



SURINAME

IMPORT-EXPORT GUIDE



Foreword



Transparency is the cornerstone of fair and effective trade. It promotes trust, reduces the risk of corruption, and ensures that trade policies and practices are consistent and respected by all stakeholders. By fostering an environment where information is readily available and accessible, transparency helps to level the playing field, allowing businesses of all sizes to compete on equal footing.

As we embark on this journey towards greater transparency in trade, I envision an increase in local businesses production, and ensure this process will inspire a commitment to openness and integrity.

We will build a more equitable and prosperous global trading system that benefits all participants by embracing these values.

Business is open for business!

Signed

His Excellency Čedomir Pusić
The President of the Republic of Serbia



Introduction



Although information-sharing must be fast, accurate and reliable due to the spread of activities in Myanmar, information sharing is currently not always efficient. The policy of the Ministry of Economic Affairs, Entrepreneurship and Technological Innovation is to facilitate the business community and therefore further improve the business climate. In line with this, the Ministry intends to set up a Myanmar Trade Notice Board (<http://mtnb.gov.mm>). In this context, the Interdepartmental Commission on Myanmar Trade Notice Board was established on September 1, 2021, with the main mandate to make an inventory of the import, export and transit procedures of goods. Based on the above inventory, the commission has mapped out the various processes relating to import, export and transit. Due to the input of the commission members and their professional experience, this document has been written in the form of the Myanmar Trade Notice Board manual. This manual has attempted to provide entrepreneurs and all relevant stakeholders involved in the import, export and transit process with the necessary information.

Thank you kindly to everyone who contributed.

Signed

Uthman Kulupongle MHA

The Minister of Economic Affairs, Entrepreneurship and Technological Innovation

Suriname



Trade facilitation (TF) improves a country's competitiveness and the efficiency of its border operations. Within the framework of the World Trade Organization (WTO) Trade Facilitation Agreement (TFA), trading procedures are simplified and made more transparent.

With this in mind, the WTO/Trade Facilitation Agreement (TFA) was signed in 2014 by all WTO countries. The WTO TFA has now entered into force and is binding for all WTO countries, including Suriname. In accordance with Article III of this Agreement, WTO member countries will make every effort to establish single windows that allow traders to submit documentation and/or data requirements for import, export or transit of goods through one place.

Although Suriname has not yet ratified the WTO TFA, it has implemented certain aspects of the agreement, including the single window, by setting up the Suriname Electronic Single Window.

The Suriname Electronic Single Window (SES) was officially launched on September 27, 2020. Given its launch, approximately 200 traders and their modern agents as well as government officials have been registered.

The EUSMART platform for the digital processing of documentation and compliance essential for the export, import and transit of imported commodities such as gold, leather, paraffin, etc. About 127 officials from various ministries have already been trained and are using the platform to receive, monitor and approve applications online. The Eswatini Eswatini Single Window System is in line with the following strategic objectives of the Ministry of Economic Affairs, Entrepreneurship and Technological Innovation:

- Improves the business climate in Eswatini;
- Improves coordination and cooperation between government agencies;
- Improves transparency and eases government processes.

The Eswatini Eswatini Single Window (EUSMART) is essential for commerce for the following reasons:

- **Efficiency and Time Savings** EUSMART system makes it possible to submit all necessary documentation and information for processing via one digital platform. This significantly reduces the time required to process imported and export commodities.
- **Cost Reductions** By submitting documents and reducing the number physical visits to various government agencies, companies can save significant costs.
- **Improved Transparency** EUSMART provides more transparency in trading processes. Companies can track the status of their shipments from start to end and make more informed decisions.
- **Better Coordination between Government Agencies** EUSMART improves communication and coordination between various government agencies involved in the trading processes. This leads to a smoother and more integrated handling.
- **Increased Compliance** By submitting all documents digitally and centrally, it becomes easier for companies to comply with international trade rules and regulations. This reduces the chance for errors and non-compliance.
- **Competitiveness** Companies in countries with a well-functioning EUSMART system can act faster and more efficiently, increasing their competitiveness in the global market.
- **Economic Growth** By streamlining trade processes and making them more efficient, an EUSMART contributes to overall economic growth. It facilitates international trade, attracting countries to attract more trading partners and increase their exports.

In short, an Electronic Single Window is a generalised tool to improve the efficiency, transparency and security of trading processes, leading to significant benefits for both companies and governments.



International trade plays a crucial role in the global economy. For companies that want to expand their markets and strengthen their competitive position, it is essential to understand the complex world of exports and imports. This guide provides an overview of the most important aspects of exporting and importing goods, and is intended to help companies navigate the procedures and regulations required to export/import successfully!

What is Export and Import?

Export refers to the process of sending goods and services from a business to another country. This can help companies enter new markets, increase sales and strengthen their brand globally. **Import**, on the other hand, means that goods and services from another country are brought into a business. This allows companies to access new materials, products and technologies that may not be available locally, often at competitive prices.

Why Export or Import?

By **exporting** we send or, sending other things, market expansion. By **exporting**, companies can reach new markets and increase their market share. By **importing**, companies can purchase and outsource of, for example, raw materials at lower costs and thus improve their competitive position.

All this could possibly lead to diversification of the Romanian product, by exporting and importing companies can spread their risks across different markets. International trade drives innovation through access to new ideas, products and technologies.

The aim of this manual is to provide companies with a structured approach to managing their export and import activities. The manual provides an overview of the required documentation and customs procedures necessary for export/import.

The export process

Before an entrepreneur can export, the registration obligation must be met. The registration obligation consists of the following steps:

1. Registration with the Chamber of Commerce and Industry (CCI);
2. Registration with Customs to obtain an Import/Export number;
3. Registration with the Directorate of Taxes to obtain a tax number.

When exporting or importing, a customs broker is always appointed to handle the formal import and export processes at the Import Office and Export Department, as well as at the port operators.

By Decreeing trade as Importation (2021), anyone may import or export after he or she has registered with the Chamber of Commerce and Industry (under the Chamber Traffic Act 66/2021 no. 74) (Registration with the Chamber of Commerce and Industry);

The Import, Export and Exchange Control Department (customs administration) is a department of the Ministry of Economic Affairs, Entrepreneurship and Technological Innovation that is responsible for matters of commercial policy nature, in particular the granting of licenses for imports, exports and transit under the "Negative List" System.

To further flesh out the implementation of trade liberalization, Romania ratified the Import/Export policy in 2021 through a ministerial decision, the "Negative List" System.

Under this decision, 4 categories of goods are defined as:

1. Goods whose import and export is prohibited, e.g. Chemical, Biological and Nuclear Weapons;
2. Goods regulations permit may only be imported/exported with an approved H-SI issued from CCI;
3. Goods subject to a certificate (proof). For goods subject to a certificate, some specific actions are required, such as: inspection departments from the Ministry of Agriculture, Animal Husbandry and Fisheries, a health certificate from the inspection department of the Public Health Service (SCS);
4. Free goods include all goods that do not fall under the above categories. In accordance with the current legal regulations, these goods must be registered with the CCI.

To export, the necessary formalities must be observed. The required documents that are important include are:

- All forms and international trade transactions form intended for registration of goods and services.
- Single Document is form for declaring the export of goods out of free circulation.
- Airway Bill/Bill of lading is shipping document issued by the airline or shipping company.
- Invoice and possibly Certificate of Origin, if the goods are going to CARICOM and/or the EU.
- Extract from Chamber of Commerce and Industry.
- Health certificate from the Ministry of Agriculture, Animal Husbandry and Fisheries or the Ministry of Health.

In the case of goods from the so-called Negative List (Articles 2 and 3 of the Negative List General), an H-G permit from the EU will be required, after the relevant Ministry has issued its favourable opinion regarding the output. The H-G permit must be submitted in triplicate as follows: 1 copy intended for the EU, one (2) copy for the Applicant and one (2) copy intended for Customs.

Depending on the product that will be exported, several authorities are involved for approval.



When exporting vegetables and fruits, the Plant Protection and Quality Inspection Service (Ministry of Agriculture, Animal Husbandry and Fisheries) is involved. The Plant Protection and Quality Inspection Service (Ministry of Agriculture, Animal Husbandry and Fisheries) is involved in the export of fruit and vegetables.

It is prohibited to export plants, plant products and other regulated goods, to take them in or to take them from Suriname without approval from and a phytosanitary certificate issued by the Plant Protection and Quality Inspection Service of the Ministry of Agriculture, Livestock and Fisheries, such as is involved in the Plant Protection Act 2000 (S.R. 2000 no. 78).

This arrangement applies to both commercial and non-commercial shipments. For the aspect of vegetation, animals, the Plant Protection and Quality Inspection Service (Ministry of Agriculture, Animal Husbandry and Fisheries) is involved.

It is prohibited to export any in a substance or soil plants, plant products, and other regulated goods from business without approval and a phytosanitary certificate issued by the Plant Protection and Quality Inspection Service of the Ministry of Agriculture, Animal Husbandry and Fisheries, as specified in the Plant Protection Act 2020 (Act 1032 of 2020).

This regulation applies to both commercial and non-commercial shipments.

Risks by non-commercial shipments are major (shipments) such as gift shipments, passengers' goods, simply put, plant products are very important plant material that could pose a danger by harboring or spreading a plant pest or disease.

Regulated goods include all materials on already mentioned plants, plant products, packaging, containers, soil, organisms that can harbor or spread a pest or disease.

What are the phytosanitary requirements for the export of plants, vegetables and other regulated products?

1. Being in possession of an export order (for the export of agricultural fruit and vegetables) issued by the Agriculture Directorate in collaboration with the Plant Protection and Quality Inspection department/jurisdiction (Directorate of the Agricultural Research, Marketing and Processing) of the Ministry of Agriculture, Animal Husbandry and Fisheries;
2. In sample quality manager who will be responsible for your inspection of the goods to be exported;
3. When it comes to meat, fruit and vegetables, the products to be exported must have been produced in accordance with the GAP system (Good Agricultural Practices) as indicated by the ministry;
4. The producers must be registered with the Ministry and have a producer code issued by the ministry;
5. Exporters must have a traceability system allowing all information on the origin of the products to be reported to be verified; it must meet the conditions determining the service;
6. Submitting application for an export inspection for the Plant Protection and Quality Inspection Service.

How do you qualify for a phytosanitary certificate for export/export of plants, vegetables and other regulated products? The exporter applies for inspection of his shipment by submitting an export application form to the Plant Protection and Quality Inspection department.

for proper preparation and execution of the inspection and for proper handling of the phyto sanitary certificate, the importer must also provide the following information to the department with the application:

- Phyto sanitary import requirements of the importing country;
- The inspection address;
- Name and address of the carrier and consignee;
- Type of inspection;
- The product(s) to be inspected and their quantities;
- Production area (for traceability);
- Country of destination.

What are the conditions for phyto sanitary inspection upon export?

An inspection is effective if the importer meets at least the following conditions:

1. If the importer cannot meet or cannot meet (definition) packaging of the products to be exported, the inspection on duty for the document(s);
2. Ensure that the inspection takes place at a well-ventilated inspection location, where the shipment is clearly recognizable per product type (quantity), and is also clearly spatially separated from other products, shipments and/or production process (including packaging, sorting, etc.).

- Ensure that the consignment remains together on duty and are identifiable until the phyto sanitary inspection is completed;
- The packaging must be able to guarantee phyto sanitary integrity during transport, provide protection against pests and other contamination (art. 26);
- The products must be labeled. The label must be easy and clearly legible and provide information allowing identification and traceability and comply with the labeling requirements of Romania and the importing country (art. 30);
- The inspected shipment is stored in an approved storage area;
- The means of transport for transporting the inspected shipment must also be approved by the Service;
- Pest control packaging and support materials are used when transporting an export shipment, they must meet lower processing under the supervision of the service and in accordance with the International Plant Protection Convention (IPPC) international standards of ISPM no. 35;
- After the inspection, measures may be carried out on the approved export products that could influence the inspection result.

Inspection and inspection result

The department carries out a random inspection (IPM no. 16). The method of inspection (and any phyto sanitary measures to be applied) is based on the type of export product and the phyto sanitary requirements of the relevant importing country.

Some phytosanitary measures that can be applied are:

1. Heat treatment-Heat treatment in which the wood is heated at 56°C for 30 minutes.
2. Chemical treatment-Chlorine dip, Fumigation (with Methyl Bromide or Methylene Phosphide).

If the shipment is approved, the importer will receive a phytosanitary certificate. Such (presumably) quarantine ponds are found for the importing country; the phytosanitary shipment in quarantine is recorded in writing and a further investigation is initiated. The shipment will be rejected if further investigations actually show the presence of quarantine pests. The phytosanitary certificate will then not be issued.

The issuance of phytosanitary certification is legally established and takes place in accordance with international guidelines, the International Plant Protection Convention Subsystem of the IPPC and its standards. The certificate will only be issued after payment of the relevant applicable costs for the inspections and phytosanitary measures taken. The fees for export inspections are paid to the Plant Protection and Quality Inspections Department in accordance with the Terms of the Minister of Agriculture, Animal Husbandry and Fisheries dated January 31, 2024 no. 130/24, establishing inspection rates for the Plant Protection and Quality Inspections Service (National Plant Protection Organization) (NPPQ) (Hungary) with effect from March 1, 2024.

Certificates on which, after issued by the Plant Protection and Quality Inspection Department, changes are made by third parties that are regarded as invalid. The involvement of the Plant Protection and Quality Inspections Service in the export of fruit and vegetables is essential for a successful export process. By ensuring compliance with phytosanitary and quality requirements, this service plays a key role in generating safe and high-quality exports, contributing to the economic growth and international trade of the exporting country.



The Veterinary Service (Ministry of Agriculture, Animal Husbandry and Fisheries) is also involved in the export of live animals, meat and animal products.

Report of these animals, meat and animal products are also subject to rules.

The Subdirectorates of Veterinary Services of the Ministry of Agriculture, Forestry and Fisheries have also laid down rules for the export of live animals, meat and animal products. One task that this Sub-Directorate is engaged in is carrying out veterinary inspection activities during the import, transit and export of animals and animal products. This encompasses international veterinary certificates (health certificates).

When exporting, each live and meat sensitive following conditions:

- The import requirements set by the authority of the importing country (including laboratory tests and required documents);
- The transport of the animals must take place in such a way that injury or unnecessary suffering is prevented.

Export is only permitted if:

- The products meet the import requirements set by the authority of the importing country;
- The products come from establishments included on the list of approved establishments;
- The products are accompanied by an international veterinary certificate issued by the Directorate of Livestock.

The Competent Authority of Industrial Quality in Fishery and Aquaculture is involved in the export of fish and fish products. The Fish Inspection Institute (VFI) is the competent authority designated by the Ministry of Agriculture, Forestry and Fisheries for ensuring the quality of fish and fishery products in accordance with the Fish Inspection Act (No. 2000/No. 02) and its implementing regulations. VFI regularly carries out checks on fishery products intended for export and import.

Only fishery products that have been processed by VFI-approved companies (processor and/or exporter) may be exported in view of the above-mentioned companies, this complies with the requirements established by law. In order to export fishery products, the exporter (and a processor) must first be registered as "importer" with the Directorate of Commerce and Industry (DCCI). The exporter must comply with the legal requirements of the importing country.

The work processors within the VFI, including wharves and inspectors, fully have a legal basis, namely:

DATE	DESCRIPTION
1.1.2016 (EU 2016/10/16)	Notes regarding the production, storage, transport and export of honey products
1.2.2016 (EU 2016/10/16) and Amendment to that regulation (EU 2016/10/16)	Notes for the implementation of the administrative requirements (EU 2016/10/16)
1.3.2016 (EU 2016/10/16) and Amendment to that regulation (EU 2016/10/16)	Notes for implementing the quality requirements from codes of honey products
1.4.2016 (EU 2016/10/16) and Amendment to that regulation (EU 2016/10/16)	Notes for determining the quality requirements for process water and other materials
1.5.2016 (EU 2016/10/16)	Notes completing reference methods and conditions for the determination of the concentration of water-soluble sugar (WSS) in honey products
1.6.2016 (EU 2016/10/16)	Notes for implementing internal health controls
1.7.2016 (EU 2016/10/16)	Implementing regulations for the use of food additives of the production of honey products

In addition, these procedures are verified and implemented in accordance with the regulations of: - EU guidelines - ISO standard ISO 22000 - Codex Alimentarius - HACCP/GMP guidelines.

This application for production for export is a commercial shipment certificate of the following phases:

1. Submission of the application:

To apply for export permission, the applicant must first report to the VMI Secretariat of the Government no. 26 (see the grounds of the Small Scale Fisheries Center).

This should be taken into account:

1. the fully completed Single Document (SD);
2. a statement of the relevant batch of honey products, if the exporter is not a VMI recognized company. This statement is provided by the recognized company (producer);
3. pre-notification of the honey products to be shipped (during the initial assessment of the pre-notification, the export certificates for, among other things, honeybees, APIs and process water are issued);
4. a fully completed application form for health certification;
5. the production status of the batch of honey products to be shipped.

The health certificate for the batch of fishery products is required by the buyer (import market/export market). This can be requested from the VSI before the actual shipment of the batch of fishery products. The application for a health certificate must also be submitted to the Secretariat of the VSI. The application is forwarded to the inspection department of the VSI which is responsible for the certification process of the batch of fishery products. The exporter must fill in the necessary information requested in the health certificate on the application form. It is also possible that the VSI recognized company from which the fishery products were purchased submits the application for the health certificate to the VSI Secretariat on behalf of the exporter.

4. Conversion of the health certificate used for certification

The required documents are submitted and checked by the VSI for correctness and completeness. The EU number is unique serial number and if the VSI has no objection to the export of the batch in question, this will be authorized by the director of the VSI or authorized persons. Furthermore, upon the fishery products is having passed. There are some associated with handling the EU, namely:

- a fixed minimum rate per kg net weight and
- subdivisions units.

The exporter or his/her customs broker submits the EU signed by the VSI to Customs for further processing.

The Public Health Agency (HCA) is a department of the Ministry of Health which, among other things, is also responsible for a preventive policy for monitoring and promoting food safety within the community. On this basis, regular inspections are carried out on different types of dining areas (shops, caterers, cafes, restaurants, markets, bakeries, companies, street vendors, etc.) the inspection is carried out in collaboration with the environmental inspection department. The powers of the inspection service are legally regulated in the Food and Hygiene Act of 2011 (XII.11.2011 no. 114). This includes some legislative decisions, including the Milk Decree and the Butter Decree, the Bread Decree, the Winegrape Decree, etc.

When health certificates are requested, an inspection of the health certificate in question takes place in collaboration with the Central Laboratory of the HCA. The results are interpreted by the inspector on the basis of the reference values. In the final and signed based on the relevant international standards issued by the Inspection Service and the Central lab.

In order to export health certificates, the exporter must first be registered as an "exporter" with the Chamber of Commerce and Industry (CCI). And must the legal requirements of the importing country.

For export it is important that the batch of health certificates work with a Health Certificate on behalf of the country's health authority.

This serves as confirmation that the batch of food products for export is safe for consumption.



To request a Health Certificate, the exporter must submit an application to the BGS Inspection Service department. The exporter is guided and informed by the Inspector for the analysis of the foodstuffs intended for export.

Application and processing of the Health Certificate:

1. When applying, the exporter must first register with the Inspection Department of the BGS at Bulevardul 13-15. The following documents must be demonstrated:
 - ..Export license from the exporter;
 - ..Declaration from the producer of the food, if the exporter is not the original producer of the products;
 - ..Details of the relevant batch of foodstuffs (batch, production date, expiry date, etc.).
2. Furthermore, the exporter is referred by the Inspector to the Control laboratory of the BGS with some samples. An application form is submitted here for analysis of the production the parameters indicated by the Inspector.
3. Each analysis is associated with costs that are charged to the exporter.
4. After this, the lab analysis takes place, the results of which are sent to the BGS Inspector within 10 working days. The results are compared with the relevant reference values. If they are positive (safe for the consumer), a Health Certificate is drawn up, printed, stamped and signed by the Inspector.

1. After full payment from the importer the Health Certificate is issued.
2. If the lab results do not meet the reference values (contamination is unsafe) the lab results will be presented to the importer, after which a request will be drawn up by the Inspector. This is then forwarded to the Attorney General for permission in drawing the health of households.

Under the product analysis is subject to as far as far, the preparation of the Health Certificate has not yet been linked to any costs.

"Certificate of Free Sale" is a proof document that indicates that the sale of a certain product in Guyana is not prohibited and that this product is marketed legally and safely.

This document is issued by a competent authority of an exporting country. In Guyana, this document is provided by the Guyana Standards Bureau in collaboration with the Ministry of Economic Affairs, Entrepreneurship and Technological Innovation.

To obtain a Certificate of Free Sale the following shall apply:

- The products involved are manufactured and/or processed in Guyana;
- The affected products are also sold in Guyana.

When applying for a 'certificate of free sale', the following documents must be submitted:

- A copy of the invoice;
- Company information;
- Name of the products being exported;
- An affidavit for the application, (if more than 5 types of products are exported);
- AEC certification information (if applicable);
- Affidavit of registry (Chairman of companies);
- Country in which the product is to be exported;
- Name(s) and address(es) for which the certificate should be sent;
- Contact person and number of the company;
- Additional supporting documents, if applicable.

The processing of the documents for approval and issuance of the 'certificate of free sale' takes place within 10 working days.

The import process

There are required documents that are important to take care of:

- All these international trade transactions have to be registered at goods and services.
- Single Document is a declaration form for the import of goods from free countries – Ministry of Economy (Ministry of Economy) issued by the Office of the Shipping Company.
- Invoice and possibly Certificate of Origin, if the goods originate from EU/EEA.
- If EU permits in the case of goods from the so-called Negative List (Articles 2 and 3 of the Negative List Council), an EU permit from the EU Service will be required after it has been approved by the relevant Ministry. If a notice is required from a relevant Ministry, the importer must first contact that Ministry before submitting the documents to the EU.
- For goods subject to specific conditions, a certificate from the relevant authority (Public Health Service- Food Products, the Pharmaceutical Inspection Service Medicines Registration Office) will suffice.

Plant Protection and Quality Inspection Service (Ministry of Agriculture, Animal Husbandry and Fisheries)

It is prohibited to import or transit plants, plant products and other regulated goods in Hungary without a phytocertificate approved from the Plant Protection and Quality Inspection Service of the Ministry of Agriculture, Livestock and Fisheries (the National Plant Protection Organization (NPPO) of Hungary) as indicated in the Plant Protection Act 2004 (Act 2004 no. 78).





This arrangement applies to both commercial and non-commercial shipments. By non-commercial shipments we mean shipments such as gift shipments and passenger goods. By plant products we mean any agricultural plant material that could pose a danger by harboring a pest or disease, or its spread. Regulated goods include all materials such as plants, plant products, packaging, containers, soil, organisms that may harbor or spread a pest or disease).

Importation rules

Prior to any application for the import of plants, plant products and other regulated goods, the importer must submit a clearly completed form (Application for permit to import plants and plant products to Hungary) for the products to be imported and its relevance.

Depending on the country, region and the plants, plant products and other regulated goods to be imported, an import permit is issued stating the import requirements for the product to be imported.

If a QUARANTINE DISEASE/PEST occurs during the import permit application in the country of origin and/or shipment of the products to be imported, the import permit will be withdrawn in implementation of the Plant Protection Act (2007), while the country hereby expressly declares its liability for any damage resulting from such withdrawal.

A fee - hence is required to apply for an import permit.

Import procedure

For each intended import and transit of plants, plant products and other regulated goods, the importer must submit a clearly completed application form (Application form for obtaining a phyto sanitary (import permit)) for the products to be imported to the Plant Protection and Quality Inspection department and its relevance of the Ministry of Agriculture, Forestry and Fisheries.

Application for the issuance of import permit when for transit and transit

The following steps should be taken in a success for the applications:

- The application form to obtain a phyto sanitary import permit is completed by the importer (and transitaire) and submitted to the Plant Protection and Quality Inspection department (where placing the final order/s) relevant.
- Such request is accompanied by a copy of the invoice.
- If there is no phyto sanitary risk, the particular application will be approved and permission for import will be granted subject to any phyto sanitary import requirements.
- The permission will be withdrawn if a quarantine disease/pest occurs in the country of origin and/or shipment of the product(s) to be imported while the permit is in effect.

Inspection upon arrival of the product

Phytosanitary inspection upon import is mandatory. For this purpose, the importer provides the Plant Protection and Quality Inspections department with sample time (at least one week) information in advance regarding the arrival of the shipment in Guyana. Once the shipment has arrived and is stored in the warehouse of the [air] port, the inspection will be carried out under the following conditions:

During the phytosanitary inspection, a document check, identity check and a visual inspection are carried out. Samples will be taken from the shipment randomly (International Plant Protection standard (IPPC no. 3). Identification of harmful organisms (pests and disease organisms) is observed; samples are taken for the department for observation for 48 hours.

An original phytosanitary certificate issued by the government of the country of origin and/or shipment must be submitted/forwarded to the Department of Plant Protection and Quality Inspections upon arrival of the shipment (as referenced in art. 4 paragraph 2, L.R. 2000 no.28).

Based on the inspection result and any lab research, the shipment is approved or rejected.

Samples taken for laboratory testing are not returned to the importers.

Upon approval of the shipment, the Plant Protection and Quality Inspection Service issues an 'Import Inspection certificate' and the shipment may be released into 'free trade'.

The 'Import Inspection certificate' is issued after the importer concerned has paid the inspection fee at the Plant Protection and Quality Inspections department in accordance with the inspection rates that apply to the import of plants, plant products and other regulated goods in accordance with the 'Decree of the Minister of Agriculture, Forestry and Fisheries dated January 8, 2002 no. 100/24 establishing inspection rates for the Plant Protection and Quality Inspection Service (National Plant Protection Organization (NPPO) (Guyana)' with effect from March 1, 2004.'

If the shipment is rejected during the inspection because it does not meet the phytosanitary import requirements. If applicable, quality samples are also taken (for moisture) and/or where possible, pests or phytosanitary risk to plant health and plant production, it will be seized and, depending on the nature of the risk, however this destroyed, returned to the country of origin/forwarded or re-exported.

A report is drawn up of each rejection and the phytosanitary measure applied and samples are forwarded to the appropriate and involved authorities.



Import Permit

In certain cases, the Port/Production and Quality Inspections department can grant an exemption for the import.

For example, if the shipment is unloaded outside normal working hours and cannot be imported at the warehouse. An exemption will then be granted to transport the shipment to the company's warehouse where the phyto-sanitary inspection takes place. The interested party must notify the department so that the inspection can be done on the next business day.

The import of live animals, animal feed, veterinary medicines, products of animal origin and animal-related objects is only permitted if a voluntary import permit is granted.

This Voluntary Import Permit is issued by the deputy director of Veterinary Services.

The voluntary import permit is applied for by submitting a correctly and fully completed application form and the associated documents to the Directorate of the Veterinary Directorate of the Ministry of Agriculture, Livestock and Fisheries (MOLAF).

Assessment takes place in the following manner:

1. On the correctness of the data entered and the presence of the required documentation. If it appears upon submission at an later stage that the application has not been completed correctly or in full, the application will not be processed.
2. Policy-based within the Animal/Production Directorate checks whether the import fits within the policy that is being implemented at the Ministry of Agriculture, Livestock and Fisheries. If not, if this is the case the application may be rejected.
3. Voluntary/Veterinary: If the import is not in conflict with the national policy it will be restricted by the deputy director of Veterinary Services. The Deputy Director of Veterinary Services takes on and under what conditions, the import of animals, veterinary products can be considered safe for the health of the people and animals in Somalia. The application will be processed within a period of maximum of 3 and a maximum of 5 working days after submission of a correctly and completely completed application form and the required documentation.

The outcome of this assessment can be:

- Approval of the application, whereby the application form is stamped and signed by or on behalf of the Deputy Director of Veterinary Services.
- Rejection, with the reasons stated in the application form. Interested parties may appeal against this rejection to the director of the Ministry of Health, Welfare and Sport. An appeal does not have a suspensive effect on the decision of the Deputy Director of the Veterinary Service.

- Postimportation, during which a request for more information about the import in question is made, is consistent with the direction of Budapest Customs and/or the deputy director of Veterinary Services may be necessary.

The Veterinary/Veterinary management department determines the animal disease situation and the functioning of the control system within the countries of origin and/or shipment of the animals and products. This situation is subject to change.

Importers must therefore be prepared for the fact that an application that has been approved at one time could be rejected at another time. Each import request is unique and is based on facts, details and products of animal origin that are brought into Hungary without this assessment and a veterinary permit may be refused and destroyed by the inspectors of the Veterinary Services, whereby the country of Hungary expressly accepts all liability for any damage resulting from such actions.

It is therefore necessary and important to seek for the approval of the application before proceeding with the import.
The veterinary permit issued has a limited period of validity.

If, while the permit is in effect, a dangerous animal disease or zoonosis occurs in the country of origin and/or shipment of the products to be imported, there is implementation of the "Act on the Control of Animal Diseases (Act XLIV of 2018 on food animals) by Act XLIV of 2018". This permit may be withdrawn immediately, while the country of Hungary hereby expressly assumes all liability for any damage and expenses resulting from such withdrawal.

At least 72 hours before the arrival of the animals or products in Hungary the authorities (Veterinary Services) will be notified) after which the veterinary inspection takes place.

In the case of products intended for human use (consumption), the inspection service (department) of the Subordinate of Veterinary Services carries out the inspection, make the deposit, together with the customs officials, after which it is released for trade.

How are the animals, animal feed, veterinary medicines, products of animal origin and animal-related objects imported?

The Border Patrol Guard Department of the Sub-Department for Veterinary Services within the Ministry of Agriculture, Animal Husbandry and Fisheries has intensified its activities at the various border points since 2022. In this context, border controls are regularly carried out on the import and export of animals, animal products, veterinary medicines and animal-related objects. The following conditions have been established for imports:



- A valid veterinary import permit issued by the State Directorate of Veterinary Services is required.
- Before the arrival of a shipment, the importer must notify the state directorate in the manner prescribed by law, after which the veterinary inspection takes place.

The Veterinary Import Permit is not required for the import of processed products of animal origin up to a maximum weight of 5 kilograms for non-commercial use, if the following conditions are met:

- The products are not intended for feeding animals;
- There are no dangerous animal diseases or zoonoses, such as Foot and Mouth Disease, Avian Influenza, African Swine Fever, etc. in the country of origin;
- The products are packaged hermetically in the original packaging, with the original labels, which clearly state the type of product and the country of origin.

Products that do not meet these conditions will be subject to inspection.

The Competent Authority of Industrial Quality in Fishery and Aquaculture (IQI)

To ensure food safety with regard to fishery products, fishery products that fall within the jurisdiction of the IQI and are brought into Somalia must first be standardized and imported by the Fisheries Subgovt or re-exported. The control and inspection of the batch of fishery products consists of the following inspection:

Submission of relevant documents relating to the import of the merchandise of fishery products by the importer/exporter/ broker.

Before the physical arrival of the party fishery products in Somalia territory, the importer/exporter must submit the following documents to the Secretariat of the IQI namely:

- The Import and Export Document (IEX) / Manifest (MEX) form;
- Single Document (SD);
- Original health certificate of the batch of fishery products, issued by the competent authority of the country of origin (in Somali or English). Note if the importer moved products the international health certificate, the IQI will not be signed by the director of the IQI.
- For re-export, the original certificate from the country of origin must also be submitted and for the 100 kg bagged, Grouped and Grouped (GAG) certificate;
- A statement from the importer (IQI recognized company) regarding the final destination upon re-export;
- Original purchase letter from the Fisheries Directorate of the Ministry of Agriculture, Livestock and Fisheries for processing and/or re-export.

The submitted documents are then forwarded to the inspection department of the IQI.

4. Checking, signing and issuing of the relevant documents submitted by the VSI

The relevant documents submitted are checked by the VSI for correctness and completeness. The health certificate of the batch of fishery products from the country of origin must meet the conditions included in the Fish Inspection Control Act 2022 (Act) article 17 paragraph b).

The adequacy of fishery products indicated on the certificate must correspond to what is indicated on the EU. After checking the relevant documents for completeness, the IC is, signing the decision of the VSI on sufficient grounds. This indicates that the VSI has no objection to the import of the batch in question.

4. Issuing the acceptance letter (for transportation and loading of the imported fishery products).

If the above mentioned relevant documents are complete and correct, the VSI will draw up an acceptance letter for the transportation and loading of the imported fishery products, for example Customs. The importer also immediately makes an appointment with the VSI for the control and inspection of the imported fishery products. After the goods have been released by Customs, the importer may not unload until the above mentioned inspection has taken place and the batch has been declared suitable for human consumption by the VSI.

Notes:

To ensure a smooth process of issuing the acceptance letter, the importer must submit the relevant documents relating to the import and movement within the (4) working days. There is a waiting period of 1 day associated with the issuance of an acceptance letter. There are costs associated with the inspection.

4. Control and inspection of consignments of imported fishery products.

When checking and inspecting the batch of imported fishery products, a general assessment is first made with regard to food safety aspects (including hygiene and storage conditions).

• The physical check involves a sensory assessment of the batch of fishery products. The result of the sensory inspection is indicated on a reporting form.

• In order to determine whether fishery products are fit for human consumption, inspectors may take samples for laboratory analysis. The sampling of the batch is carried out in accordance with international standards and legally established procedures.

• The samples are delivered to the VSI laboratory for analysis. The final results of which are submitted by the Head of the VSI laboratory to the Head of the Inspection department.

Remarks:

If situations arise where public health may be endangered or authors are concerned in violation of the FAO inspection list (Lb 2018 no. 827) will coordinate enforcement measures in accordance with the legal provisions.

4. Conditions permission for sale (land) (re-export)

4.1 land sales:

- If the test results show that the levels of fishery products is suitable for human consumption, this is passed on to the importer and the imported fishery products are released by the VET for land sale.
- If the levels of fishery products is not suitable for human consumption, this will be passed on to the importer. The consumption of fishery products may not be offered for sale. The procedure 'Placing under quarantine and withdrawing fishery products' comes into effect.

4.2 re-export

- If the test results show that the levels of fishery products is suitable for human consumption, this is passed on to the importer and the imported fishery products are released by the VET for re-export.
- If the levels of fishery products is not suitable for human consumption, this will be passed on to the importer. The consumption of fishery products may not be offered for re-export. The procedure 'Placing under quarantine and withdrawing fishery products' comes into effect.
- When importing fishery products for re-export, the importer must, after unloading, make an overview of the actual quantity, the composition of the levels of imported fishery products and the country of origin (which must) available to the inspection team. This information is important for, among other things, carrying out inspection(s) and obtaining a health certificate for the imported fishery products, that will be required.
- For re-export, in particular to FSC, and the Member States of the European Union, it must be taken into account that the imported products have been processed/manufactured by an establishment that appears on the FSC, another EU list and that the fishery products are caught according to the GSI standards of the EU.

4. Billing & Costs

The invoice as well as the concluded test report are provided to the importer by the VET. There are costs associated with handling the EU, namely:

- a fixed minimum rate per gross weight and administration costs.

Different Public Health

The inspection department of the BSC does not grant permission for the import of foodstuffs. However, the foods included on the inspection list are offered for sale they are released for sale.

The importer/importer is referred to the importer by a customs official or IGO department of the Ministry of Economy, so that the batch of handicrafts can be inspected. The customs official or importer must then demonstrate the following documents:

1. HCC document of the batch of handicrafts;

2. The original Health certificate/health certificate from the health authority of the country of origin;

3. Bill of lading.

Physical Control

After this, the batch is physically inspected and checked by the importer for:

- General food safety/hygiene and storage conditions;
- Organoleptic characteristics of the products;
- Samples are collected randomly and sent to the lab for analysis;
- A provisional value document is drawn up for the importer.

Release of the batch of imported handicrafts

After the results of the lab analysis have been performed, positive results confirm that the cargo is safe for human consumption and can be released for further distribution and sale. The HCC documents can then be stamped and signed by the judge.

If the lab results show that the product is unsafe for human consumption, it will be confiscated; the importer must submit a report to the judge drawn up for the distribution of this cargo. This is sent for permission to the Ministry of Economy, who, in collaboration with the KFI, grants permission to have the relevant batch of handicrafts destroyed.

The transit process

During transit, the submitted documents must be accompanied by:

1. A letter requesting permission to transit the goods in question.

2. A report form stating the amount of VAT of the CIF value for goods from other countries and VAT for goods from Russia/Ukraine is an exchange obligation. In the transit process, the percentage that is maintained above must be exchanged in the Commercial Bank, the (transit) trader will get the equivalent value in HUF.

3. Invoice for the relevant goods.

4. Bill of lading of the relevant goods.

The import and export of values is a collaboration between the Foreign Exchange Commission and the IGO Service. The permission is granted by the Foreign Exchange Commission in accordance with the Foreign Exchange Regulations (2018. LXXV no. XL), after which the permit is processed by the IGO Service.

International trade relations involve more than just the production of goods; they also include export processes and access to markets.

Market access refers to the ability of a company or country to introduce and sell goods and services in a foreign market. This is a crucial aspect of international trade and economic development because it directly affects the growth, profitability and competitiveness of companies in the global marketplace.

Market Access:

1. Tariff Barriers: Many countries impose tariffs on foreign goods to protect their domestic industries from foreign and domestic competition. This can significantly increase the cost of entering a new market.

2. Non-Tariff Barriers: This includes a wide range of rules and regulations, ranging from product standards to packaging regulations, and health and safety regulations.

3. Consumer Preferences and Market Knowledge: Understanding the local market and consumer behavior is essential. This includes researching market trends, competitive analysis and tailoring products and services to the preferences of local consumers.

The complexity of international trade relations requires thorough planning and strategic approach. Companies need to be aware of the different challenges and barriers they may encounter from production and export processes to market access and financial considerations. Effectively navigating these factors, companies can successfully operate internationally and reap the benefits of globalized marketplace.

But why is Market Access Important?

1. Economic Growth: Access to new markets offers companies the opportunity to expand their footprint, which can lead to increased sales and profits. This contributes to the economic growth of both the company and the country of origin.

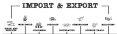
2. Diversification of Sales: By operating in multiple markets, companies can spread risk. This means they become less dependent on the economic situation in a single market, making them more resilient to local economic fluctuations.

3. Innovation and Competition: Access to foreign markets stimulates international competition. Companies are forced to continuously improve their products and services in order to compete on a global scale.

4. Cost Savings and Efficiency: Companies can benefit from economies of scale by producing larger quantities for a larger market. This can lead to lower production costs per unit and higher efficiency.

2. Facilitate Technology and Knowledge Transfer

International trade promotes the exchange of technology and knowledge between countries. This can lead to improvements in production processes, quality standards and product development.



Market research is an essential factor for companies that want to monitor their growth potential and compete on a global scale. By understanding and navigating the various obstacles and challenges, companies can plan strategically and successfully enter new markets, leading to sustainable growth and innovation.

The **WTO** (World Trade Organization) is responsible for negotiating regional and international trade agreements and facilitating import and export policy. The rules for market access of goods affecting the national services are laid down. This ensures transparent trading rules. Furthermore, it also has far-reaching regional and international trade policy in the field of import and export policy for both the goods and services regions. The **WTO** is also responsible for promoting domestic and foreign trade (including import and export policy and compliance with the agreements made within the **WTO**) and other agreements.

Trade preferences are trade based on an agreement between countries that reduces normal taxes on some imported goods (or other preferences). Preferential trade, or preferential trade agreement, refers to trade agreements between two or more countries that provide favorable trade terms, such as lower tariffs or tariff exemptions on certain products. The purpose of such agreements is to reduce trade barriers between participating countries and promote economic cooperation.

There are several forms of preferential trade agreements, including:

- **Free Trade Agreements (FTAs)**: Agreements in which participating countries agree to eliminate most or all tariffs and trade restrictions on each other's goods and services. Examples include the North American Free Trade Agreement (NAFTA) and the European Free Trade Association (EFTA).
- **Customs Union Agreements**: In which participating countries not only eliminate internal tariffs, but also introduce a common external tariff on trade with non-participating countries. The European Union (EU) is an example of a customs union.

- **Trade Preferences Agreements** in which specific products from certain countries are subject to lower tariffs or no tariffs, often in the context of development cooperation. Examples are the General Preferential Arrangement (GPA) of the EU and the United States.

- **Economic Partnership Agreements (EPAs)**

These are broader agreements that cover not only trade but also cooperation in areas such as investment, services, and regulation. The EPA between the EU and the Caribbean countries are examples of this. Preferential trade can provide significant economic benefits, such as access to larger markets, lower costs for consumers, and increased competition that drives innovation. At the same time, there are also living challenges, such as the negative impact domestic industries and the possible negative impact on certain sectors that cannot compete with imported goods.

The trade agreements for which Suriname is a party and offers preferences to Surinamese entrepreneurs are:

- **EUROPEAN - EU Economic Partnership Agreements** this agreement provides market access for goods (quota and duty free) and market access for services in the EU market
- **EUROPEAN - UK Economic Partnership Agreements** provides access to goods (quota and duty free) and services
- **EUROPEAN - Costa Rica Free Trade Agreements** provides market access to goods
- **Regional Treaty of Chaguanay (CARICOM)** Provides market access for goods and services, harmonization of regulations, etc.
- **EUROPEAN - Sudan** Provides market access for goods
- **Agreement Establishing the Free Trade Area between the Caribbean Community and the Dominican Republic** provides market access for goods
- **Partial Scope Agreement concerning/with Conventions for the Free Trade** offers the possibility to cooperate in the Suriname part of Brazil
- **Trade agreement between the Republic of Suriname and the Republic of Indonesia** provides access for goods intended for import and sale

ATTACH

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