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**INDUSTRY STANDARD OF THE
PEOPLE'S REPUBLIC OF CHINA**

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Codes of permit for hygiene for entry-exit special products

入出境特殊物品卫生许可规程

(English Translation)

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Foreword

Appendix D of this standard is a normative appendix. Appendix A, Appendix B, Appendix C, Appendix E, Appendix F, and Appendix G are informative appendices.

This standard is proposed and managed by the National Certification and Accreditation Administration.

This standard was drafted. Beijing Entry-Exit Inspection and Quarantine Bureau of the People's Republic of China.

1 Scope

This standard specifies the procedures for the work permit for the entry and exit of special items.

This standard applies to the health and quarantine examination and approval of entry and exit special items.

2 Requirements

2.1

The implementing authority for the sanitation permit for entry and exit special items (hereinafter referred to as the implementing agency) is the inspection and quarantine organ designated by the AQSIQ.

2.2

The implementing agency shall base the conditions,, procedures, deadlines and all materials required for the submission of the special permit for entry and exit.

The documentary and application model texts are published in the office and published on

the official website.

3.1 Application

The owner, applicant or his agent of special items entering or leaving the country shall submit special items for entry and exit in accordance with the application procedures publicly disclosed by the implementing agency.

Hygiene permit and submit the appropriate materials.

3.2 Acceptance

3.2.1

The directly affiliated inspection and quarantine institution shall meet the scope of the sanitation permit for special articles, complete the application materials, conform to the legal form, or the applicant If all the application materials for correction are submitted in accordance with the requirements of the inspection and quarantine institution, they shall be accepted.

3.2.2

The application for the sanitation permit for entry and exit special items decided to be accepted shall be issued by the directly affiliated inspection and quarantine agency.

The Administrative License Application Acceptance Decision (see Appendix A), stamped with the special seal of the administrative license of the inspection and quarantine institution and dated.

3.2.3

If the entry and exit products are not in the scope of special item management, the applicant shall be notified immediately that the application will not be accepted.

3.2.4

If the applicant's application does not fall within the scope of the authority of the inspection and quarantine organ, it shall immediately make a decision not to accept the application and inform the application.

The person applies to other relevant administrative agencies.

3.2.5

For applications that are not accepted, the special inspection and quarantine agency shall issue a special seal and date for the administrative license of the inspection and quarantine institution.

The "Qualification Supervision, Inspection and Quarantine Administrative License Non-acceptance Decision" (see Appendix B).

3.2.6

If there is any error in the application materials that can be corrected on the spot, the applicant should be allowed to correct on the spot.

3.3 Review

1) Relevant competent authorities include. health administrative department, human genetic resources management office, drug supervision department, environmental protection department and His administrative department.

3.3.1

The following information should be reviewed for microorganisms, human tissues, organs and blood entering and leaving the country.

--- The relevant competent authority 1) issued the original and a copy of the permit for entry and exit (see Appendix C);

--- The scientific name (Chinese and Latin) and biological characteristics (Chinese and English) of pathogenic microorganisms contained in special articles file;

--- Entry special items containing or possibly containing grade 3 to 4 pathogenic microorganisms, and containing or may contain pathogens that have not yet been classified Special items for living organisms, using the P3 laboratory qualification certificate provided by the unit;

---Special items for research purposes, original documents for research projects approved by the employer or applicants for research projects and domestic Original and photocopy of the agreement of the foreign cooperation agency;

--- The donor health certificate or inspection report issued by the qualified hospital of the transplanted country (or region) provided by the applicant.

3.3.2

Entry biological products and blood products used for the treatment, prevention and diagnosis of human diseases shall be examined and issued by the State Drug Administration.

Import registration certificate.