers: -

a. Submit the application,

The Client shall submit the application as per the DGSM requirement Ref: MQS-AFCG Ver:00

b. Review

Examine the conformity requirements obtained during the determination stage to determine whether the product is in conformity or not.

c. Decision on Certification

Granting, Maintaining, Extending, Reducing, Suspending, and Withdrawing Certification.

d. Attestation, Licensing

- Issuing a Certificate of Conformity or other statement of Conformity (attestation).
- Granting the right to use Certificates or other statements of Conformity.

9 Sequence of Certification Cycle

The following is the sequence of the Certification Cycle, which is followed on a day-to-day basis: the client submits all the required files, such as the test report, labeling, and the manufacturer's certificate of composition, through either email or the MUSCAT QUALITY SERVICES software.

The Product Certification Assessor/Certification Manager checks all the documents to ensure that the tests comply with the regulation requirements to issue the Certificate of Conformity.

10 Application for Certification

The Client has to provide the following Information to MUSCAT QUALITY SERVICES:

- Legal name of the Applicant, Address of its Registered Office, Contact Details.
- Name and function of the person acting as contact to **MUSCAT QUALITY SERVICES** and authorized to sign on behalf of the applicant
- Business address and Contact Details.
- Manufacturer Details
- Designation of Product(s) for which Certification is requested:
 - Description of Product(s), including Technical Specifications,
 - Standard(s) and other Normative Document(s) to which Certification is requested: number, title, year of issue.
- Manufacture of Product(s):
 - Place(s) (physical address(es)) of Manufacture of the Product(s).

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11.0.2 Technical Evaluation

After approval of the application review form, the required documents are uploaded via e-mail or the official website of **MUSCAT QUALITY SERVICES**.

The Product Certification Assessor/Certification Manager performs a preliminary review of the documents and fills in the Technical Evaluation Report Ref: MQS-CCP Ver:00 Form before the conformity process, to ensure that all requirements are valid in order to save time and effort.:

Form 2: Checklist

Form 2: Checklist

11.0.3 Review

Once all evaluation activities are completed, the documents are reviewed by the **Reviewer/ Product Certification Assessor/Certification Manager** then the form is completed with the review decision, whether the application conforms or not, to be forwarded to the **Certification Manager/ CEO** for final decision.

The review is made by a Reviewer/ Product Certification Assessor/Certification Manager who has not been involved in the Evaluation Activities. The Review and Decision may be made by the same person or group of persons, depending upon the situation:

9. Summary of Product Evaluation:

Evaluation Findings:	
Non-Conformance of Products:	
Evidence of correction provide:	
Verification of Corrections:	

10. Evaluator's Recommendation:

☐ Product in Compliance with the requirements		
☐ Not complying or fails to comply with the requirement		
Evaluator's Name:	Signature:	Date:

11. Certification Review:

☐ Recommendation of approval of Product certification		
☐ Recommendation of rejection of Product certification		
Reviewer's Name:	Signature:	Date:

Form 3: Review checklist

11.0.4 Decision

When the outcome of the review is positive, a decision is made to grant Certification. When the outcome of the review is negative, a decision is made not to grant Certification. The Client is informed with the reasons for the negative decision. The decision is made by a **Certification Manager/CEO** who has not been involved in the Evaluation Activities. The Review and Decision may be made by the same person or group of persons, depending upon the situation. The decision of certification filled by **Certification Manager/CEO**:

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11.Certification Decision

☐ Grant Product Certification		
☐ Not to grant Product Certification		
Decision Maker name:	Signature:	Date:

Form 4: Decision Checklist

11.0.5 Attestation

Following the decision to grant Certification, **MUSCAT QUALITY SERVICES** issues a statement of Conformity.

Under this Scheme, the statement of Conformity is in the form of a Certificate.

In addition, the Certified Client may place the Scheme's Certification Mark on the Product, subject to a Licensing Agreement being entered into with **MUSCAT QUALITY SERVICES**.

11.1 Contents of Certificate of Conformity

The following information is provided in the Certificate of Conformity as a minimum;

- Certificate Number
- Name and address of **MUSCAT QUALITY SERVICES**
- Name and Address of Client (certificate holder)
- Statement of Conformity, including:
 - Name and Unique Designation (Trade Mark) of Product
 - Production Sites and Country of Origin
 - Applicable Standard(s) and other Normative Document(s) (including test report number)
- If applicable, reference to the Accreditation or Recognition Status of **MUSCAT QUALITY SERVICES** (Means status of ISO/IEC 17065 Accreditation)
- Date of Issue of Certificate
- Date of Expiry of Certificate
- Legally binding Signature(s) of Person(s) authorized to sign on behalf of **MUSCAT QUALITY SERVICES**.

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11.2 Licensing Use of Certificates and Marks of Conformity

11.2.1 General

The use of the Certificate and Mark of Conformity is controlled through the Certificate of Conformity issued by **MUSCAT QUALITY SERVICES** to each Organization which uses them on, or in conjunction with, Certified Products.

The Organization holding the Certificate of Conformity (referred to in this clause as the Licensee) may be different from the Client to which the Certificate was issued. Circumstances under which a different Organization might be involved include the following:

- The Client sub-contracts the manufacture of the Product, including the placing of the mark on the Product, to another Organization (the Manufacturer would need to be a Licensee)
- a Customer of the Client applies its own label, including the mark, to the Product under an agreement with the Client (the customer would need to be a Licensee)
- other cases.

In all cases, the Client ensures that **MUSCAT QUALITY SERVICES** has access to the Licensee's premises for Evaluation as agreed.

11.2.2 Mark of Conformity

Not currently applicable for DGSM.

11.3 Suspending or withdrawing a Certificate of Conformity

11.3.1 Suspension

The applicability of the Certificate of Conformity to a specific Product may be suspended for a limited period, for example, in the following cases:

- if it shows that the Corrective Action taken on the non-conformity issued is not effective or satisfactorily discharged.
- if a case of improper use of the Certificate (e.g., misleading publications or advertisements) is not solved by suitable retractions and appropriate Corrective Actions by the Client.
- if there has been any other contravention of the Product Certification Scheme or the Procedures of **MUSCAT QUALITY SERVICES**.

The Client is prohibited from identifying as certified any product that has been manufactured or placed on the market under a suspension of the certificate of conformity as applicable to that product.

A certificate of conformity may also be suspended after mutual agreement between **MUSCAT QUALITY SERVICES** and the Client for a limited period of non-production or for other reasons.

An official suspension of a certificate of conformity is confirmed by **MUSCAT QUALITY SERVICES** in a registered letter to the Client (or by equivalent means).

The Client may give notice of appeal, Ref: MQS-CAP, and **MUSCAT QUALITY SERVICES**, when considering the appeal, may or may not (depending on the nature of the case) decide to proceed with its decision to suspend the certificate of conformity.

MUSCAT QUALITY SERVICES indicates under which conditions the suspension would be removed, such for

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example, corrective action taken in accordance with Misuse of the Certificate of Conformity.

At the end of the suspension period, **MUSCAT QUALITY SERVICES** investigates whether the indicated conditions for re-instituting the certificate of conformity have been fulfilled.

On fulfillment of these conditions, the suspension is removed by notifying the Client.

11.3.2 Withdrawal

Apart from the Suspension of the Certificate of Conformity is withdrawn in the following cases:

- a. If the Client fails to comply with the due settlement of Financial Obligations.
- b. if there is any other contravention of the Certification Agreement.
- c. if inadequate measures are taken by the Client in the case of Suspension.

In the above cases, **MUSCAT QUALITY SERVICES** has the right to withdraw the Certificate of Conformity by informing the Client in writing concerning the specification of a time limit.

The Client may give notice of appeal Ref: MQS-CAP, and **MUSCAT QUALITY SERVICES**, when considering the appeal, may or may not (depending on the nature of the case) decide to proceed with its decision to withdraw the certificate of conformity.

Prior to withdrawal of a certificate of conformity, **MUSCAT QUALITY SERVICES** decides upon the consequences in relation to products covered by the certificate, whether a clearance of the stock of certified products is permissible within a short period of time. **MUSCAT QUALITY SERVICES** decides if other actions are required, including informing the clients.

Furthermore, the Certificate of Conformity may be withdrawn in the following cases:

- If the Client does not wish to maintain the Certificate of Conformity.
- If the Standard or Rules are changed and the Client either will not or cannot ensure Conformity with the New Requirements.
- If the Product is no longer made or the Certificate of Conformity goes out of business.
- On the grounds of other provisions specified in the certification Agreement.

11.4 Extending or Reducing the Scope of Certification

11.4.1 Extending the Scope of Certification

Purpose:

To include additional products, models, or manufacturing sites under the same certification.

Procedure:

1. The client submits a formal application for scope extension, indicating the new products or variants.
2. **MUSCAT QUALITY SERVICES** conducts a review and evaluation of the additional documentation, test reports, and conformity evidence.

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3. If the new scope uses the same product family and regulatory requirements, only partial evaluation may be performed.
4. If the product category or standard differs, a full evaluation (determination, review, decision) is carried out.
5. Upon satisfactory evaluation results, the Certification Manager shall approve the scope and issue an updated Certificate of Conformity reflecting the extended scope. Where applicable, a new certificate may be issued and the previous one withdrawn, depending on the specific conditions.

11.4.2 Reducing the Scope of Certification

Purpose:

To remove specific products or variants when the certified client no longer maintains compliance.

Procedure:

1. Reduction may occur when:
 - the client requests it;
 - a product or variants fail to meet requirements;
 - non-conformity persists after the corrective-action deadline; or
 - a production site closes.
2. The Certification Manager recommends the scope reduction based on evaluation findings.
3. A revised certificate shall be issued reflecting the reduced scope, and all corresponding records shall be updated in **MUSCAT QUALITY SERVICES** certification database. If applicable, a new certificate may be issued and the previous one withdrawn, depending on the specific conditions.

11.5 Refusing Certification

Purpose:

To reject an application when evidence of conformity is insufficient.

Procedure:

1. The Certification Reviewer identifies failure to meet regulatory or standard requirements, such as client participating in illegal activities, having a history of repeated non-compliances with certification/product requirements, or similar client-related issues.
2. A refusal notice is sent to the applicant stating the reasons and applicable clauses.
3. The applicant may reapply after addressing deficiencies.
4. All refusal decisions are documented and retained for impartiality review.

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11.6 Misuse of the Certificate of Conformity

MUSCAT QUALITY SERVICES takes action when unauthorized, incorrect, or misleading use of the certificates is found.

Incorrect references to the certification scheme or misleading use of certificates found in advertisements, catalogues, etc., are dealt with by suitable actions, which could include legal or corrective action or publication of the transgression.

12 Publicity by Client

The Client has the right to publish the fact that:

- a. an identified Product has been Certified
- b. the Client has been authorized to use a valid Certificate of Conformity

In every case, the Client takes sufficient care of its publications and advertising so that no confusion arises between Certified and Non-Certified Products.

The Client does not specify any function or make any claim or the like in user information that could lead purchasers to believe that performance of the Product or its use is covered by the Certification when in fact it is not.

In this example the Scheme requires **MUSCAT QUALITY SERVICES** to approve instruction books, manuals and other user information accompanying the Product.

13 Changes affecting Certification

13.1 Changes to Product Requirements

When a standard or another Normative Document that is part of the Certification Requirements is changed, there are a number of factors that have to be considered when fixing up the date on which the New Product Requirements of the changed document will come into force (effective date reflecting the transition period).

The effective date of obsolescence of a Standard or other Normative Document is communicated by **MUSCAT QUALITY SERVICES** to all applicable Clients to allow them adequate time to take appropriate action.

In those cases when the Standard development Organization responsible for the Standard or other Normative Document defines the transition period until which the superseded document is valid, this date defines the obsolescence of the superseded document unless otherwise stated by law or by the Scheme.

Further factors that are considered when choosing the effective date include, but are not necessarily restricted to, the following:

- a. Compliance with Regulations or Contractual Obligations
- b. The urgency of complying with revised Health, Safety, or Environmental Requirements
- c. The length of time and financial costs for retooling and manufacturing a Product complying with the revised requirements
- d. The extent of stock on hand and whether it can be reworked to meet the revised requirements

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- e. Avoidance of unintentional commercial advantage given to a particular manufacture or design
- f. Operational constraints of **MUSCAT QUALITY SERVICES**.

13.2 Changes to Scheme Requirements

MUSCAT QUALITY SERVICES advises their clients if necessary, of changes in the Scheme requirements, such as:

- a. Test and examination Procedures where these are not contained in the standards or other Normative documents that specify the Product Requirements.
- b. Conditions for licensing of the certification mark
- c. Qualification criteria and Procedures for Conformity Assessment bodies participating in the Scheme.

13.3 Changes by Client

The Client informs **MUSCAT QUALITY SERVICES** about any intended modification to the Product, which may affect the Conformity of the Product. **MUSCAT QUALITY SERVICES** determines whether the announced changes require another Initial Testing and Assessment or other further investigations. In such cases, the Client is not permitted to release Products under the Certificate resulting from such changes until **MUSCAT QUALITY SERVICES** has notified the Client accordingly.

A client wishing to extend the scope of certification to additional types or models of products, to the same specified requirements as the products for which a certification is already granted, applies to **MUSCAT QUALITY SERVICES** using an application form. In such cases, **MUSCAT QUALITY SERVICES** may decide to require or select test samples of the additional types of products to determine that they comply with the specified requirements. If the tests are successful, the scope of certification is extended.

If the Client wishes to apply the Certification to additional types of Products, but to different specified requirements, it will be necessary to perform only those parts of the Original Application Procedure which do not cover the new circumstances.

14 Confidentiality

MUSCAT QUALITY SERVICES is responsible for ensuring that the Confidentiality of Information is maintained by its Employees and those of its subcontractors concerning all information obtained as a result of their contacts with the Client; this applies also to information obtained at the application stage.

15 Impartiality

MUSCAT QUALITY SERVICES is responsible for ensuring that impartiality as per **MUSCAT QUALITY SERVICES** Impartiality policy

16 Product Liability

In this Scheme, all questions related to Product Liability are dealt with on the basis of the relevant Legal System(s).

17 Complaints and Appeals

The Client has a right to complain to **MUSCAT QUALITY SERVICES** about aspects of the service provided. The Client may also appeal to **MUSCAT QUALITY SERVICES** against its decisions on issuing, maintaining, extending, suspending and withdrawing Certification. All the Complaints and Appeal will be entertained as per the defined Procedure.

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18 Reference

18.1 “ISO/EC 17065 Conformity assessment – Requirements for bodies certifying products, processes and services”

18.2 MQS -PR Ver:00 Procedure for granting, for maintaining, for extending or reducing the scope of, for suspending, for withdrawing or for refusing certification”

19 formats / Exhibits

19.1 MQS – AF Ver:00: Application Form

19.2 MQS – CCP Ver:00: Checklist for Cosmetics and Personal Care Products

19.3 DGSM – CPS-2: Cosmetics and Personal care products Stage2

19.4 MQS – CA: Certification Agreement

19.5 MQS – CAP: Complaints and Appeals

قرار وزاري

رقم ٢٠٢١/١٩٩

باعتبار مواصفة قياسية خليجية مواصفة قياسية عمانية ملزمة

استنادا إلى المرسوم السلطاني رقم ٢٠٢٠/٩٧ بتعديل مسمى وزارة التجارة والصناعة إلى وزارة التجارة والصناعة وترويج الاستثمار وتحديد اختصاصاتها واعتماد هيكلها التنظيمي، وإلى القرار الوزاري رقم ٢٠١٩/١٢٨ باعتبار مواصفة قياسية خليجية مواصفة قياسية عمانية ملزمة، وبناء على ما تقتضيه المصلحة العامة.

تقرر

المادة الأولى

تعتبر المواصفة القياسية الخليجية رقم (١٩٤٣ GSO) الخاصة بمتطلبات السلامة في مستحضرات التجميل ومنتجات العناية الشخصية، مواصفة قياسية عمانية ملزمة.

المادة الثانية

تفرض غرامة إدارية لا تتجاوز (١٠٠٠) ألف ريال عماني على كل من يخالف أحكام هذا القرار، وتضاعف الغرامة في حال تكرار المخالفة.

المادة الثالثة

يلغى القرار الوزاري رقم ٢٠١٩/١٢٨ المشار إليه، كما يلغى كل ما يخالف هذا القرار، أو يتعارض مع أحكامه.

المادة الرابعة

ينشر هذا القرار في الجريدة الرسمية، ويعمل به بعد انقضاء (٦) ستة أشهر من تاريخ نشره.

صدر في: ١١ من جمادى الأولى ١٤٤٣هـ

الموافق: ١٦ من ديسمبر ٢٠٢١م

قيس بن محمد بن موسى اليوسف

وزير التجارة والصناعة وترويج الاستثمار



نوع الوثيقة	إجراءات الموافقة على مستحضرات التجميل ومنتجات العناية الشخصية
المرجع	د م - م ك - إ م - 1 - 2022
الإصدار	2023/01/8م
تاريخ النفاذ	2023/06/01م

الاعتماد

عماد بن خميس الشكيلي
مدير عام المواصفات والمقاييس
2023/01/08م




متابعة التعديلات

رقم الإصدار	تاريخ الاصدار	عدل من	وصف التعديل
02	2023/01/08م	-	-



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قائمة مستحضرات التجميل ومنتجات العناية الشخصية المحدثّة الخاضعة بدءاً من 1 يونيو 2023م

List of updates regulated cosmetics & personal care products since 1 June 2023

رمز النظام المنسق HS code	المنتجات Products
33049990	1. Leave on cosmetic products and personal care products which is intended to be applied on the skin (face , body, hands, feet , external genital organs act...)
	منتجات التجميل والعناية الشخصية (لغير الشطف) الموجهة للاستعمال على البشرة (الوجه ، الجسم ، اليد ، الأقدام ، الأعضاء التناسلية الخارجية)
	1. Creams with exceptions of cream for make-up creams such as foundation 2. Lotions \ emulsion 3. Butter \ balm 4. oil
	1. الكريم باستثناء الكريم الخاص بالمكياج مثل كريم الأساس 2. اللوشن/ المستحلب 3. الزبدة / الهلام 4. الزيوت
33049990	5. Products for external intimate hygiene
	5. منتجات نظافة المناطق الحساسة الخارجية
-	Products for care of the teeth and mouth
	منتجات العناية بالفم والأسنان
33061010	6. All types of Toothpaste
	6. معاجين الأسنان بكافة أنواعها
33069010	7. All types of mouthwash & breath freshener.
	7. منتجات غسل الفم وتعطير رائحته بكافة أنواعها
-	Care products intended for use on hair (head, face , bread)
	المنتجات الموجهة للاستعمال على الشعر (الرأس ، الوجه ، اللحية)
33059030	8. Different types of hair dyes (black, brown, other color)
	8. صبغات الشعر بأنواعها المختلفة (اللون الأسود واللون البني وأي لون آخر)
33052000	8. Hair straightener \ Relaxer
	9. منتجات فرد وتثبيت الشعر
33049990	9. Bleaching products
	10. منتجات التفتيح
33051000	10. Shampoo
	11. الشامبو
33079010	11. Chemical hair depilatories
	12. مزيلات الشعر الكيميائية
33042000	12. Eye Kohl
	13. كحل العين
33049920	13. All types of sunscreen products
	14. منتجات الوقاية من الشمس بكافة الأنواع
33072000	14. all types of deodorant & antiperspirant
	15. مزيلات ومضادات العرق بكافة الأنواع



General Condition	شروط عامة
1. The applicant company must be licensed to import and sell cosmetics and personal care products .	1. يجب أن تكون الشركة المتقدمة حاصلة على ترخيص يخولها بإستيراد وبيع مستحضرات التجميل ومنتجات العناية الشخصية .
2. The procedure in this document are applicable to the products listed in the table below.	2. تطبق الإجراءات المذكورة في هذه الوثيقة على كافة المنتجات الواردة في الجدول أدناه.
3. Only the products mentioned in this document are subject to conformity assessment procedures* *The products that requires obtaining approval from the General Directorate for Standardization and Metrology.	3. المنتجات المذكورة في هذه الوثيقة هي فقط الخاضعة لإجراءات التحقق من المطابقة* *المنتجات التي يتطلب إستيرادها وتسويقها الحصول على الموافقة من المديرية العامة للمواصفات والمقاييس
4.The company must obligate with the all requirements in Technical Regulation No. 1943 regarding “safety requirements in cosmetics and personal care products “	4. تلتزم الشركة بإستيفاء كافة المتطلبات المنصوص عليها في اللائحة الفنية رقم 2021/1943 الخاصة بمتطلبات السلامة في مستحضرات التجميل ومنتجات العناية الشخصية .
5. Ingredients used in cosmetics and personal care products must meet the restrictions in the latest update of the European regulation EC No. 1223/2009	6. يجب أن تستوفي المكونات المستعملة في مستحضرات التجميل ومنتجات العناية الشخصية القيود المفروضة وفقا لأخر تحديث للنظام الأوروبي (EC. No 1223/2009)
6. The company must submit the test report according. Appendix 1	7. تلتزم الشركة بتقديم تقرير الاختبار محتويا الفحوصات المطلوبة وفقا لنموذج المعد في هذه الوثيقة .



8. One certificate of conformity is accepted for the products of the same category* and same brand. *The products that similar to each other and have same function & characteristics.	8. قبول شهادة مطابقة واحدة للمنتجات التي تندرج تحت نفس الفئة * و نفس العلامة التجارية. *هي المنتجات التي تكون متشابهة ومماثلة لبعضها البعض وتشارك في الوظيفة والخصائص.
9. Products that contain medical claims or contain ingredients according to MOH gridlines require contact of General Directorate of Pharmacy and Drug Control at the Ministry of Health	9. المنتجات التي تحتوي على إدعاءات طبية وعلاجية أو تحتوي على مكونات بالنسب المذكورة في دليل الصحة (وفقا لآخر تحديث) يتطلب مراجعة المديرية العامة للصيدلة والرقابة الدوائية بوزارة الصحة.
10. Conformity certificates are issued for every product that is manufactured in factories in different countries	10. شهادة المطابقة تصدر لكل منتج تم تصنيعه في مصانع في دول مختلفة.
11. The Directorate with the market survey authorities, carry out the necessary activities that required to ensure suppliers meet the requirements in Technical Regulation No. 1943, including taking samples of products for testing and analysis in laboratories approved by the Directorate, with the supplier bearing the costs of testing and analyzing.	11. تقوم المديرية بالتنسيق مع سلطات مسح السوق بالأنشطة والتدابير اللازمة للتحقق من إستيفاء المزودين للمتطلبات المنصوص عليها في اللائحة الفنية رقم 1943 ومن ضمنها سحب عينات من المنتجات وإدخالها للفحص والتحليل في المختبرات المعتمدة من المديرية، مع تحمل المزود لتكاليف الفحص والإختبار.
12. The applicant shall responsible for the authenticity of the submitted documents.	12. يتحمل مقدم الطلب كافة المسؤولية حول صحة الوثائق المقدمة.
13. This document is subject to improvement in the future according to the circumstances.	13. هذه الوثيقة عرضة للتحديث مستقبلا وفقا للمستجدات



خطوات الحصول على الموافقة لمستحضرات التجميل ومنتجات العناية الشخصية

الخطوة الأولى : الحصول على شهادة المطابقة

(تتكرر عند إنتهاء صلاحية الوثائق)

- تصدر شهادة مطابقة واحدة للمنتجات من نفس الفئة ونفس العلامة التجارية من إحدى جهات منح شهادات المطابقة المحلية المسجلة فقط في مكتب الاعتماد بالمديرية العامة للمواصفات والمقاييس ويمكن التواصل معهم على الأرقام (24774754 & 24774855) أو على البريد الإلكتروني (omani.accredit@gmail.com)
- 1. تصدر شهادة المطابقة (السارية لمدة عام من تاريخ الأصدار أو بإنتهاء صلاحية تقرير الاختبار) وذلك بتقديم المستندات التالية:
 - 1.1 تقارير اختبار (سارية لمدة عامين من تاريخ الاصدار) من مختبرات **محلية** مسجلة بالمديرية حاصلة على الاعتماد في المواصفة الدولية (آيزو 17025) الخاصة بالمتطلبات العامة لكفاءة مختبرات الفحص والمعايرة في مجال مستحضرات التجميل ومنتجات العناية الشخصية ، مع إمكانية النظر في إستثناء بعض الحالات حسب تقييم المديرية.
 - 2.1 بطاقة البيانات الإيضاحية للمنتج وفقا لبند البيانات الإيضاحية للائحة الفنية رقم 1943 الخاصة بـ " متطلبات السلامة في مستحضرات التجميل ومنتجات العناية الشخصية "
 - 3.1 شهادة مختومة من الصانع تبين المكونات الداخلة لكل مستحضر مع نسبتها ووظيفتها.



الخطوة الثانية : التقديم عبر بوابة استثمر بسهولة

(تكرر عند انتهاء صلاحية الوثائق)

تقديم شهادة المطابقة السارية عبر بوابة استثمر بسهولة التابعة لوزارة التجارة والصناعة وترويج الاستثمار عبر الموقع الإلكتروني <https://www.business.gov.om/ieasy/wp/en>

الخطوة الثالثة : تقديم المستندات التالية عبر نظام النافذة الإلكترونية الواحدة (بيان) عبر الموقع (وتكرر عند إستيراد كل شحنة)

<https://www.customs.gov.om/dgcportal/ar/home/>

1. إقرار المستورد بالمطابقة (مفتاح لدى المديرية)
2. فاتورة الشراء مختومة وموقعة.
3. تصريح الموافقة على الأستيراد الحاصل عليه من بوابة استثمر بسهولة الخاص بتصاريح مستحضرات التجميل الخاضعة والواردة في فاتورة الشراء .

في حالة عدم وجود أحد الوثائق المطلوبة عند وصول الشحنة في المنافذ الحدودية :

- 1- يتقدم المستورد بطلب مكتوب مرفقا به نسخة من فاتورة الشراء للإفراج المشروط (المؤقت) عن الشحنة متضمنا رقم فاتورة الشراء إلى دائرة المطابقة بالمديرية العامة للمواصفات والمقاييس ودفع الرسوم المقررة و قدرها (50 ريال عماني) وإرفاقه في نظام بيان بعد مراجعة الدائرة له.
- 2- قيام المستورد بتقديم الوثائق الغير مقدمة.
- 3- الموافقة على الإفراج النهائي عن الشحنة على ضوء الوثائق المقدمة بالقبول أو الرفض (في حالة الرفض يتم إعادة تصدير الشحنة مباشرة إلى مصدرها أو إتلافها بالتنسيق مع الجهات المعنية مع تحمل المستورد التكاليف)
- 4- تسديد الرسوم المقررة لفك التحريز عن الشحنة برسوم وقدرها (50 ريال عماني)



Steps of cosmetics & Personal care products approval

Step 1: obtain a certificate of conformity

(Repeat when documents expire)

- One certificate of conformity for the products of the same category and the same brand is issued by one of the local certification bodies registered only in the accreditation office of the General Directorate of Standard (contact them at Tel. No. 24774855/24774854 or E- mail omani.accredit@tejarah.gov.om)

1- certificate of conformity (valid for one year from the date of issue or the date of expiry of the test report) is issued by submitting following documents:

- 1.1 test reports (valid for 2 years from issue date) issued by **local** laboratories which must be registered by DGSM , accredited and certified in the international standard (ISO 17025 general requirements for the competence of testing and calibration laboratories) with the scope in cosmetics and personal care products. With the possibility of considering the exception of some cases according to the evaluation of this Directorate.
- 1.2 product label according to the Gulf Standard (GSO 1943) Safety Requirements in Cosmetics and Personal Care Products.
- 1.3 List of product ingredients issued by manufacture showing ingredients percentage and function.



Step 2: Apply for Permit via Invest Easy:

(Repeat when documents expire)

submitting the certificate of conformity (valid for one year from issue date) via invest easy portal
<https://www.business.gov.om/ieasy/wp/en>

Step 3: Submit the following documents via Custom Electronic System (Bayan):

(Repeat when importing each shipment)

The following documents must be attached to the Bayan system for the final approval of import:

- 3.1. The importer's declaration of conformity (available in DGSM)
- 3.2. The commercial invoice
- 3.3. Permit Certificate which you get from Invest Easy Portal for the products mentioned in the commercial invoice.



In the absence of one of the required documents when the the shipment arrive at the border

- 1- The importer submits a letter, attached with a copy of the commercial invoice, for the temporary release of the shipment, including the invoice number, to the Conformity Department of the General Directorate of Standards and Metrology, and pays the fees **(50 OR)** then attach it to a statement system after the department reviews it.
- 2- The importer should submit the unsubmitted documents (the importer take the all responsibility in the event of disposing of the shipment or part of it before obtaining the approval of the Directorate)
- 3- Approval of the final release of the shipment, in case of the documents submitted are rejected will be re-exported or destroyed with coordination with the concerned authorities)
- 4- Pay the fees for final release of consignment with a fee of **(50 OR)**.



جدول يوضح أمثلة على فئات المنتجات

Example list of Product Category

فئة المنتجات Product Category	أمثلة على فئة المنتجات
كريمات الجلد Skin Cream	كريم مرطب ليلي ، كريم مرطب نهاري ، كريم مرطب للجسم ، كريم مرطب لليد ، كريم مرطب للقدم
	كريم تفتيح اليد ، كريم تفتيح الجسم ، كريم تفتيح القدم
	كريم ضد التجاعيد للوجه ، كريم ضد التجاعيد اليد... الخ
	أخرى
لوشن الجلد Skin Lotion	لوشن مرطب ليلي ، لوشن مرطب نهاري ، لوشن مرطب للجسم ، لوشن مرطب لليد ، لوشن مرطب للقدم
	أخرى
الزبدة Butter	زبدة للوجه ، زبدة لليد ، زبدة للقدم ، زبدة للجسم
	أخرى
الزيوت Oils	زيت للمساج ، زيت للوجه ، زيت الجسم الخ
	أخرى
منتجات نظافة المناطق الحساسة الخارجية Products for external intimate hygiene	غسل نسائي
	رغوة تنظيف نسائية
	أخرى
معاجين الأسنان Toothpaste	معجون أسنان ضد التسوس ، معجون للتبييض ، معجون للأسنان الحساسة الخ
	جل تنظيف الأسنان
	أخرى
منتجات غسل الفم وتعطير رائحته	غسل الفم التجميلي
	بخاخ الفم التجميلي



Others	أخرى	mouthwash & breath freshener.
All types of Black hair dyes	صبغة شعر أسود ودرجاته	صبغات الشعر Hair Dyes
All types of brown hair dyes	صبغة شعر بني ودرجاته	
Red , cupper, blond, white, purple hair dyes ect	صبغة شعر أحمر ونحاسي وأشقر و أبيض البنفسجي الخ....	
Others	أخرى	
Hair protein	بروتين الشعر	منتجات فرد وتثبيت الشعر Hair straightener & hair Relaxer
Hair keratin	كيراتين الشعر	
Hair straightener cream	كريم فرد الشعر	
Others	أخرى	
Hair bleaching cream	كريم تشقير الشعر	منتجات التشقير Bleaching products
Hair bleaching powder	بودرة تشقير الشعر	
Hair bleaching gel	جل تشقير الشعر	
Others	أخرى	
Shampoo for normal hair, Shampoo for oily hair , Shampoo for dry hair , Shampoo for colored hair , Shampoo for anti-dandruff	شامبو للشعر العادي ، شامبو للشعر الدهني ، شامبو للشعر الجاف ، شامبو للشعر المصبوغ ، شامبو مضاد للقشرة الخ	الشامبو Shampoo
Others	أخرى	مزيلات الشعر الكيميائية Chemical hair Depilatories
Hair depilatories cream	كريم إزالة الشعر	
Hair depilatories paste	معجون إزالة الشعر	
Hair depilatories foam	رغوة إزالة الشعر	
Others	أخرى	كحل العين Eye Kohl
Eye pencil	قلم العين	
Eye powder	بودرة العين	
Eye gel	جل العين	
Others	أخرى	منتجات الوقاية من الشمس sunscreen products
Sunscreen cream	كريم الوقاية من الشمس	
Sunscreen gel	جل الوقاية من الشمس	
Sunscreen spray	بخاخ الوقاية من الشمس	
Others	أخرى	مزيلات العرق ومضادات العرق Deodorants and antiperspirant
Deodorants/ antiperspirant spray	بخاخ مزيل/ مضاد للعرق	
Deodorants/ antiperspirant cream	كريم مزيل / مضاد للعرق	
Roll-on (gel) deodorant/antiperspirant	الكرة الدوارة (جل) مزيل/ مضاد العرق	
deodorant/antiperspirant stick	ستيك مزيل/ مضاد العرق	
Others	أخرى	



المراجع:

- 1- القرار الوزاري رقم 2021/199 باعتبار مواصفة قياسية خليجية (GSO1943) مواصفة قياسية عمانية ملزمة.
- 2- القرار الوزاري رقم 2021/190 بإصدار اللائحة الفنية لنظام المطابقة.
- 3- القرار الوزاري رقم 13/2012 بتحديد رسوم الخدمات والأختبارات والتحليل التي تؤديها المديرية العامة للمواصفات والمقاييس

للاستفسار التواصل

For contact

1- أرقام الهواتف - Tel. No

24774858

24774883

24774861

2- البريد الإلكتروني - Email

qcd.chemical@tejarah.gov.om



Product Test Report		
Test Report no. :		TR issued date :
Type of sample :		
Sample description		
Sampled by:	Sample condition & Temp. :	Sample Quantity:
Customer Name & address:		
Sample Date received :	Date of analysis start :	Date of Finished analysis
Production Date :	Batch No. :	
General Safety Tests for all Product Categories		
Test	Test Result	Unit
Physical Test		
1. Homogeneity		
2. Filthy/ Decomposed Substance		
Chemical Test		
1 pH Value		
2 formaldehyde		%
Heavy Metals Test		
1 Lead		ppm
2 Arsenic		
3 Cadmium		
4 Antimony		
5 Mercury		
Microbiological Tests		
1 Lard & Lard Derivative		CFU/ g
2 Total Aerobic mesophilic Count (Bacteria plus Yeast and Mould)		
3 Escherichia coli		
4 Staphylococcus aureus		
5 Pseudomonas aeruginosa		
6 Candida albicans		
Additional Tests		
Skin Brightening/ Whitening Products		
1 Hydroquinone		



2	Hydrogen Peroxide		%
Liquid Mouth Freshener			
1.	Safrole		%
Toothpaste			
1	Safrole		%
2.	Calcium monofluorophosphates		
3.	Magnesium fluorosilicate		
4.	Strontium chloride hexahydrate		
Hair Straighteners			
1	Calcium hydroxide		%
2	Lithium hydroxide		
Hair Shampoo			
1	quinine		%
2	Selenium disulphide (anti-dandruff shampoo)		
Chemical Depilatories			
1	Alkali sulphides		%
2	Alkaline earth sulphides		
Antiperspirant			
1	Aluminium zirconium chloride hydroxide		%
2	aluminium zirconium chloride		

تعميم بتطبيق المواصفة المحدثة ل 1943 External Inbox x

Summarise this email



qcd.chemical <qcd.chemical@tejarah.gov.om>

to info@sccom.com, me, contact@tawtheeqconformity.com, ali@tawkeed.org, Tadqeeq, Amr, Laila, Imad, Said, omani.accredit

21 Oct 2024, 10:52 ☆ ↩ ⋮

Translate to English X

الأفاضل بشركات منح شهادات المطابقة المحلية المحترمين
تحية طيبة وبعد،،

انطلاقاً من حرص وزارة التجارة والصناعة وترويج الإستثمار متمثلة في المديرية العامة للمواصفات والمقاييس على توفير منتجات آمنة وسليمة في السوق المحلي، نود إفادكم إلى أنه تم تحديث المواصفة القياسية (GSO1943/2021) الخاصة بمتطلبات السلامة في مستحضرات التجميل ومنتجات العناية الشخصية والصادرة بالقرار الوزاري 199/2021

وعليه ندعوكم بتطبيق اخر تحديث للمواصفة GSO1943/2024 وهي حالياً متوفرة في متجر المواصفات العمانية باللغتين العربية والإنجليزية.

وتفضلوا بقبول فائق الاحترام

المديرية العامة للمواصفات والمقاييس

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