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欣海报关

2021

Customs Consulting of Xinhai Newsletter in September

\ New version of Administrative Measures on Registration and Filing of Medical Devices
\ List of elements of spot check of import and export commodities other than legal inspection in 2021
\ Summary of emergency preventive measures
\ Analysis of the new CIQ policies in September

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Background of New Policy

- Measures for the administration of registration and filing of medical devices
- Administrative measures for registration and filing of in vitro diagnostic reagents

It is an effective supporting measure of the Regulations

- On February 9, 2021, Premier of the State Council, Li Keqiang signed State Council Order No.739, promulgating the new Regulations on Supervision and Administration of Medical Devices. In order to implement the new Regulations, meet the reform requirements of the examination and approval system for medical devices, establish a more scientific supervision and management system for medical devices, and strengthen the registration management of medical devices, the State Food and Drug Administration organized the revision of the original Administrative Measures for Registration of Medical Devices and Administrative Measures for Registration of in Vitro Diagnostic Reagents.

Examination and approval system reform

- Add a special chapter on special registration procedures, and stipulate the registration procedures for innovative products and the priority registration procedures.
- The emergency registration procedures are stipulated, and the inclusion scope and supporting policies of each procedure are defined.

Optimize scientific and efficient review and approval procedures

- Adjusted the relevant requirements of clinical evaluation of medical devices, and made clear the situation of exemption from submitting clinical evaluation data and the requirements of implied permission for clinical trial approval.
- Implement the responsibilities of all aspects of medical device registration and record management, and strengthen the connection of medical device registration acceptance, review and system verification.

Clear supervision responsibilities

- Improve the relevant provisions on risk control and on-site inspection of clinical trials, and establish a responsibility interview system.
- To strengthen the implementation of the main responsibility of medical device registrants and filers, it is required to strengthen the life cycle quality management of medical devices, and take responsibility for the safety, effectiveness and quality controllability of medical devices in the whole process of development, production, operation and use according to law.

The "streamline administration, delegate power, strengthen regulation and improve services" reform is simplified.

- Simplify the requirements for registration and filing materials such as overseas listing documents and inspection reports.
- Adjust the requirements for inspection reports of Class II and Class III medical devices, and make it clear that applicants for registration can submit self-inspection reports.

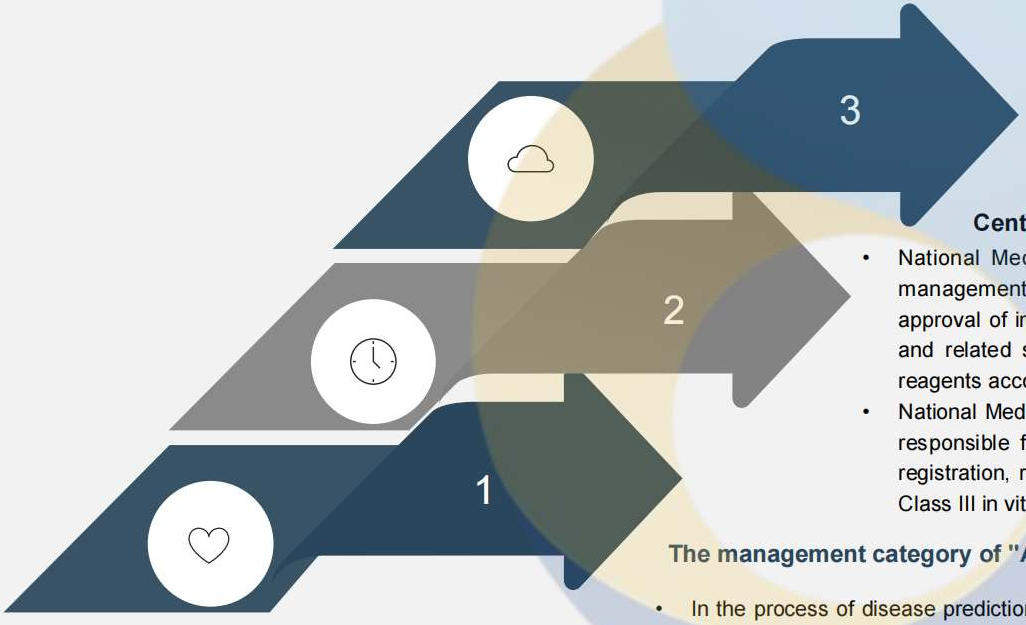
Administrative measures on registration and filing of medical devices (hereinafter referred to as "Administrative Measures")

Adjustment purpose	Adjustment measures	Rules of management measures
Fully implement the system of medical device registrants and filers	The main responsibility of medical device registrants and filers shall strengthen the quality management of the whole life cycle of medical devices, and take responsibility for the safety, effectiveness and quality controllability of medical devices in the whole process of development, production, operation and use according to law.	Article 9 of the Administrative Measures
Implied permission for clinical trials	Within 60 working days from the date of acceptance and payment of the application for examination and approval of clinical trials, if the applicant has not received the opinions of the device examination center (including the notice of expert consultation meeting and the notice of supplementary information) on the premise that the reserved contact information and mailing address are valid, the applicant can carry out clinical trials.	Article 40 of the Administrative Measures
Extended clinical trials	Medical devices that are undergoing clinical trials for the treatment of diseases that are seriously life-threatening and have no effective treatment means may benefit patients after medical observation. After ethical review and informed consent, they can be used for other patients with the same condition free of charge in institutions that conduct clinical trials of medical devices, and their safety data can be used for medical device registration applications.	Article 46 of the Administrative Measures
Condition approval regulations	For the treatment of rare diseases, serious life-threatening diseases without effective treatment means, and urgently needed medical devices such as responding to public health events, the drug supervision and administration department may make a conditional approval decision, and specify in the medical device registration certificate the validity period, the research work to be completed after listing, the completion time limit and other related matters.	Article 61 of the Administrative Measures

Administrative measures on registration and filing of medical devices (hereinafter referred to as "Administrative Measures")

Adjustment purpose	Adjustment measures	Rules of management measures
Simplify the requirements for registration and filing materials	Innovative medical devices that are not listed in the country (region) where the applicant or the filer is registered or produced do not need to submit the certificate that the competent department of the country (region) where the applicant or filer is registered allows the medical devices to be listed and sold. So as to encourage "global new" products to be listed in China as soon as possible.	Article 18 of the Administrative Measures
Adjust the inspection report requirements of Class II and Class III medical devices	The inspection report of medical device products submitted for registration or filing may be the self-inspection report of the applicant and the filing person, or the inspection report issued by a qualified medical device inspection institution entrusted.	Article 32 of the Administrative Measures
Medical device registration adds "product development" stage regulation	At the stage of product development, the applicant shall prepare product technical requirements, product specifications and labels, conduct non-clinical research on medical devices according to the applicable scope and technical characteristics of the products, conduct inspection according to the product technical requirements, and submit inspection reports.	Article 24-32 of the Administrative Measures
Exempt from submitting clinical evaluation data	(1) The working mechanism is clear, the design is finalized, the production process is mature, and the medical devices of the same variety that have been on the market have been used clinically for many years, and there is no serious adverse events record, and the routine usage is not changed; (2) Other non clinical evaluation can prove that the medical device is safe and effective.	Article 34 of the Administrative Measures
Three cases can be registered first	(1) The applicant legally owns the patent right of core technology invention in China through its leading technological innovation activities, or legally obtains the patent right of invention in China or its right to use through transferee, and the time for applying for the registration procedure of innovative products is within 5 years from the announcement date of patent authorization. (two) the applicant has completed the preliminary research of the product and has basically finalized products, the research process is true and controlled, and the research data is complete and traceable; (3) The main working principle or action mechanism of the product is the first in China, the performance or safety of the product has been fundamentally improved compared with that of similar products, and it is at the international leading level in technology and has significant clinical application value.	Article 73-75 of the Administrative Measures

Administrative measures for registration and filing of in vitro diagnostic reagents (hereinafter referred to as "Administrative Measures")



In vitro diagnostic reagent registration/filing agency

- The first kind of in vitro diagnostic reagents shall be subject to product record management. Class II and Class III in vitro diagnostic reagents shall be subject to product registration management.
- Import the first kind of in vitro diagnostic reagents for filing, and the filer shall submit the filing materials to National Medical Products Administration. Imported Class II and Class III in vitro diagnostic reagents shall be examined by National Medical Products Administration, and after approval, a medical device registration certificate shall be issued.

Centralized management department

- National Medical Products Administration is in charge of the registration and filing management of in vitro diagnostic reagents nationwide, organizes the review and approval of imported second and third types of in vitro diagnostic reagents, and records and related supervision and management of imported first type of in vitro diagnostic reagents according to law;
- National Medical Products Administration Medical Device Technical Evaluation Center is responsible for the technical evaluation of the application for registration, change of registration, renewal of registration, etc. of Class III in China and imported Class II and Class III in vitro diagnostic reagents.

The management category of "Administrative Measures"

- In the process of disease prediction, prevention, diagnosis, treatment monitoring, prognosis observation and health status evaluation, reagents, kits, calibrators, quality control products and other products used for in vitro detection of human samples can be used alone or in combination with instruments, appliances, equipment or systems. In vitro diagnostic reagents for blood source screening and in vitro diagnostic reagents labeled with radionuclides are not within the scope of administration measures.

Administrative measures for registration and filing of in vitro diagnostic reagents (hereinafter referred to as "Administrative Measures")

Basic Requirement

- The registration and filing of in vitro diagnostic reagents shall comply with the relevant requirements of the classification rules and catalogue of in vitro diagnostic reagents (Food and Drug Administration [2013] No.242);
- The applicant and the filing person shall establish a quality management system related to product development and production, and maintain effective operation.

Registration Filing

- Enterprises applying for registration/filing shall prepare technical requirements for in vitro diagnostic reagent products applying for registration or filing;
- The main raw materials and production process requirements should be specified in appendix form in the product technical requirements of the third kind of in vitro diagnostic reagents.
- When the same registration application includes different packaging specifications, only one packaging specification product can be inspected;
- The inspection report submitted for registration or filing can be either a self-inspection report or an inspection report issued by a third party.

Clinical Trial

- Clinical trials of in vitro diagnostic reagents refer to systematic research on the clinical performance of in vitro diagnostic reagents in the corresponding clinical environment.
- To carry out clinical evaluation of in vitro diagnostic reagents, clinical trials should be conducted to prove the safety and effectiveness of in vitro diagnostic reagents and the situation that they can be exempted from clinical trials [Article 37 of the Administrative Measures];
- The catalogue of Class II and Class III in vitro diagnostic reagents exempted from clinical trials shall be formulated, adjusted and published by the State Food and Drug Administration [Notice of the State Food and Drug Administration on Issuing the Catalogue of in vitro diagnostic reagents exempted from clinical trials (No.70 of 2021)].

Special Registration

- Special registration procedures include: innovative product registration procedures, priority registration procedures and emergency registration procedures;
- The special registration procedure here is consistent with the special registration procedure of the Administrative Measures for Registration and Filing of Medical Devices.
- If the design, raw materials, production process, application scope and use method of the registered in-vitro diagnostic reagent products of Class II and Class III are substantially changed, which may affect the safety and effectiveness of the in-vitro diagnostic reagent, the registrant shall apply to the original registration department for registration change; Other changes shall be filed with the original registration department within 30 days from the date of change.

Announcement on no longer issuing GSP certificate of origin for goods exported to Eurasian Economic Union

According to the report of the Eurasian Economic Commission, the Eurasian Economic Union decided not to grant GSP tariff preference to Chinese products exported to the Union from October 12, 2021. The relevant matters are hereby announced as follows:

1. Since October 12, 2021, the Customs will no longer issue GSP certificates of origin for goods exported to Eurasian Economic Union member countries.

2. if the consignors of goods exported to Eurasian Economic Union member countries need the certificate of origin, they can apply for the issuance of non preferential certificate of origin.

Hereby announce,

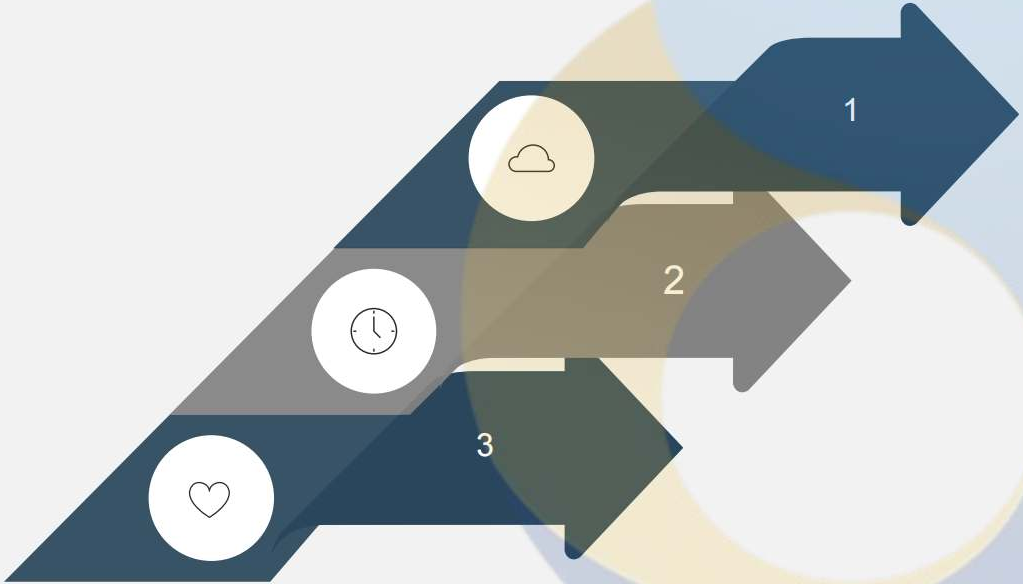
General Administration of Customs

September 23rd, 2021

What is GSP tariff preference?

- GSP, is a kind of tariff system, which refers to the general, non-discriminatory and non-reciprocal tariff system given by industrial developed countries to manufactured goods and semi-manufactured goods exported from developing countries or regions.
- This is after the Japanese Ministry of Finance no longer granted GSP tariff preference to Chinese goods exported to Japan since April 1, 2019, the newly added export goods exported to Eurasian Economic Union member countries have cancelled the issuance of GSP certificate of origin.

Announcement on no longer issuing GSP certificate of origin for goods exported to Eurasian Economic Union



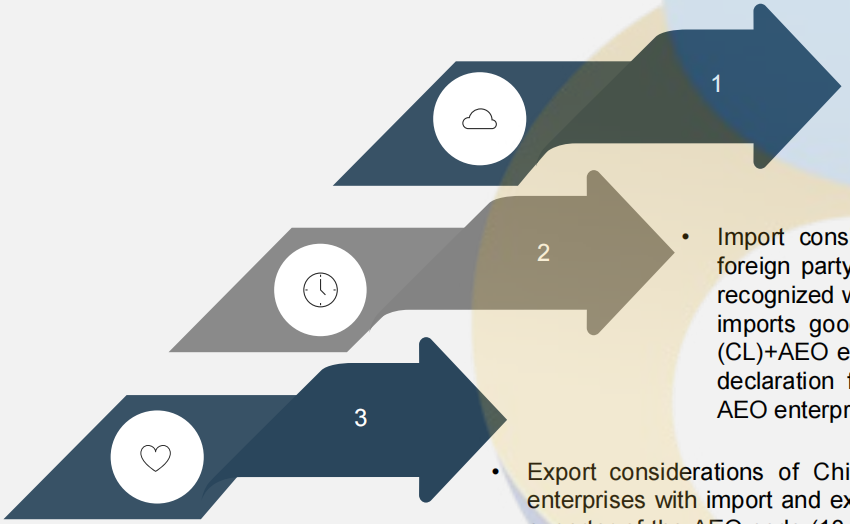
What are the member countries of Eurasian Economic Union?

Include Russia, Kazakhstan, Belarus, Kyrgyzstan and Armenia.

How should export enterprises respond and reduce the impact of this policy?

It is suggested that relevant enterprises seek diversified development strategies: pay attention to the promotion and implementation of various FTA policies, make full use of the FTA signed between China and ASEAN, Chile, Australia, Switzerland and other countries and regions, apply for various certificates of origin from the customs, and enjoy preferential tariffs of importers. At the same time, China is accelerating the negotiation process of the China-Japan-Korea Free Trade Area and the Regional Comprehensive Economic Partnership Agreement (RCEP). Once these two free trade agreements are established, a more comprehensive and mutually beneficial trade arrangement will be reached.

Announcement on implementation of China-Chile customs "Certified Operators" (AEO) mutual recognition



- According to the provisions of the Mutual Recognition Arrangement, China and Chile mutually recognize each other's "certified operators" of the other's customs to provide customs clearance facilities for the import and export goods of the AEO enterprises of both parties. Among them, the Chilean customs recognized the Chinese customs senior certification enterprises as mutual recognition AEO enterprises, and the Chinese customs recognized the Chilean customs "certified operators" as mutual recognition AEO enterprises. The announcement shall come into force on October 8.

- Import considerations for Chinese enterprises: It is recommended to confirm with the foreign party in advance whether the overseas consignor is an AEO enterprise mutually recognized with China, and obtain the AEO code as soon as possible. When an enterprise imports goods from an AEO enterprise in Chile, it is required to fill in "country code (CL)+AEO enterprise code (10 digits)" in the column of "overseas consignor" in the import declaration form, for example, "CL0123456789". After confirming the identity of Chile's AEO enterprise, the Chinese customs will provide relevant facilitation measures.

- Export considerations of Chinese enterprises: Chinese customs senior certification enterprises with import and export trade with Chile need to inform the Chilean importer or exporter of the AEO code (10-digit enterprise code registered with the Chinese customs of AEOCN+, for example, AEOCN0123456789), which will fill in the declaration in accordance with the regulations of the Chilean customs. The Chilean customs will provide relevant convenience measures after confirming the identity of the Chinese customs AEO enterprise.

Details of spot check elements of import and export commodities other than legal inspection in 2021

Announcement No.60 of the General Administration of Customs in 2021 (Announcement on Carrying out Spot Check Inspection of Import and Export Commodities Other than Statutory Inspection Commodities in 2021).

According to the Import and Export Commodity Inspection Law of the People's Republic of China and the relevant provisions of its implementing regulations, the General Administration of Customs has decided to carry out spot checks on some import and export commodities other than the legally inspected commodities from the date of this announcement. See Annex for the scope of spot checks.

The random inspection shall be carried out in accordance with the Administrative Measures for Random Inspection of Import and Export Commodities (promulgated by Order No.39 of the former General Administration of Quality Supervision, Inspection and Quarantine and amended by Order No.238 of the General Administration of Customs).

Hereby announce,
General Administration of Customs
August 12th, 2021

How to deal with unqualified spot checks？

- Imported goods: if the items related to personal and property safety, health and environmental protection are involved, the customs shall order the parties to destroy them, or issue a return treatment notice to go through the formalities of returning goods; Other unqualified items can be processed technically under the supervision of the customs, and can be sold or used only after passing the re-inspection by the customs;
- Export commodities: Unqualified commodities can be technically treated under the supervision of the customs, and only those that pass the re-inspection by the customs can be exported; Those who fail to pass the technical treatment or pass the re-inspection by the customs after the technical treatment shall not be exported.

Details of spot check elements of import and export commodities other than legal inspection in 2021

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Commodity	HS Code	Import/ Export	Trade Mode	Sampling quantity (must be of the same specification and model)
Dishwasher	8422110000	Import	General trade	2/batch
Air purifier	8421391000	Import	General trade	2/batch
Electronic toilet	8516799000 8543709990	Import	General trade	2/batch
Food processor	8509802000	Import	General trade	2/batch
Induction cooktop	8516601000	Import	General trade	2/batch
Printer	8443321100、8443321200、8443321300、8443321400、8443321900	Import	General trade	2/batch
Children garment	6104620090、6104630090、6105100090、6105200090、6105900000、6106100090、6106200090、6106900000、6115950019、6115960000、6115990000、6116910000、6116920000、6116930010、6116930090、6116990000、6201921000、6201931000、6202921000、6202931000、6203320090、6203330000、6204320090、6204330000、6204420000	Import	General trade	6115950019、6115960000、6115990000、6116910000、6116920000、6116930010 6116930090、6116990000 5 units ; other 3 units
Stationery	9608100000、9608200000、9608301000、9608302000、9608309000、9608400000、9608500000、9609101000、9609102000、9609200000、9609900000、4820200000	Import	General trade	Writing pens (with caps), 12/batch Writing pens (without caps), 12/batch Oil pastels, etc. (set), 3 sets Oil pastels, etc. (unit), 6 Book, 3 copies
Imitation ornaments	7117110000、7117190000、7117900000	Import	General trade	6/batch
Automobile interior parts	8708299000、3926300000、4016910000、8708995900	Import	General trade	5/batch （Sample length is at least 500mm）

Details of spot check elements of import and export commodities other than legal inspection in 2021

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(Continued from the above table)

Commodity	HS Code	Import/ Export	Trade Mode	Sampling quantity (must be of the same specification and model)
Helmet	6506100090	Import	General trade	2/batch
Car seat	9401209000	Import	General trade	2/batch
Plates, dishes, basins, cups and similar articles of paper or cardboard	4823400000	Import	General trade	20/batch (30 pieces in each batch for small samples)
Festival light string	9405300000	Export	General trade	3/batch
LED lighting	8539501000 8539502000 9405409000	Export	General trade	6/batch
Child's bike	9503001000	Export	General trade	3/batch
Children's scooter	9503001000 9503009000 9506990000 8503001000	Export	General trade	2/batch
Cloth toys, bamboo toys, plastic toys, electric toys	9503002100 9503002900 9503006000	Export	General trade	3 toys other than electric toys; 4 electric toys
Plastic food contact products	3924100000	Export	General trade	20/batch (30 pieces in each batch for small samples)

Summary of emergency preventive measures taken by the General Administration of Customs for overseas enterprises

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Country	Overseas manufacturers	Specific notice/announcement
Russia Korea	South Korean manufacturer SAJO SEAFOOD CO., LTD (The registration number is KP-154) Russian manufacturer Tongchi International Co., LTD (The registration number is CH-889)	As Covid-19 nucleic acid test was positive from two outer packaging samples of a batch of frozen narrow cod imported from South Korea and Russian manufacturers, according to the regulation of Announcement No.103 of General Administration of Customs in 2020, the national customs continued to suspend the import declaration of products from South Korean manufacturer SAJO SEAFOOD CO., LTD (registration number KP-154) for 4 weeks from September 11, and Russian manufacturer Tongchi International Co., LTD (registration number CH-88).
Indonesia	Manufacturer PT. NAGA LAUT TIMUR (The registration number: CR 690-14)	As Covid-19 nucleic acid test was positive from three outer packaging samples of two batches of frozen squid imported from Indonesia, according to the regulations of General Administration of Customs Announcement No.103 of 2020, the national customs suspended the product import declaration from PT. NAGA LAUT TIMUR (registration number CR 690-14), an Indonesian aquatic product manufacturer, for 2 weeks from September 4.
Peru	Manufacturer enterprise PRODUCTORA ANDINA DE CONGELADOS S.R.L.（The registration number isP234-SUL-POAD）、SEAFROST S.A.C.（The registration number is P107-PAI-SEAF）	As Covid-19 nucleic acid test was positive from three outer and one inner packing samples of three batches of frozen squid imported from Peru, according to the regulation of General Administration of Customs Announcement No.103 of 2020, the national customs suspended the import declaration of products from Peru aquatic products manufacturer Productora Andina de Congelados S.R.L. (registration number P234-SUL-POAD) for five weeks, and SEAFROST S.A.C.
Russia	Manufacturer enterprise Polluks JSC “Okeanrybflot”（The registration number is CH-16F）	As Covid-19 nucleic acid test was positive from one outer packing sample of a batch of frozen narrow cod imported from Russia, according to the regulation of Announcement No.103 of General Administration of Customs in 2020, the national customs suspended the import declaration of products from Russian manufacturer Polluks JSC “Okeanrybflot "(registration number: CH-16F) for one week from September 9.
India	Manufacturer enterprise M/s. Malpefresh Marine Export Private Limited（The registration number is 1557）、M/s. Castlerock Fisheries Pvt. Ltd.（The registration number is 277）	As Covid-19 nucleic acid test positive was detected from two outer packaging samples of two batches of frozen South American white shrimp imported from India, according to the regulations of General Administration of Customs Announcement No.103 of 2020, the national customs has suspended the import of products from Indian aquatic products manufacturers M/S. Malpefresh Marine Export Private Limited (registration number 1557) and M/S. Castle Rock Fisheries PVT. Ltd. (registration number 277) since September 23.

Summary of emergency preventive measures taken by the General Administration of Customs for overseas enterprises

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Country	Overseas manufacturers	Specific notice/announcement
Indonesia	Manufacturer enterprise UD. PIALA（The registration number is CR 174-32）、PT. KARYA MINA PUTRA（The registration number is CR 554-14）and PT. SANJAYA INTERNASIONAL FISHERY（The registration number is CR 513-12）	Covid-19 nucleic acid positive was detected from an outer packaging sample of a batch of frozen Penaeus monodon imported from Indonesia. Covid-19 nucleic acid test was positive in 5 outer packaging samples of a batch of frozen hairtail. Covid-19 nucleic acid positive was detected in an inner packaging sample of a batch of frozen squid. According to the regulations of General Administration of Customs Announcement No.103 of 2020, the national customs suspended the acceptance of Indonesian aquatic products manufacturers UD. PIALA (registration number CR 174-32), PT. KARYA MINA PUTRA (registration number CR 554-14) and PT. SANJAYA INTERNASIONAL FISHERY (registration number) from September 9.
Pakistan	Manufacturer enterprise M/s. Orb Exim Corporation (Pvt). LTD（The registration number is T-594/07）	As Covid-19 nucleic acid was positive from an inner packing sample of a batch of frozen sardines imported from Pakistan, according to the regulation of Announcement No.103 of General Administration of Customs, the national customs suspended the import declaration of products from M/S. Orbexim Corporation (PVT). Ltd (registration number: T-594/07), a Pakistani aquatic product manufacturer, for one week from September 9.
Myanmar	Manufacturer enterprise Ocean World Coldstore（The registration number is YGN/132/OW/DOF）	As Covid-19 nucleic acid test is positive from one outer packaging sample of a batch of frozen hairtail imported from Myanmar, according to the regulations of Announcement No.103 of General Administration of Customs, the national customs has suspended the import declaration of products from Ocean World Coldstore (The registration number YGN/132/OW/DOF), a Myanmar aquatic product manufacturer, for one week since September 9.
Myanmar	Manufacturer enterprise Myanmar Golden Phoenix Industrial Co., Ltd（The registration number is YGN/023/MGPI/DOF）	Covid-19 nucleic acid test was positive from one outer packaging sample of a batch of frozen hairtail imported from Myanmar. Covid-19 nucleic acid positive was detected in 8 outer packages and 1 inner package sample of a batch of frozen squid, etc. According to the regulation of Announcement No.103 of General Administration of Customs, the national customs has suspended the import declaration of Myanmar Golden Phoenix Industrial Co., Ltd (The registration number YGN/023/MGPI/DOF) for 2 weeks since September 14.
Myanmar	Manufacturer enterprise Ocean World Coldstore（The registration number is YGN/132/OW/DOF）	As Covid-19 nucleic acid test was positive from one outer packaging sample of a batch of frozen hairtail imported from Myanmar, according to the regulations of Announcement No.103 of General Administration of Customs, the national customs continued to suspend the import declaration of products from Ocean World Coldstore (registration number YGN/132/OW/DOF), a Myanmar aquatic product manufacturer, for 4 weeks from September 24th.

Summary of emergency preventive measures taken by the General Administration of Customs for overseas enterprises

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Country	Overseas manufacturers	Specific notice/announcement
Russia	Russian processing fishing boat Seawind-1 PJSC “NBAMR” (Public Joint Stock Company “Nakhodka Active Marine Fishery Base”)（The registration number is CH-21P）、Manufacturer enterprise Oktyabrskiy-1 Co LLC fish plant № 2（The registration number is CH-734）	Covid-19 nucleic acid test was positive from four outer packing samples of a batch of frozen narrow cod imported from Russia. Covid-19 nucleic acid positive was detected in one outer packaging sample of a batch of frozen Pacific COD. According to the regulations of Announcement No.103 of General Administration of Customs in 2020, since September 14, the whole country has suspended the import declaration of products from the Russian processing fishing boat Seawind-1 PJSC "NBA MR" (Public Joint Stock Company "Nakhodka Active Marine Fishery Base") (registration number CH-21P) and the production enterprise Oktyabrskiy-1 Co LLC fish plant № 2 (registration number CH-734) for one week.
Pakistan	Manufacturer enterprise M/s. Neelam Seafoods（registration number is T-386/2015）and M/s. Arsala Seafoods（The registration number is T-480/2005，T-23/08）	As Covid-19 nucleic acid test was positive from one outer package and one inner package sample of two batches of frozen cuttlefish imported from Pakistan, according to the regulation of General Administration of Customs Announcement No.103 of 2020, the national customs has suspended the acceptance of M/S. Neelam Seafood (registration number T-386/2015) and M/S. Arsala Seafood (registration number T-44) from Pakistani aquatic products manufacturers since September 16.
Myanmar	Manufacturer enterprise Ocean World Coldstore（The registration number is YGN/132/OW/DOF）	As Covid-19 nucleic acid test wass positive from one outer packaging sample of a batch of frozen hairtail imported from Myanmar, according to the regulations of Announcement No.103 of General Administration of Customs, the national customs has suspended the import declaration of products from Ocean World Coldstore (registration number YGN/132/OW/DOF), a Myanmar aquatic product manufacturer, for one week since September 17.
Vietnam	Investment commerce fisheries corporation-INCOMFISH Corporation, Seafood & foodstuff processing factory No 5（The registration number is HK 187）	As Covid-19 nucleic acid test was positive from one outer packaging sample of a batch of dried small male fish imported from Vietnam, according to the regulations of Announcement No.103 of General Administration of Customs, the national customs has suspended the acceptance of Vietnamese aquatic products enterprise Investment Commerce Fisheries Corporation-Inconfish Corporation since September 19. Import declaration for seafood & foodstuff processing factory no 5 (The registration number: HK 187) is 1 week.

Analysis of the new CIQ policies in September

Category	Announcement No.	Comments
Supervision of animal and plant products	Announcement of Ministry of Agriculture and Rural Affairs of General Administration of Customs No.71, 2021	Announcement on preventing Botswana highly pathogenic avian influenza from being introduced into China. From September 14, 2021, it is prohibited to import poultry and related products directly or indirectly from Botswana, including products originating from unprocessed poultry or processed poultry that may still spread epidemics.
	Announcement of Ministry of Agriculture and Rural Affairs of General Administration of Customs No.67, 2021	Announcement on preventing highly pathogenic avian influenza from Benin from being introduced into China. From September 1, 2021, it is prohibited to import poultry and related products directly or indirectly from Benin, including products from poultry that are unprocessed or processed but may still spread epidemics.
	Announcement No.66 of the General Administration of Customs in 2021	Announcement on quarantine requirements for imported fresh citrus plants from south africa. From August 31st, 2021, South African fresh citrus which meets the requirements will be allowed to be imported. The fresh citrus exported to China includes Orange (scientific name: Citrus sinensis, English name: orange), Grapefruit (scientific name: Citrus paradisi, English name: grapefruit), Lemon (scientific name: Citrus limon, English name: lemon) and orange (scientific name: Citrus reticulata, English name: Mandarin). It must be citrus produced in South Africa. It is required to standardize the approved orchards, packaging plants and quarantine treatment facilities, quarantine pests, orchard management, packaging plant management, packaging requirements, quarantine treatment requirements, pre-export quarantine, plant quarantine certificate requirements, entry inspection and quarantine and unqualified treatment.
	Announcement No.65 of the General Administration of Customs in 2021	Announcement on inspection and quarantine requirements for imported Uzbekistan prunes. From August 26th, 2021, Uzbekistan dried plums that meet the requirements are allowed to be imported. Dried Chinese plums refer to fresh plums produced in Uzbekistan, which are processed by selecting, cleaning, soaking and drying. The requirements are standardized from six aspects, such as the requirements of production enterprises, phytosanitary requirements, phytosanitary certificates, food safety requirements, packaging requirements and transportation requirements.

Analysis of the new CIQ policies in September

Category	Announcement No.	Comments
Supervision of animal and plant products	Announcement No.64 of the General Administration of Customs in 2021	Announcement on plant quarantine requirements of imported kazakhstan forage barley powder. From August 20, 2021, the import of Kazakhstan forage barley powder that meets the requirements will be allowed. Feed barley powder refers to the fine powdery feed raw material (whole barley powder containing bran) obtained by processing barley produced in Kazakhstan. Barley used to produce forage barley powder exported to China shall come from warehousing enterprises registered by the General Administration of Customs of the People's Republic of China. The requirements are standardized from five aspects: enterprise registration, quarantine pests, product production and export requirements, entry inspection and quarantine, and treatment of nonconformities.
	Announcement of Ministry of Agriculture and Rural Affairs of General Administration of Customs No.63, 2021.	Announcement on preventing dominican african swine fever from being introduced into China. Since August 18, 2021, it is forbidden to import pigs, wild boars and their products directly or indirectly from Dominica. Once found, they will be returned or destroyed.
Import and export food safety	The General Administration of Customs Import and Export Food Safety Risk Warning Form No.1 of 2021	Warning circular on strengthening inspection and quarantine supervision of casings exported from france to china. From September 1, 2021, the import declaration of products from the French casing manufacturer "BOYAUDERIE BLESOISE SAS FOLLET" (French registration number FR 41018004 CE) will be suspended. Suspension of the veterinary hygiene certificate of French casings exported to China issued by Viviane MARIAU, the veterinary officer signed by France.
Administrative approva	General Office of the Ministry of Commerce (Notice on Doing a Good Job in Applying for Export License of Automobile and Motorcycle in 2022).	From September 10, 2021, enterprises exporting automobiles and motorcycles can apply for the qualification of export enterprises in 2022 online through the unified platform of the data center of the Quota and License Bureau of the Ministry of Commerce. The local commerce authorities shall complete the preliminary examination of the application materials for export license of automobile and motorcycle enterprises before October 11th and submit them to the Ministry of Commerce online.
	Announcement No.15, 2021 of the Ministry of Agriculture and Rural Affairs of the State Forestry and Grassland Bureau.	The newly adjusted List of National Key Protected Wild Plants has been listed in 455 species and 40 categories of national key protected wild plants, including 54 species and 4 categories of national first-class protected wild plants and 401 species and 36 categories of national second-class protected wild plants. Among them, 324 species and 25 classes are managed by the competent forestry and grassland departments, and 131 species and 15 classes are managed by the competent agricultural and rural departments.

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