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Welcome to Unilever’s Supplier Quality Manual. Over the years, we have published many handbooks, standards and manuals for our suppliers, but this is the first time we have provided a common, global guideline. We recognize that if we are to realize the potential of Unilever’s global scale, we have to speak with one voice to our partners around the world, rather than with many, sometimes contradictory voices.

This manual is your introduction to Unilever. You will learn about our vision to double the size of our business in a responsible and sustainable way, while still delighting our customers and consumers every time they use one of our products. You will get an overview of the processes, quality standards and information systems we use to manage quality across our supply base and will find links to take you to some of these systems.

Our success is intimately linked to your success......collaboration, simplification, transparency and clarity are the principles which must guide our partnership.

Thanks for your commitment

Marika Lindstrom
VP Procurement Operations
Global Markets

Kerry Easter
Chief Quality Officer
INTRODUCING UNILEVER

Unilever is one of the world’s leading marketers of fast-moving consumer goods. Our products are sold in over 190 countries and used by 2 billion consumers every day.

With more than 400 brands focused on health and wellbeing, no company touches so many lives in so many different ways.

Our portfolio ranges from nutritionally balanced foods to indulgent ice creams, affordable soaps, luxurious shampoos and everyday household care products. We produce world-leading brands including Lipton, Knorr, Dove, Axe, Hellmann’s and Omo, alongside trusted local names such as Blue Band, Pureit and Suave.
Unilever’s corporate mission – helping people to look good, feel good and get more out of life – shows how clearly our business understands 21st century consumers and their lives. The spirit of this mission has been a thread that has run throughout our history and quality has been an integral part of our business and our brands since its foundation.

We encourage you to visit our website www.unilever.com for more information.
Our Vision is to double the size of our business, whilst reducing our environmental footprint and increasing our positive social impact.

This vision statement places our consumers and the environment at the center of everything we do.
ABOUT THIS MANUAL

We clearly understand that we cannot deliver this Vision in isolation. We will only succeed if we effectively collaborate with all our business partners across our extended supply chain.

The purpose of this manual is to communicate our Quality work processes and operating framework and to share Unilever’s Quality requirements in a simple, clear and easy to use format.

We believe that in applying the processes and standards outlined in this manual, you will be able to continuously improve the quality of the materials you provide to Unilever.

SCOPE

Supplier in the context of this manual is identified as any third party contracted to supply materials to be used in the manufacture of Unilever products. Material in the context of this manual refers to raw materials, chemicals, ingredients or packaging components Unilever purchases from a supplier.

UPDATES

In the spirit of continuous improvement, we will evolve and improve this manual over time. You will be able to access the most up-to-date version of the manual through our SupplierNet portal.
UNILEVER’S STRATEGY
Our simple but compelling Vision is being brought to life across our supply chain in the form of our Supply Chain Blueprint. This entails building coherent and integrated strategies in all our product categories, business functions and country operations.

Within the area of Quality, three strategic thrusts have been launched across Unilever focusing on our customers and consumers, quality fundamentals, and our people and culture.
From a Procurement perspective, five strategic thrusts have been launched covering supplier capacity & capability, innovation & technology, responsible sourcing, quality & service, and value creation.

This manual represents the integration of our Quality and Procurement strategies and lays out the framework which we are implementing with all of our suppliers.
OUR SUPPLIER QUALITY FRAMEWORK

The Supplier Quality framework encompass how we select and contract with our suppliers, how we collaborate and integrate through information transparency, and how we collectively monitor and improve our performance.

This manual provides more details in each of the following areas:

- Supplier quality approval process
- Specification agreement
- Certificate of analysis
- Shelf life & date of manufacture
- Traceability, lot coding & records retention
- Storage, transportation & delivery requirements
- Performance monitoring & notification
- Performance measurement & improvement

Appendix 1: Calculating Remaining Shelf Life
Appendix 2: Information Systems
Our Supplier Quality Approval (SQA) process is applicable to all of Unilever’s materials suppliers. Each of your manufacturing facilities must successfully complete this qualification step and subsequently maintain approved status in order to supply to Unilever.

Our SQA process uses certification audits against globally recognized certification standards for suppliers of high and medium risk materials, or self-assessment questionnaires for suppliers of low risk materials. Certification audits are conducted by licensed audit companies, not by Unilever personnel. By successfully completing the certification audit or the self-assessment, you have demonstrated that you have the necessary Quality processes and practices in place to provide us with materials that are safe for consumers to use and are in compliance with regulatory requirements.

Our SQA process is managed through the Unilever Supplier Qualification System, USQS. USQS is administered by a third party contracted by Unilever. You will be asked to register each of your manufacturing facilities in USQS. Thereafter, all steps in the SQA process will be managed in the system.
Material Risk & SQA Assessment Process

Unilever assess a risk level for each material we purchase. Risk level can be assessed as high, medium or low. Risk dimensions include susceptibility to microbiological, chemical or physical contamination, quality performance factors and sensorial factors.

All of your manufacturing facilities must register in USQS and provide details of the material they supply to us. The risk level associated with the material then determines the type of audit your facility must successfully complete in order to be permitted to supply us.

Facilities that produce high or medium risk materials are treated the same in terms of audit certification and must obtain certification against a Unilever accepted external audit protocol. Facilities that produce low risk materials must complete and pass the Unilever Supplier Self Assessment audit (SSA).

If you have facilities that produce different materials with different risk levels, each facility will have to go through a different type of audit.

For a facility that produces more than one type of material for supply to Unilever, the overall risk and applicable audit process is dictated by the material risk and material type. For example, if a facility produces 2 low risk materials and 1 high risk material, the high risk material dictates the audit requirements for the facility.
On-site audits are conducted by 3rd party audit companies against globally recognized standards which are suitable for certifying your quality management systems, work processes and operating procedures.

A full list of audit standards that we accept can be found on USQS. Of this full list, there are audit standards that we prefer you to use. This preferred list is shown on the next page. In SupplierNet and USQS, a guidance document is available to help you select the appropriate audit standard for your facility.

If the audit protocol has the option for certification, this must be selected.

Some of the audit protocols result in a statement of compliance being issued at the end of the audit process instead of a certificate. This is fully accepted by us.

We do not accept ISO9001 certification.
**SQA AUDIT STANDARD INFORMATION FOR HIGH AND MEDIUM RISK MATERIALS**

**UNILEVER-PREFERRED AUDIT PROTOCOLS**

<table>
<thead>
<tr>
<th>Activity</th>
<th>Standards</th>
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- BRC Global Standard For Food Safety, v6  
- IFS Food v6  
- SQF Code |
| Manufacture of Raw Material for Home & Personal Care | - BRC Consumer Products  
- European Federation for Cosmetics Ingredients (EFCI) – GMP for Cosmetic Ingredients |
| Manufacture of Food Packaging Components       | - FSSC 22000  
- BRC IoP v4  
- ISO 22000 + PAS 223:2011  
- IFS PACsecure |
| Manufacture of Non-Food Packaging Components   | - BRC IoP v4 |

We periodically review the list of audit standards we are willing to accept. Please ensure that you only select audit standards from the information provided in USQS.
SQA audits must be performed by licensed audit companies that are certified by the audit standard owner. We will not accept any certification from an audit company that only holds a “Provisional License”. We recommend and prefer that you work with one of the following audit companies:

- Intertek
- DNV-GL
- Bureau Veritas
- Control Union
- SGS

Working with one of our preferred audit companies is advantageous because they are able to:
1) Conduct a wide range of SQA audits
2) Have auditors located across the world
3) Work directly within USQS.

Through USQS, you have the option to allow these companies to contact you directly to clarify any audit details such as auditing costs, schedules, audit formats and duration etc.

You are responsible for all costs related to the SQA audits, close-out of non-conformances and certification.
SQA AUDIT PROCESS FOR HIGH AND MEDIUM RISK MATERIALS

Your facility may already be certified and therefore approved to supply Unilever. Before you initiate the SQA process for the first time, check if your facility can demonstrate the following:

1. An current audit certificate against any of the audit standards accepted by Unilever with a scope that is applicable to the materials we procure
2. Provided by a licensed audit company
3. With at least 3 months remaining before expiration and a next certification audit booked

If so, you will be considered SQA approved once the certificate has been uploaded to USQS and validated by our service provider. If you cannot demonstrate this, you must select an audit standard and one of the preferred audit companies to perform the required audit or, initiate the re-approval process through USQS. The basic steps in the SQA process are shown on the next page.

Receipt of the certificate from the appropriate certification body is confirmation that your facility has met the requirements of the particular scheme your facility has been assessed against. The uploaded documentation will be reviewed by the USQS service provider to ensure that the audit matches one of the accepted audit protocols and the documentation has not expired. As soon as the USQS service provider confirms that the audit submitted is in accordance with our SQA requirements, you will be considered approved.

Pre-audits or pre-assessment reports, alone, do not demonstrate compliance to the SQA requirements and are therefore not accepted.

Maintaining SQA approval on an ongoing basis will be managed through USQS and our service providers.
Step 1: USQS Service Provider start process for audit booking based on risk and type of material purchased by Unilever

Step 2a: Supplier select audit protocol
Step 2b: Supplier select audit company

Step 3: Audit Company provide quotation and contract

Step 4: Supplier and Audit company to agree audit schedule

Step 5: Audit Company conduct audit

Step 6: Audit Company issue initial audit report (as per audit standard requirements)

Step 7: Supplier correct audit non conformances and write a corrective action plan (CAP)

Step 8: Audit Company validate CAP and issue certificate or statement of compliance (as per audit standard requirements)

Step 9: Supplier upload the certificate or statement of compliance to USQS

Step 10: USQS service provider change status in USQS based upon audit results

Step 11: USQS service provider escalate if audit failure

Note: Where your input or action is required, you will be contacted by USQS system generated e-mail and given weblinks to directly access the USQS.
If your facility produces low risk materials and does not have a current and applicable audit certificate, you must complete the Supplier Self-Assessment audit (SSA) instead of a full audit.

The SSA is available in the USQS database and is completed on-line. The SSA is a set of audit questions, relevant for the materials you supply to us and must be completed for each of your facilities.
Once your facility has been successfully approved, the facility must maintain an approved status in order to continue doing business with us. As the audit certification or self-assessment expiration date approaches, you will receive an email alert from USQS to initiate the re-approval process. Re-approval frequency is shown below.

### MATERIAL RISK

<table>
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<tr>
<th>HIGH AND MEDIUM</th>
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<tr>
<td>Certification audit</td>
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### GENERAL INFORMATION

All audit documentation and related supplier information will be stored in a secured database managed by a third party, under contract to Unilever. Only that third party and Unilever personnel are authorized to access the information in the database. Unilever-preferred audit companies do not have access to supplier information.

The risk level we assign to materials is subject to change; if any change results in you requiring approval to a new requirement (for example, external audit instead of self-assessment), you will be promptly notified.
SPECIFICATION AGREEMENT
Unilever’s specification forms the basis of our quality relationship with you for each material purchased. Specifications are developed independent of supplier and manufacturing facility. The specification defines all the properties of the material that you have to assure:

- consumer safety requirements
- product performance requirements
- visual & sensory requirements
- manufacturing & handling requirements
- storage, shelf life and transportation requirements

The specification identifies quality properties, target values and tolerances, and the required data accuracy and test methods to be used to measure each of these properties. The specification also defines the minimum required shelf life of the material. For certain materials, storage and transportation conditions (for example, temperature limits, humidity limits, etc.) and prohibited contact materials will also be defined in the specification.

Certain properties are defined as being “critical” for either consumer safety or quality performance reasons. You must confirm conformance for these critical properties for each production lot delivered to us in the form of a certificate of analysis, CoA. CoA requirements are covered in more detail later in this manual.
The material specification and testing methods must be agreed as part of the contracting process. An agreed specification implies that you understand all of our quality requirements for each respective property.

It is your responsibility to assure the quality of supplied materials and compliance to the agreed specification up to the point of receipt, or the point of use.

If changes are made to any quality properties, target values and tolerances, data accuracy or test methods, shelf life or storage and shipping conditions then a revised specification must be agreed.

Specification agreements will either be processed through SupplierNet or through our regular communication channels, for more information please refer to appendix 2.
CERTIFICATE OF ANALYSIS
A Certificate of Analysis (CoA) must be provided according to the agreed specification for each delivered lot. The CoA may be provided ahead of the delivery but must be available to our receiving location no later than the time of delivery. In some countries, we have the capability to receive electronic CoA data. You should check with your Unilever contacts to confirm if this is available.

The CoA must reference your unique manufacturing lot code for the delivered lot.

The CoA must be complete, legible and signed by your Quality representative (following local legislation where required). The CoA must reference our specification number and revision.

The CoA must record target values, tolerances and actual test results to the specified level of accuracy for each critical quality property identified in the agreed specification for the delivered lot. Each test result must reference the test method used and test method revision number.

Where you contract an external laboratory to perform analytical testing, the laboratory must be certified to a publicly available standard.

If a CoA is not provided with a delivery, we will generate a Supplier Non-conformance Report and reserve the right to reject the delivery.
SHELF LIFE AND DATE OF MANUFACTURE
SHELF LIFE, DATE OF MANUFACTURE & SAMPLES RETENTION

SHELF LIFE & DATE OF MANUFACTURE

You must provide the date of manufacture of the delivered lot for use in Unilever inventory control systems.

Our specifications define the minimum required shelf life of the materials. You must provide the corresponding maximum shelf life of the materials for each delivered lot.

Storage and stability data to support the maximum shelf life might be required during any qualification process.

Remaining shelf life of each delivered lot at the time of receipt at our facility must be at least 70% of minimum required shelf life. For an example calculation please refer to Appendix 1.

SAMPLES RETENTION

You must retain samples of the material lots delivered to us. Those samples must be retained for a time period equal to the maximum shelf life of your material or the time period denoted by local regulations (whichever is longer).
TRACEABILITY, LOT CODING & RECORDS’ RETENTION
You must have a traceability system in place that is capable of identifying unique lots of materials delivered to us which can be linked back to the uniquely identifiable lots of component parts used in your manufacturing process. You must ensure that your suppliers also maintain a traceability system with similar capabilities.

You must be capable of tracing materials one step downstream and one step upstream in your supply chain.

You must be able to complete a traceability mass balance (materials, ingredients and packaging components received, produced, lost or reworked, product released) within 4 hours of a request being received from a Unilever representative.

Where local legislation is more restrictive than our requirements, these local requirements must be applied.
LOT CODING & RECORDS’ RETENTION

LOT CODING

All material lots delivered to our manufacturing facilities must be uniquely coded in such a way that the tracing of individual lots as defined by you (for example, batch, shift, line or day) can be achieved.

RECORDS RETENTION

Manufacturing records for each material lot (for example, processing data, analytic data, materials consumed, etc.) must be retained for a time period equal to the maximum shelf life of your material or the time period denoted by local regulations (whichever is longer).
STORAGE, TRANSPORTATION AND DELIVERY REQUIREMENTS
You must have implemented storage and transportation systems to ensure that the safety, quality, and security of materials is maintained at all stages of manufacturing from receipt of raw materials through to delivery of products to Unilever.

If you use a third party warehousing facility to store raw materials, packaging materials, semi-finished or finished products, you must conduct periodic assessments to ensure that these requirements are met.

You must be capable of monitoring and controlling storage and transportation conditions according to the agreed specification, for example, temperature control, humidity control, etc.

Delivery vehicles must be checked for suitability, condition, cleanliness and integrity prior to loading. Loading procedures must be in place to ensure product will be stable and protected from damage for its transportation to Unilever.
You must conform to our inbound quality requirements for shipping containers (for example, bulk tankers, intermediate bulk containers, drums, big bags, bags, etc.).

Where non-dedicated bulk tankers and reusable containers are used, you must provide documentation with each delivery to verify the cleanliness.

You must ensure that during the shipping and storage of raw materials, they will be protected from cross-contamination from other materials, for example allergenic, chemical, physical and odor contamination, etc.

Unilever quality requirements for shipping containers are available to download from SupplierNet.
PERFORMANCE MONITORING & NOTIFICATION
Our receiving facilities will assess the conformance of each delivered lot against the agreed specification. This assessment can occur at the point of receipt or the point of use.

As part of the contracting and initial set-up process, you will have already been requested to provide an email address and contact name we will use in our systems. If any non-conformance is identified, a Supplier Non-Conformance Report, SNCR, will be generated by our receiving facility and you will be notified via an email to this address.

Examples of non-conformances are:
- Missing CoA
- Incomplete CoA
- Critical parameters out of specified tolerance
- Damage to the material (for example, punctured, crushed, wet etc.)
- Unacceptable condition of containers used to deliver the material (for example, odor contamination, molds, presence of insects, dirty, etc.)
- Unacceptable bar-coding (for example, cannot be read by scanner, illegible, missing etc.)
- Unacceptable labeling or printing (for example, poor positioning, no label, damage etc.)

Additionally, materials may be rejected if they are unsafe for our personnel to unload or handle.
SUPPLIER RESPONSE TO A SNCR NOTIFICATION

The SNCR will contain the following information:
- Description of the non-conformance
- Purchase order number for the delivery
- Date that the materials were received
- Quantity of non-conforming materials (units or weight)
- Total down time (in minutes) caused by the non-conforming materials
- Specific date that the supplier is expected to respond
- Name of the Unilever person who created the SNCR
- Where available, photographs or samples of the defect

You must analyse the SNCR data associated with the non-conforming lot, identify root cause(s), and implement actions to prevent the non-conformance from re-occurring.

You must then return the completed Corrective Action Template to the generic Unilever SNCR email address within 10 business days of receiving the SNCR notification. The email address is provided in the SNCR notification email sent by the Unilever location.
The Corrective Action Template summarizes key information and actions you have identified:

- Root cause(s) of the non-conformance
- Number of previous incidents of the same non-conformance
- Short term corrective actions you have already implemented
- Preventive measures you will be implementing to avoid re-occurrence
- Confirmation that you have identified and quarantined all materials from the same non-conforming lot

We will review the corrective action template response for completeness and will monitor subsequent deliveries for any re-occurrence of the non-conformance.
PERFORMANCE MEASUREMENT & IMPROVEMENT
We use the following key performance indicators to assess your performance at individual facility, regional and global level:
- Number of your facilities approved in accordance with the SQA program
- Number of agreed specifications
- Number of SNCRs

From these key measures of performance, more granular measures will be used to identify both local and cross-network improvement opportunities. These may include:
- Number of consumer safety SNCRs (for example, allergen contamination incidents, microbiological contamination incidents)
- Number of foreign matter SNCRs
- Your response time to provide corrective action information for SNCRs
- Number of SNCRs where corrective actions have not been effective in resolving the non-conformance
- Number of materials with all critical properties demonstrated to be under statistical control

All the information used to assess performance is based upon the data available to both of us and will be the basis of performance scorecards we will periodically review with you. You must be proactively analyzing performance data to drive continuous improvement.
We have a performance improvement culture which requires a relentless focus on loss elimination. Improvement goals are set with a zero-based vision and mindset – zero incidents, zero defects, zero losses. The quality information generated through the performance monitoring and notification process forms the basis for identifying improvement opportunities and loss elimination projects.

Overall performance improvement is driven in two ways:
- SNCR analysis and defect resolution at a local level as described in Performance Monitoring & Notification section
- Structured, visible and sponsored focused improvement projects (FI) designed to resolve high impact or chronic quality losses. Examples of high impact or chronic quality losses include:
  - SNCRs being repeatedly raised for the same non-conformance
  - consumer safety impact
  - consumer and customer complaints
  - market place incidents and recalls
  - operational productivity and cost impact

FI projects must be managed through a formal continuous improvement tool (for example, major kaizen, CAPDO Cycle, PDCA, FMEA, 6 sigma, etc). A good practice would be to nominate a FI project leader and ensure visibility of progress and performance improvement to your leadership team. We will periodically review the FI project performance with you at local, regional and potentially global level.
APPENDICES
APPENDIX 1: CALCULATING REMAINING SHELF LIFE ON DELIVERY

Remaining shelf life at the time of delivery to a Unilever facility must be at least 70% of the minimum required shelf life defined in our specification.

EXAMPLE CALCULATION

Minimum Required Shelf Life defined in Unilever specification: 180 days
Date of manufacture provided by supplier: 01/01/2014
Maximum Shelf Life provided by supplier: 365 days

Delivered to Unilever facility on 31/05/2014

Calculated expiration date = 01/01/2014 + 365 days = 31/12/2014

Remaining shelf life for material delivered on the 31st of May, 2014
= 31/12/2014 - 31/05/2014 = 214 days

Remaining shelf life/Minimum required shelf life = 214/180 = 119%
UNILEVER SUPPLIER QUALIFICATION SYSTEM, USQS

USQS is a global system Unilever uses to manage the qualification process for all of our suppliers. We have contracted a third party to administer the day-to-day operation of the system and the basic qualification workflow embedded in USQS.

All suppliers facilities should be registered in USQS and have access to the system to upload certification data and find more detailed documents which support the SQA process described in this manual.

Access can be requested by contacting Unilever Procurement.
SUPPLIERNET

SupplierNet is a global collaboration platform Unilever and its supply partners can use to share information and work on improvement activities. It provides news and information in addition to customized reports, enabling our suppliers to improve their operational and commercial performance across the markets they serve. All SNCR information and supplier responses are accessible via SupplierNet. Each supplier can only access its own data and is not able to access performance data for any other suppliers on SupplierNet.

Link and Unilever contact to request access: https://supplier.unileverservices.com/Pages/Home2013.aspx
Access can be requested by emailing snet.admin@unilever.com

SupplierNet includes reporting at the global and cluster level, searchable focus improvement projects, and material specification agreement. All suppliers are invited to regularly visit this website to review performance against the established KPI’s, and analyze the transactional data to identify the classification of events.
Suppliers should access the “Using SupplierNet” section in the training centre for additional information https://supplier.unileverservices.com/Training/Pages/TrainingHome.aspx

SupplierNet is also our online global portal for suppliers and provides access to additional information such as news, business alerts, policies and training.
# SUPPLIER QUALITY MANUAL CONTROL PAGE

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<td>Kerry Easter, Marika Lindstrom</td>
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## Revision History

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