Safety Data Sheet

Reference No. 1027

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1. Chemical product and company identification

Product name PACKTEST Silver Model WAK-Ag

Company name KYORITSU CHEMICAL-CHECK Lab., Corp.
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Dept. in charge Sales Department

Recommended uses and restrictions Reagent for water quality measurement

2. Hazards identification

[GHS Classification**]**

Physical hazards: Classification not possible (no data for GHS classification available)

Health hazards:

Specific target organ toxicity (repeated exposure): Category 2 (kidneys)

For those health hazards not listed above are not classified or classification not possible (no data for GHS classification available)

Environmental hazards:

Hazardous to the aquatic environment, short-term (acute): Category 3
Hazardous to the aquatic environment, long-term (chronic): Category 3
Harmful effects on the ozone layer: Classification not possible

[GHS labeling elements**]**

[Signal word**]**

Warning

[Hazard statements**]**

May cause damage to kidneys through prolonged or repeated exposure.
Harmful to aquatic life.
Harmful to aquatic life with long lasting effects.

[Precautionary statements**]**

Keep out of reach of children and store in the dry and dark place at room temperature.
Carefully read instructions before use and do not use for other purposes.
Wear personal protective equipment if necessary.
Do not inhale reagents.
Wash contaminated clothing.
Wash hands well before and after handling.
Avoid release to the environment.

12. Ecological information

No data on mixture is available. Data on each substance are shown.

Tetrabromofluorescein sodium, 1,10-Phenanthroline monohydrate and Polyethylene:

No eco-toxicological information available.

Ethylenediamine-N,N,N',N'-tetraacetic Acid Tetrasodium Salt Tetrahydrate
(data on Ethylenediaminetetraacetic acid (CAS 60-00-4) are shown) :

Hazardous to the aquatic environment (Acute): Category 3

It was classified in Category 3 from 96-hour LC50 = 41 mg/L for fish (Lepomis macrochirus) (EU-RAR, 2005, etc.).

Hazardous to the aquatic environment (Long-term): Category 3

"If chronic toxicity data are used, then it is classified as ""Not classified"" due to 21-day NOEC = 5.5 mg/L for crustacea (Daphnia magna) (Results of Aquatic Toxicity Tests of Chemicals conducted by Ministry of the Environment in Japan (Ministry of the Environment, 2002), etc.), although it is not rapidly degradable (a 4-week degradation rate by BOD: 0% (Biodegradation and Bioconcentration Results of Existing Chemical Substances under the Chemical Substances Control Law, 1994)).

If acute toxicity data are used for a trophic level for which chronic toxicity data are not obtained, then it is classified in Category 3 due to being not rapidly degradable (a 4-week degradation rate by BOD: 0% (Biodegradation and Bioconcentration Results of Existing Chemical Substances under the Chemical Substances Control Law, 1994)), and 96-hour LC50 = 41 mg/L for fish (Lepomis macrochirus) (EU-RAR, 2005, etc.).

By drawing a comparison between the above results, it was classified in Category 3. "

Hazardous to the ozone layer: Classification not possible

This substance is not listed in the Annexes to the Montreal Protocol.

GHS classifications as a mixture are shown below.

[Hazardous to the aquatic environment, short-term (acute)]

Classified as Category 3 (-, Harmful to aquatic life.) because more than or equal to 2.5% of Category 2 substance.

[Hazardous to the aquatic environment, long-term (chronic)]

Classified as Category 3 (-, Harmful to aquatic life with long lasting effects.) because more than or equal to 2.5% of Category 2 substance.

[Harmful effects on the ozone layer]:

Classification is not possible because each of the substances is not described in Annex to Montreal Protocol.

13. Disposal considerations

Always dispose of in accordance with local regulations.

14. Transport information

In addition to precautionary measures regarding handling and storage, avoid rough handling so as not to break containers. It is recommended to ship by air because under high temperature for long period may lead to deterioration.

UN classification and number:	Not applicable (this product contains less than or equal to 1% of 1,10-Phenanthroline monohydrate)
Civil Aeronautics Act:	Not regulated as per IATA
Fire Service Act:	Not applicable
Total weight of the product:	ca.140 g/kit

15. Regulatory information

Poisonous and Deleterious Substances Control Act: Not applicable

PRTR Act: Ethylenediamine-N,N,N',N'-tetraacetic Acid Tetrasodium Salt Tetrahydrate is applicable as “Class I Designated Chemical Substances No. 595 Ethylenediaminetetraacetic acid and its potassium and sodium salts.

Industrial Safety and Health Act: Not applicable

3. Composition/ information on ingredients

Discrimination of single substance or mixture: Mixture

Reagent name	K-1 reagent				
Chemical name	Tetrabromofluorescein sodium	1,10-Phenanthroline monohydrate (o-Phenanthroline)	Ethylenediamine-N,N,N',N'-tetraacetic Acid Tetrasodium Salt Tetrahydrate	Other (not regulated)	Polyethylene
Content	<0.01%	< 0.02%	<5%	<5%	>89.97%
Chemical formula	$C_{20}H_6Br_4Na_2O_5$	$C_{12}H_8N_2 \cdot H_2O$	$C_{10}H_{12}N_2Na_4O_8 \cdot 4H_2O$	-	$(C_2H_4)_n$
METI No. (reference number under CSCL in Japan)	(5)-1511	(5)-3915	(2)-1265 2-(4)-113	-	(6)-1
CAS No.	548-26-5	5144-89-8	13235-36-4	-	9002-88-4

4. First-aid measures

If reagents or test solutions;
- Enter in eyes: Immediately rinse eyes thoroughly.
- Contact with skin: Immediately wash out contaminated site with plenty of water.
- Enter into mouth: Immediately rinse mouth with plenty of water.

If ingested or in case any symptoms appear after above measures, immediately get medical advice or treatment.

5. Fire-fighting measures

Extinguishing methods: Cut off ignition sources and extinct by a suitable media.
Suitable extinguishing media: Water (mist), powder, carbon dioxide, dry sand.

6. Accidental release measures

In case of outdoor use: avoid spill of reagent or waste solution.
In case of indoor use: if spilled on a table or floor, wipe off immediately spilled reagent and dispose of them.

7. Handling and storage

Handling: Care should be made so that reagents and test solutions will not contact with eyes and skin and to avoid ingestion.
Especially for outdoor use, ensure to bring back reagents, waste solutions after the measurement and used containers.
Storage: Avoid direct sunlight and store in a well-ventilated, dry and dark place at room temperature.

8. Exposure controls and personal protection

Administrative control level
- Working environment standard: Not established

Occupational exposure limits
- Japan Society for Occupational health: Not established
- ACGIH (TLVs): Not established
- OSHA (PEL): Not established

Protective equipment: Recommended to wear protective glasses and gloves.

9. Physical and chemical properties

Physical state:	Tube containing powder reagent 1.1 g x 50 tubes/kit, aluminum laminate packaging each of 5 tubes
Color:	Light gray color (powder), semi-transparent (polyethylene tube)
Odor:	No odor
pH:	6

Melting point, boiling point, flash point, ignition point, lower explosion limit, vapor pressure, density, relative density, solubility, Pow, kinetic viscosity: not available as a mixture.

10. Stability and reactivity

Avoid leaving in a place where high temperature, humid or under direct sunlight. Stable under normal use conditions and no dangerous reactions under specific conditions are expected. No information on hazardous decomposition product is available.

11. Toxicological information

No data on mixture is available. Data on each substance are shown.

Tetrabromofluorescein sodium:
Acute toxicity: Oral-rat: LD_{50} = 2,344 mg/kg, Interperitoneal-rat LDL_0 = 500 mg/kg,
Intravenous injection-rabbit: LDL_0 = 300 mg/kg (RTECS)
Other data : Not available

1,10-Phenanthroline monohydrate (data on anhydride (CAS No.66-71-7) are shown):
Acute toxicity: Intravenous injection- mouse: LD_{50} = 18 mg/kg, Interperitoneal-moue: LD_{50} = 75 mg/kg
Other data : Not available

Ethylenediamine-N,N,N',N'-tetraacetic Acid Tetrasodium Salt Tetrahydrate
(data on Ethylenediaminetetraacetic acid (CAS 60-00-4) are shown) :
Acute toxicity (Oral) : Not classified
It was classified as "Not classified" in the Classification JIS (Category 5 in UN GHS classification) based on LD50 values for rats of > 2000 mg/kg (EU-RAR 49 (2004)), 2,580, 4,500 mg/kg (the above, Initial Risk Assessment Report Ver.1.1, 14 (NITE, CERI, NEDO, 2007)).
Acute toxicity (Dermal) : Classification not possible
No data available.
Acute toxicity (Inhalation: Gases) : Not applicable
Solid (GHS definition)
Acute toxicity (Inhalation: Vapours) : Classification not possible
No data available.
Acute toxicity (Inhalation: Dusts and mists) : Classification not possible
Classification not possible due to lack of data. Besides, it is reported that there were no dead animals after 8-hour inhalation of a dust atmosphere at 20 or 80 deg C (Initial Risk Assessment Report Ver.1.1, 14 (NITE, CERI, NEDO, 2007)).
Skin corrosion/irritation : Not classified
It was classified as "Not classified" because it is reported that it was not irritating in a Draize test with rabbits (Initial Risk Assessment Report Ver.1.1, 14 (NITE, CERI, NEDO, 2007)), and there was mild irritation at 24 hours in another test by 20-hour application to the ear of one rabbit (EU-RAR 49 (2004)).
Serious eye damage/eye irritation : Category 2B
It was classified in Category 2B because it is described that strong irritation, mild edema, and strong corneal opacity were seen in a test in which 50 mg was applied to the rabbit eye, but there were no findings at 8 days (EU-RAR 49 (2004)).
Respiratory sensitization : Classification not possible
No data available.
Skin sensitization : Classification not possible
No data available. Besides, for the disodium salt of this substance, it is reported that in a guinea pigs maximization test (OECD TG406), a positive rate was 30% (3/10) in the first challenge after 24 hours and 10% (1/10) in a second challenge after 7 days (EU-RAR 49 (2004)), and the substance was assessed as not sensitizing in another guinea pig maximization test (Initial Risk Assessment Report Ver.1.1, 14 (NITE, CERI, NEDO, 2007)).
Germ cell mutagenicity : Not classified

As for in vivo tests on the disodium salt of this substance, negative results were obtained in all of a dominant lethal test by drinking water administration to mice (in vivo heritable germ cell mutagenicity test), a chromosomal aberration test with spermatogonial cells after intraperitoneal administration to mice (in vivo heritable germ cell mutagenicity test), and micronucleus tests with bone marrow after oral or intraperitoneal administration to mice (in vivo somatic cell mutagenicity tests) (all, EU-RAR 49 (2004)). Therefore, this substance was classified as "Not classified." Also, as for in vitro tests, this substance was negative in an Ames test and positive in a mouse lymphoma test, and there were negative results in an Ames test and a mouse lymphoma test on the trisodium salt of this substance (EU-RAR 49 (2004), Mutagenicity Test Data of Existing Chemical Substances based on the toxicity investigation system of the Industrial Safety and Health Law Supplement 2 (2000)). Besides, this substance was reported to be positive in a chromosomal aberration test with bone marrow and spleen cells of mice (in vivo somatic cell mutagenicity test) (Initial Risk Assessment Report Ver.1.1, 14 (NITE, CERI, NEDO, 2007)), but it was not used for classification based on expert judgment, claiming that test details such as administration route and doses are unknown, and the reproducibility of results is questionable for the test.

Carcinogenicity : Classification not possible

No data available. Besides, in 103-week diet administration tests with rats and mice on the trisodium salt of this substance, there were no occurrences of treatment-related tumors in rats or mice, but it is mentioned that the substance was not tested at the maximum tolerated dose (EU-RAR 49 (2004)).

Reproductive toxicity : Category 2

It is reported that after gavage administration to rats on gestational days 7-14, there were no effects in fetuses at the dose where effects such as death, diarrhea, and behavioral suppression were seen in parent animals (Initial Risk Assessment Report Ver.1.1, 14 (NITE, CERI, NEDO, 2007)), while it is reported that cleft palate, brain and eye defects, and skeletal anomalies occurred in fetuses in a test by diet administration to rats after day 6 of gestation (Teratogenic (12th, 2007)), although there is no description of general toxicity in parent animals, and incidences of malformations in fetuses were also reported after intraperitoneal or intramuscular administration to pregnant rats (Initial Risk Assessment Report Ver.1.1, 14 (NITE, CERI, NEDO, 2007), JECFA 796 (1993)). Therefore, it was classified in Category 2.

Specific target organ toxicity - Single exposure : Classification not possible

No data available. Besides, as for effects of the related substance in humans, as acute symptoms when the disodium salt of EDTA (Na2EDTA) was intravenously administrated as an antidote for lead poisoning, numbness and tingling pain, arising around the hands and mouth, were reported (Initial Risk Assessment Report Ver.1.1, 14 (NITE, CERI, NEDO, 2007)).

Specific target organ toxicity - Repeated exposure : Category 1 (kidney)

As effects in humans, it is described that renal tubular defects were seen when EDTA or its salts (sodium, calcium disodium) were ingested for a long time and in a large amount (Initial Risk Assessment Report Ver.1.1, 14 (NITE, CERI, NEDO, 2007)). Therefore, it was classified in Category 1 (kidney). Besides, the formulation of CaNa2EDTA, the related substance, is commercially sold as an antidote for lead poisoning, side effect information includes renal tubular defects after long-term administration of tablets, transient proteinuria for infusion, and renal tubular defects after long-term infusion, and it is described in other remarks that acute and large-amount administration may cause serious consequences such as death due to renal toxicity (Environmental Risk Assessment for Chemical Substances vol. 3 (Ministry of the Environment, 2004)).

Aspiration hazard : Classification not possible

No data available.

Polyethylene:

Acute toxicity:

Oral: Rat LD_{50} > 7,950 mg/kg (used 7,950 mg/kg for the calculation of ATEmix below)

Carcinogenicity: IARC Group 3 (not classifiable as to carcinogenicity to humans).

Other data: Not available

GHS classifications as a mixture are shown below.

[Specific target organ toxicity (repeated exposure)]

Classified as Category 2 (Warning, May cause damage to kidneys through prolonged or repeated exposure.) because 1 to 10% of category 1 substance is contained.

[Acute toxicity], [Skin corrosion/ irritation], [Serious eye damage/ eye irritation], [Respiratory or skin sensitization], [Germ cell mutagenicity], [Carcinogenicity], [Reproductive toxicity], [Specific target organ toxicity (single exposure)], [Aspiration hazard]

Not classified or classifications are not possible because of lack of data.

16. Other information

Reference literature

15,911 Kagaku Shouhin, The Chemical Diary Co., Ltd. (2011)
Safety Data Sheet No. JW050672, Wako Pure Chemical Corporation. (2007.09.07)
Safety Data Sheet No. JW160086, Wako Pure Chemical Corporation. (2008.08.27)
NITE, GHS Classification, ID m-nite-60-00-4_v1, Ethylenediaminetetraacetic acid
Material Safety Data Sheet No.051110033, TOSOH CORPORATION (2004.07.09)
Koukuu Kikenbutsu Yusou Houreisyu, Ed. MLIT, HOUBUN SHORIN CO., LTD. (2019)
JIS Z 7252:2019 Classification of chemicals based on “Globally Harmonized System of Classification and Labelling of Chemicals (GHS)” (Japanese Industrial Standards Committee)
JIS Z 7253:2019 Hazard communication of chemicals based on GHS-Labelling and Safety Data Sheet (SDS) (Japanese Industrial Standards Committee)
UN GHS (tentative translation, forth revised version), GHS Kankei Syocho Renraku Kaigi (2011)
Ministry of Economy, Trade and Industry, GHS Classification Guidance for Enterprises 2013 Revised Edition (2013)

NOTE) This information is not always exhaustive and use with care.
This data sheet only provides information but any description cannot be warranted.
Descriptions may possibly be changed because of new findings or modification of the current knowledge.
Precautions only cover normal handling.
This English SDS is prepared in the cooperation with the Chemicals Evaluation and Research Institute (CERI), Japan.