Inspection and confirmation of compliance with the standard provided for in Articles 5 to 19 of the Ministerial Ordinance on Good Laboratory Practice for Agricultural Chemicals

(Notification: 30 Shouan No.4215, issued on November 30, 2018 by Director-General, Food Safety and Consumer Affairs Bureau, the Ministry of Agriculture, Forestry and Fisheries)

DISCLAIMER:

The translations of notification on this page are not official. Only the original Japanese texts of notification have legal effect, and the translations are to be used solely as reference materials to aid in the understanding of the original texts. Neither FAMIC nor the government of Japan shall be held responsible for the accuracy, reliability or currency of the translations provided on this website, or for any consequences resulting from the use of information on this website. For all purposes of interpreting and applying notices to any legal issue or dispute, users should consult the original Japanese texts. The translations may be revised in the future. The data contained on this page may be reprinted or reproduced, provided that the disclaimer is accompanied and the source is shown.

Revision: 1 Shouan No.914, issued on June 28, 2019 Revision: 2 Shouan No.4310, issued on Jan 5, 2021 Confirmation and Inspection on compliance with the standard provided for in Articles 5 to 19 of the Ministerial Ordinance on Good Laboratory Practice for Agricultural Chemicals

1 Purpose

Pursuant to Article 3, paragraph (2) of the Agricultural Chemicals Regulation Act (Act No. 82 of 1948), the test results specified in Article 2 (hereinafter referred to as "Specified Test Results") of the "Ministerial Ordinance of the Good Laboratory Practice for Agricultural Chemicals" (hereinafter referred to as "Ministerial Ordinance") must be derived from tests conducted in accordance with the standard provided for in Articles 5 to 19 of the Ministerial Ordinance as necessary to ensure the reliability of the results (hereinafter referred to as "Agricultural Chemicals GLP Standards").

This notification is to provide for the procedures of inspection and confirmation of compliance with the Agricultural Chemicals GLP Standards.

2 Confirmation of GLP Compliance by the Director-General, Food Safety and Consumer Affairs Bureau

- (1) Confirmation of compliance with the Agricultural Chemicals GLP Standards must be made in accordance with either (i) or (ii) below:
 - (i) The Director-General, Food Safety and Consumer Affairs Bureau, Ministry of Agriculture, Forestry and Fisheries (hereinafter referred to as the "Director-General, Food Safety and Consumer Affairs Bureau") confirms that the test facility conducting studies is in compliance with the Agricultural Chemicals GLP Standards (hereinafter referred to as a "Confirmation of GLP Compliance") at least once in every three years; or
 - (ii) A government agency or an organization representing it confirms that the test facility conducting studies is in compliance with standards of GLP (limited to the programs that have been confirmed to be in compliance with GLP principles of the Organization for Economic Co-operation and Development (OECD) or the programs of the countries which have concluded bilateral arrangements with Japan), on the basis of which the Director-General, Food Safety and Consumer Affairs Bureau decides that the test facility conducting studies is in compliance with the Agricultural Chemicals GLP Standards.
- (2) Regardless of (1), (i), if any doubt arises, as the result of the examination in Article 3, paragraph (4), Article 7, paragraph (2) or Article 8, paragraph (4) of the Act or on the basis of other information, as to the reliability of the specified test results provided in Article 3, paragraph (2), Article 7, paragraph (1) or Article 8, paragraph (3) of the Act, and when the notice thereof is issued by the Director-General, Food Safety and Consumer Affairs Bureau, the test facility must undergo a confirmation of GLP compliance by the Director-General, Food Safety and Consumer Affairs Bureau, the test facility must undergo a confirmation of eliminate the doubts regarding the reliability of the Specified Test Results.

3 Application for Confirmation of GLP Compliance

- (1) When conducting studies subject to a Confirmation of GLP Compliance in Article 2, (1), (i), a person or, in case of a corporation, its representative (hereinafter referred to as "GLP Confirmation Applicant.") must submit the application for Confirmation of GLP Compliance of test facility (Appended Form 1), and the applicants for the registration of agricultural chemicals under 2, (2) must submit the application for Confirmation of GLP Compliance (Appended Form 2) to the Director-General Food Safety and Consumer Affairs Bureau along with the following documents from (i) to (iv):
 - (i) Overview of the test facility (name, location, date of establishment, organizational and personnel composition (describe both the organizational affiliations in the facility and the organizational

affiliations under the Agricultural Chemicals GLP Standards), site area and building area, internal and external building layout, and the types and specifications of equipment);

- (ii) Summary of the competence to conduct Specified Tests and the numbers of the specified test completed over the past three years;
- (iii) List of Standard Operating Procedures and
- (iv) Documents concerning Test Facility Management (Test Site Management, if applicable), Study Director, (Principal Investigator), study personnel, personnel of Quality Assurance Unit and Archivist, which list the names, career backgrounds, research histories and names of academic societies and academic organizations to which they belong.
- (2) Submission of the application for Confirmation of GLP Compliance of test facility and the relevant documents in (1), can be made via Independent Administrative Agency, Food and Agricultural Materials Inspection Center (hereinafter referred to as "FAMIC"). In this case, FAMIC may append comments, etc. to the application.
- (3) The Director-General, Food Safety and Consumer Affairs Bureau may request a GLP Confirmation Applicant to submit further documents and other materials necessary for assessing GLP Compliance. In this case, the GLP Confirmation Applicant may submit them to the Director-General, Food Safety and Consumer Affairs Bureau via FAMIC.

4 Examination with document and Inspection

The Director-General Food Safety and Consumer Affairs Bureau examamines the documents submitted in 3 above, and then, if it is found to be necessary to conduct an inspection, the Director-General may have either of the followings inspect the test facility:

- (i) FAMIC;
- (ii) Those appointed by the Director-General among those who have sufficient knowledge or experience on the inspection related to test results or specimens, etc.

5 Inspection Procedure

- (1) Prior to the inspection, the Director-General, Food Safety Consumer Affairs Bureau notifies the GLP Confirmation Applicant of the date of the inspection and other necessary matters.
- (2) Inspection is conducted in accordance with the following steps.
 - (i) Inspection of the overall management of the test facility;
 - (ii) Inspection of the facility and the maintenance of apparatus;
 - (iii) Inspection of operations;
 - (iv) Inspection of the documentation of study plan, Standard Operating Procedures, final report, etc.;
 - (v) Inspection of the activity of the Quality Assurance Unit;
 - (vi) Inspection of the management of retained raw data, specimen, etc.; and
 - (vii) Inspection and verification of raw data, specimens, final reports, etc. related to the studies for audit
- (3) If it is deemed necessary as a result of the inspection, a request may be made to submit samples of test item, etc., specimens, raw data, and other necessary materials.
- (4) After the inspection is completed, the personnel who conducted the inspection (hereinafter referred to as the "Inspector") is to give advice or guidance on-site as necessary. The advice or guidance must be recorded.

6 Reporting the results of inspections

(1) The Inspector must prepare an inspection report including the following items, and submit it to the Director-General, Food Safety and Consumer Affairs Bureau after conducting the inspection:

- (i) Name(s) and affiliation(s) of the Inspector(s);
- (ii) Name and Address of the test facility inspected;
- (iii) Date(s) of inspection;
- (iv) Purpose of the inspection;
- (v) Overview of the test facility;
- (vi) Type(s) of study audited;
- (vii) Name(s) and affiliation(s) of personnel of the test facility who attended in the inspection;
- (viii) Compliance with the Agricultural Chemicals GLP Standards on the test facility and studies inspected;
- (ix) Conclusion; and
- (x) Other items
 - Improvement(s) relating to deviation(s) found in the last inspection;
 - Implementable studies
- (2) Based on the inspection report, the Director-General, Food Safety and Consumer Affairs Bureau verifies whether the test facility complies with the Agricultural Chemicals GLP Standards, and notifies the GLP Confirmation Applicant of the result.

7 Others

- (1) The representative of the test facility that has received the confirmation of the GLP compliance from the Director-General, Food Safety and Consumer Affairs Bureau under 2, (1), (i) or (2) must submit a notification concerning the change of the application for Confirmation of GLP Compliance (Appended Form 3) to the Director-General, Food Safety and Consumer Affairs Bureau, in cases where any changes to the following items occur in the contents of the application for Confirmation of GLP Compliance of Test Facility (Appended Form 1):
 - (i) Address or name of the GLP Confirmation Applicant; or
 - (ii) Name or address of the test facility.
- (2) The representative of the test facility that has received confirmation of GLP compliance from the Director-General, Food Safety and Consumer Affairs Bureau under 2, (1), (i) or (2) is promptly to submit the notification of business abolition (Appended Form 4) to the Director-General, Food Safety and Consumer Affairs Bureau, in cases where the GLP operation of a test facility is to be closed.

Supplementary provisions (December 1, 2018)

1 This notification applies on December 1, 2018. However, it does not apply to the test results listed below.

- Following test results initiated before September 30, 1984: Acute oral toxicity, acute dermal toxicity, acute inhalation toxicity, skin irritation, eye irritation, skin sensitization, acute delayed neurotoxicity, 90-day repeated dose oral toxicity, 21-day repeated dose dermal toxicity, 90-day repeated inhalation toxicity, 28-day repeated dose delayed neurotoxicity, 1-year repeated dose oral toxicity, carcinogenicity, reproductive toxicity, teratogenicity and mutagenicity;
- (2) Following test results initiated before August 28, 1997: Human safety of microbial pesticides;
- (3) Following test results initiated before September 30, 1999: Properties, stability, degradability, etc. of active ingredient (excluding test results regarding color, shape and odor, and bioconcentration);
- (4) Following test results initiated before January 31, 2001: Acute neurotoxicity, repeated dose oral neurotoxicity, detoxification methods or emergency medical treatment, metabolism in animals, metabolism in plants, behavior in soil, behavior in water, fish acute toxicity, Acute immobilization on daphnids, Reproduction on daphnids, and growth inhibition on algae;

- (5) Following test results initiated before January 9, 2003: Analysis of composition of an agricultural chemical's TGAI, and analysis of dioxins in an agricultural chemical TGAI;
- (6) Following test results initiated before March 31, 2005: Fish (incubated larvae) acute toxicity, Acute immobilization on daphnids (adults), effects of coexisting organic substances on fish acute toxicity / acute swimming inhibition of the genus Daphnia, Acute toxicity on freshwater shrimps (Paratya compressa compressa and Paratya compressa improvisa) and amphipoda, and Acute immobilization on chironomid larvae;
- (7) Following test results initiated before October 1, 2007: Bioconcentration;
- (8) Following test results initiated before March 31, 2011: Residues in crops;
- (9) Following test results initiated before November 13, 2014: Residues in livestock; and
- (10) Following test results initiated before March 31, 2017: Analysis of composition of an agricultural chemical TGAI used in toxicity tests, toxicity for additives and impurities, and Analysis methods on an agricultural chemical TGAI
- 2 "The Good Laboratory Practice for Agricultural Chemicals" (The notification: 11 Nousan No.6283, issued on October 1, 1999 by Director-General of Agricultural Production Bureau, the Ministry of Agriculture, Forestry and Fisheries; hereinafter referred to as the "former notification") is hereby abolished.
- 3 With regard to an application for the confirmation of the GLP compliance based on 4 of the former notification filed prior to the enforcement of this notification, and for which no notification has been made for the results of the confirmation of the GLP compliance on 7, (2), of the former notification, the procedure for the confirmation of the GLP compliance then in force must remain applicable.
- 4 With regard to test results that may not be in compliance with the standards set out in Articles 5 to 19 of the Ministerial Ordinance on GLP for Agricultural Chemicals under the provision of paragraph (2) of the supplementary provisions of the Ministerial Ordinance on GLP for Agricultural Chemicals, and that are subject to the Agricultural Chemicals GLP Standards in the former notification (other than those listed in 1, (1) to (10)), the provisions of the former Notification (including the attached standards) is to remain in force.

Supplementary provisions (June 28, 2019)

- 1 This notification applies on July 1, 2019.
- 2 Notwithstanding the provisions of the preceding paragraph, the provisions for revising relating to the following studies applies on April 1, 2020: Physical and chemical properties of formulation products in the physical and chemical properties area; dermal absorption and Exposure of operator to agricultural chemicals applying them in the field in the toxicity area; growth inhibition on algae and cyanobacteria, growth inhibition on Lemna sp., acute oral toxicity on avian, acute dermal toxicity on honeybees (adults), acute oral toxicity on honeybees (adults), chronic oral toxicity on honeybees (adults), oral toxicity on honeybees (larvae) and residues in pollen and nectar in the adverse effects on aquatic and terrestrial organisms area; provided however, that it does not preclude the application of the provisions of the revised notification on the Confirmation of GLP Compliance, etc. to these studies on or before March 31, 2020.



(Japanese Industrial Standard A4)

Remarks

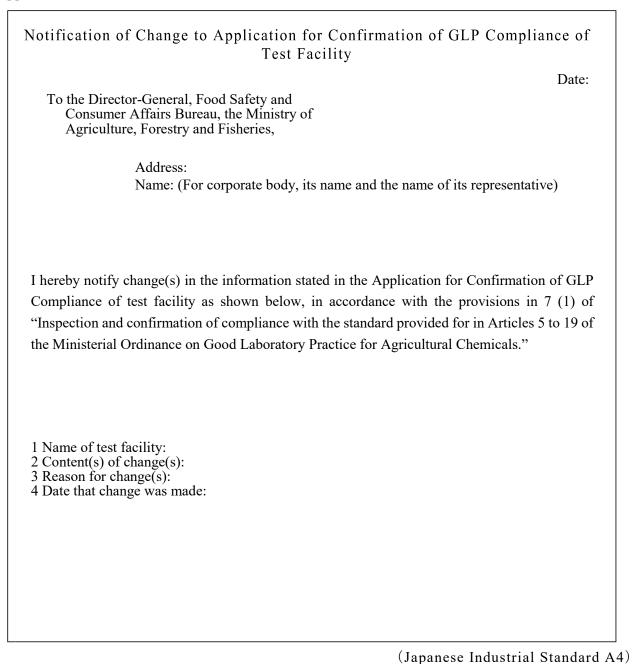
- (Note 1) The documents, etc. provided in "3: Application for Confirmation of GLP Compliance" are to be attached as related documents.
- (Note 2) For Notes 1 and 2 above, write them in English alongside.
- (Note 3) Refer to appendix for the area in 3 and 4 above.

Application for Confirmation of GLP Compliance of Test result
Date: To the Director-General, Food Safety and Consumer Affairs Bureau, the Ministry of Agriculture, Forestry and Fisheries, Address: Name: (For corporate body, its name and the name of its representative)
I have been notified on [yymmdd] that there are doubts regarding the reliability of the test result indicated below. In order to have the doubts eliminated, I hereby apply for inspection with relevant documents attached.
1 Title of the test result(s) concerning the doubts: 2 Name of test facility: 3 Address of test facility:

(Japanese Industrial Standard A4)

Remarks

- (Note 1) The documents, etc. provided in "3: Application for Confirmation of GLP Compliance" are to be attached as related documents.
- (Note 2) For 2 and 3 above, write them in English alongside.



7

Notification of Business abolition	
To the Director-General, Food Safety and Consumer Affairs Bureau, the Ministry of Agriculture, Forestry and Fisheries, Address: Name: (For corporate body, its name and the name of its representative	Date: e)
I hereby notify the discontinuation of the business of GLP test facility as follows, in acc with the provisions in 7 (2) of "Inspection and Confirmation of compliance with the s provided for in Articles 5 to 19 of the Ministerial Ordinance on Good Laboratory Prac Agricultural Chemicals".	tandard
 Name of test facility: Contents of abolishment: Reason for abolishment: List of place of transfers of final reports, specimens, raw data, etc. after abolishmet (3) (Scheduled) date of abolishment: 	nt:

(Japanese Industrial Standard A4)

Remarks

(Note) About 2 (2), the list may be submitted as an attached document of the application.

Appendix

Area	Name of study
Analysis of the technical grade active ingredient (TGAI) Physical and chemical properties	Analysis of composition of an agricultural chemical TGAI
	Analysis methods on an agricultural chemical TGAI
	Physical and chemical properties of an active ingredient
	Physical and chemical properties of formulation products
Toxicity	Acute oral toxicity
	Acute dermal toxicity
	Acute inhalation toxicity
	Skin irritation
-	Eye irritation
	Skin sensitization
	Acute neurotoxicity
<u>ہ</u>	Acute delayed neurotoxicity
	90-day repeated dose oral toxicity
	21-day repeated dose dermal toxicity
	90-day repeated inhalation toxicity
-	Repeated dose oral neurotoxicity
	28-day repeated dose delayed neurotoxicity
	Chronic toxicity
	Carcinogenicity
	Combined of Carcinogenicity and Carcinogenicity
	Reproductive toxicity
	Developmental toxicity
	Developmental neurotoxicity
	Detoxification methods or emergency medical treatment
-	Dermal absorption
	Exposure of operator to agricultural chemicals applying them in the field
	metabolism in animals
	Acute oral toxicity / infectivity
	Acute dermal toxicity / infectivity
	Acute pulmonary toxicity / infectivity
	Acute intravenous toxicity / infectivity
	Cell culture
	Repeated dose toxicity / infectivity
	Reproductive toxicity / infectivity
	Viral carcinogenicity
	Immunodeficiency inducing

Area	Item
Mutagenicity	Effect on primates
	Histopathological specimen preparation
	Reverse mutation
	Chromosomal aberration
	Micronucleus
	Gene mutation or DNA damage
Residues	Metabolism in plants
	Residues in crops
	Analysis methods on residues in crops
	Residues in processed commodities
	Metabolism in livestock
	Residues in livestock
	Analysis methods on residues in livestock
	Stability in stored commodities
Environmental	Aerobic flooding soil
behavior	Aerobic soil
	Anaerobic soil
	Hydrolysis
	Photolysis in water
	Soil adsorption
	Bioconcentration
Ecotoxicity	Fish acute toxicity
	Acute immobilization on daphnids
	Acute immobilization on daphnids (adults)
	Reproduction on daphnids
	Adverse effects of coexistent organic substances on fish acute toxicity and daphnids acute immobilization
	Acute toxicity on freshwater shrimps (Paratya compressa compressa and Paratya compressa improvisa) and amphipoda
	Acute immobilization on chironomid larvae
	Growth inhibition on algae and cyanobacteria
	Growth inhibition on Lemna sp.
	Acute oral toxicity on avian
	Acute dermal toxicity on honeybees (adults)
	Acute oral toxicity on honeybees (adults)
	Chronic oral toxicity on honeybees (adults)
	Oral toxicity on honeybees (larvae)
	Residues in pollen and nectar