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OLD Diagnostika	GmbH		SAFETY DATA				
SECTION 1: Identifica	tion of the	substance/mixture and of the com	pany/undertaking				
Catalogue no: RA	 1 Product identifier MuSK Antibody RIA (RiaRSR[™] MuSK Ab) Catalogue no: RA120/25 (REF) MK/25) 2 Relevant identified uses of the substance or mixture and uses advised against: 						
	mination o	of autoantibodies to muscle specific					
DLD Diagnostika Adlerhorst 15 22459 HAMBURO	GmbH G, GERMA 58710; Fax	:: +494055587111					
1.4 Emergency telep +49(0)40-555871		n ber: Fri, except public holidays, 8.00 – 15	5.30)				
SECTION 2: Hazards	s identifica	ation					
2.1 Classification of Classification ac		tance or mixture o Regulation (EC) No. 1272/2008 [_				
Kit Compo	nent	Hazard Classification	Hazard Statements*				
Precipitation Enh		Specific target organ toxicity – repeated exposure, Category 2	H373				
*See section 16 for ful 2.2 Label elements Labelling accord		gulation (EC) No. 1272/2008 [CLP]:				
PRECIPITATIO		CER					
Hazard pictogram		Signal word: Warning					
Hazard stateme	• •						
H373	exposure.		nged or repeated				
Precautionary s							
P260		eathe dust/fume/gas/mist/vapours/s					
P314		cal advice/attention if you feel unwel					
F301	P501 Dispose of contents/container to hazardous or special waste collection point, in accordance with local, regional, national and/or international regulation						
	nponents	not listed in section 2.1 and 2.2 oncentrations which meet the criteria					
te: 22 nd May 2023		MuSK Antibody RIA (RiaRSR [™] MuSK Ab					

according to Regulation (EC) No. 1272/2008. However, ingestion or exposure to large amounts from improper handling can be potentially hazardous.

This kit contains both animal and human proteins and should be treated as a potential biohazard. All animal and human sera have been tested to ensure the absence of infectious agents but all materials should be handled as though capable of transmitting infectious disease and disposed of accordingly.

This kit contains ¹²⁵Iodine, a radioisotope with a half-life of approximately 60 days which emits gamma radiation with a maximum energy of 35 keV. Evidence exists of mutagenic, teratogenic and carcinogenic effects by ionising radiation.

MuSK Antibody RIA kit components ingredients listed in 3.2 have not been identified as having endocrine disrupting properties according to Regulation (EU) 2017/2100 and does not meet the criteria for vPvB and PBT according to Regulation (EC) No. 1907/2006 Annex XIII.

The following precautionary phrases should be taken into consideration: P233, P260, P270, P281, P301 + P330 + P331, P302 + P352, P304 + P340, P305 + P351 + P338, P314 (see section 16 for full text).

SECTION 3: Composition/information on ingredients

3.1 Substances

Not applicable

3.2 Mixtures

Hazardous ingredients according to Regulation (EC) No. 1272/2008:

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Ingredient(s)	CAS No.	EC No.	Classification (GHS)	Conc. (v/v)	Conc. Limits
Silica (silicon dioxide)	14808-60-7	238-878-4	STOT RE 1; H372	1 – 10%	STOT RE 1 C≥10% STOT RE 2 1≤C<10%

The following kit components contain ingredients which are considered hazardous but are not present in high enough concentrations to be classified under Regulation (EC) No. 1272/2008.

Component(s)	Ingredient	Number	Classification (GHS)	Conc. (v/v)	Conc. Limits (v/v)
Reconstitution Buffer Precipitation Enhancer Anti Human IgG Wash Solution Controls	Sodium Azide	CAS No. 26628-22-8 EC No. 247-852-1	Acute Tox. 2 (Oral & Inhalation), Acute Tox. 1 (Dermal), STOT RE 2, Aquatic Acute 1, Aquatic Chronic 1; H300, H310, H330, H373, H400, H410, EUH032	<0.1%	Acute Tox. 2 (Oral & Inhalation) $C \ge 0.1\%$ Acute Tox. 1 (Dermal) $C \ge 0.1\%$ STOT RE 2 $C \ge 10\%$ Aquatic Acute 1 $C \ge 0.1\%$ Aquatic Chronic 1 $C \ge 0.1\%$

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¹²⁵I-labelled MuSK tracer, anti-human IgG, positive and negative controls contain animal proteins and/or human proteins and should be treated as potential biohazards.

 125 l-labelled MuSK is radioactive, ~30 kBq per vial (~0.811 $\mu Ci).$

The full text for the hazard statements can be found in section 16.

SECTION 4: First aid measures

4.1 Description of first aid measures

After skin contact

Wash off skin thoroughly with water for at least 15 minutes. Remove contaminated clothing. In severe cases or if skin is broken, OBTAIN MEDICAL ATTENTION.

After eye contact

Separate eyelids with fingers and flush eye with copious amounts of water for at least 15 minutes. OBTAIN MEDICAL ATTENTION.

After Inhalation

Remove from exposure, rest and keep warm. If breathing becomes difficult, OBTAIN MEDICAL ATTENTION.

After Ingestion

If patient is conscious, wash out mouth with water and give plenty of water to drink. OBTAIN MEDICAL ATTENTION.

4.2 Most important symptoms and effects, both acute and delayed Not available.

4.3 Indication of any immediate medical attention and special treatment needed

Not available.

SECTION 5: Firefighting measures

5.1 Extinguishing media

Use water, dry powder or foam as appropriate to supporting fire.

5.2 Special hazards arising from the substance or mixture

May evolve toxic fumes in fire. Hazardous combustion products are not known for kit components but combustion products for the ingredients listed in subsection 3.2 can be found in the following table:

Ingredient	Hazardous combustion product(s)	
Silica	None	
Sodium Azide	Nitrogen oxides (NOx)	

5.3 Advice for firefighters

Wear self-contained breathing apparatus and protective clothing to prevent contact with skin and eyes.

6.1 Personal precautions, protective equipment and emergency procedures Wear appropriate protective clothing as described in subsection 8.2. Ventilate area and avoid breathing vapours, mist or gas.

6.2 Environmental precautions

Prevent further leakage or spillage if safe to do so. Prevent any reagents from entering drains.

6.3 Methods and material for containment and cleaning up

Radioactive spills should be dealt with immediately in accordance with the current local and national regulations and guidelines.

Wipe up liquid spills with absorbent paper. For solid spills, sweep up without raising dust. Once pick up is complete, wash site with detergent and water and decontaminate with a suitable disinfectant solution. Any surfaces contaminated with ¹²⁵lodine should be washed with a suitable detergent to remove all traces of radioactivity. Dispose of radioactive waste via an authorised route.

6.4 Reference to other sections

See sections 8 and 13.

SECTION 7: Handling and storage

7.1 Precautions for safe handling

Users should make themselves aware of, and observe any national and local legislation and codes of practice governing the use, storage, transportation and disposal of radioactive materials.

Radioactive materials should only be used by authorised personnel and in designated areas. Wash hands thoroughly after handling. Monitor hands and clothing before leaving the designated area. Report contamination to the responsible person and take remedial action.

Material of human origin has been tested and found non-reactive for HIV 1 and 2 and HCV antibodies and HBsAg. All animal sourced material has been obtained from animals certified as healthy and free from disease. However all potentially biohazardous components should be considered as potentially infectious. Level 2 containment should be applied.

Do not eat, drink or smoke in the laboratory. Do not pipette by mouth. Avoid skin and eye contact. Wear appropriate protective clothing as described in subsection 8.2. Avoid the use of needles or other sharp implements. Avoid prolonged or repeated exposure. Wash hands thoroughly after handling. Avoid release into drains; in case of accidental spillage, refer to section 6.

7.2 Conditions for safe storage, including any incompatibilities

Keep containers tightly closed. Store in a dry place in the box supplied at a temperature between +2 and +8°C.

7.3 Specific end use(s)

The MuSK Antibody RIA Kit is intended for professional use only and to be used solely for the purpose as specified in subsection 1.2. Refer to kit instructions for details.

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SECTION 8: Exposure controls/personal protection

8.1 Control parameters

No occupational exposure limits exist for any kit components. However, exposure limits apply to the following ingredients (see subsection 3.2 for components containing these substances):

Value	Control	Basis	
	Parameters		
Silica			
TWA	0.1 mg/m ³	UK: EH40 Workplace Exposure Limits (WEL)	
Sodium Az	ide		
TWA	0.1 mg/m ³	UK: EH40 Workplace Exposure Limits (WEL)	
STEL	0.3 mg/m ³	Europe: Commission Directive 2000/39/EC	
8.2 Exposur	3.2 Exposure controls		

Annropriate engineering cor

Appropriate engineering controls

Good laboratory practice should be followed (see Section 7). Avoid contact with skin or eyes. Wash hands after use.

Individual protection measures (personal protective equipment)

Eye/face protection

Chemical safety glasses or goggles conforming to appropriate government standards such as EN166 (EU) or NIOSH (US).

Skin and body protection

Chemical resistant gloves to be used in accordance with standard EN374 derived from Regulation (EU) 2016/425. Inspect gloves for damage prior to use and change if any sign of degradation. Proper glove removal technique must be used. Wash hands after use.

The following are suitable as protective gloves:

Glove materials: Nitrile rubber Glove Thickness: >= 0.4 mm thickness Permeation Time: >= 480 min

This recommendation is advisory only and should be evaluated by the customer for suitability in their specific situation.

Respiratory protection Local exhaust.

Environmental exposure controls

Prevent further leakage or spillage if safe to do so. Prevent any reagents from entering drains.

SECTION 9: Physical and chemical properties

9.1 Information on basic physical and chemical properties

Kit component	Appearance	Odour	рН	Solubility
¹²⁵ I-labelled MuSK Tracer	White solid	None	N/A	In water

Reconstitution Buffer	Colourless liquid	None	~7.6	N/A
Precipitation Enhancer	White suspension	None	N/A	N/A
Anti-human IgG	Yellow/brown liquid	None	N/A	N/A
Wash Solution	Colourless liquid	None	~7.6	N/A
Positive and Negative Controls	Yellow/brown liquid	None	N/A	N/A

There is no information available for the following categories: odour threshold, melting/freezing point, initial boiling point/boiling range, flash point, evaporation rate, flammability (solid, gas), upper/lower flammability or explosive limits, vapour pressure, relative vapour density, relative density, particle characteristics, partition coefficient, autoignition temperature, decomposition temperature, kinematic viscosity, explosive properties or oxidising properties.

9.2 Other information

All liquid components are miscible with water in all proportions.

SECTION 10: Stability and reactivity

10.1 Reactivity

Data is not available on the reactivity of individual kit components but is given, where available, on ingredients listed in subsection 3.2.

10.2 Chemical stability

All components of the MuSK Antibody RIA Kit have been found stable for stated shelf life when stored under the recommended conditions.

10.3 Possibility of hazardous reactions

No hazardous reactions known for kit components although, hazardous reactions occur for the following ingredients listed in subsection 3.2:

Ingredient	Hazardous Reaction
Sodium Azide	Risk of explosion and/or toxic gas formation exists with heavy metals, bromine, lead, chromyl chloride, dichloromethane, dimethylsulfate, halogenated hydrocarbon, acid, carbon disulphide, sulphuric acid, copper and nitric acid. Generates dangerous gases or fumes with acids and water, leading to the release of hydrazoic acid. Violent reactions possible with nitrates, benzoyl chloride and potassium nitrate.

10.4 Conditions to avoid

Proteins and sodium azide are heat sensitive and storage or use at the improper temperature may compromise the integrity of the kit.

10.5 Incompatible materials

No data is known for kit components but the following data is known for ingredients listed in subsection 3.2:

Ingredient	Incompatible materials
Silica	Reacts with hydrogen fluoride.
Sodium Azide	Aluminium and heavy metals.

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10.6 Hazardous decomposition products

No decomposition products are formed if kit is stored and used under the specified storage and handling conditions.

May evolve toxic fumes in fire. Thermal decomposition products are not known for the kit components but hazardous combustion products of the ingredients listed in subsection 3.2 can be found in subsection 5.2

SECTION 11: Toxicological information

11.1 Information on toxicological effects

The kit components have not been directly tested for their toxicological effects, therefore no information is known for these mixtures. The following toxicological data is known for ingredients listed in subsection 3.2:

_	-		+~~
	а	ACULE	toxicity
	-		

*Definitions can be found in section 16

Ingredient	Measurement*	Value	Species
Sodium Azide	LD ₅₀ (Oral)	27 mg/kg	Rat
	LC ₅₀ (Inhalation)	0.054 – 0.52 mg/L (4h)	Rat
	LD ₅₀ (Dermal)	20 mg/kg	Rabbit

No data available for other ingredients listed in subsection 3.2.

(b) Skin corrosion/irritation

Ingredient	Test/Result
Sodium Azide	In vitro study, human skin model test – No skin irritation

No data available for other ingredients listed in subsection 3.2.

(c) Serious eve damage/irritation

Ingredient	Test/Result			
Sodium Azide	Bovine cornea, exposure time 4 hours – No eye irritation			
New data wave lightly fair athen is one diants. Fate discussion attack 0.0				

No data available for other ingredients listed in subsection 3.2.

(d) Respiratory or skin sensitisation

Ingredient	Test/Result
Sodium Azide	Sensitisation test, (dermal).Local lymph node assay (LLNA)
	- Mouse – Result: Negative

No data available for other ingredients listed in subsection 3.2.

(e) Germ cell mutagenicity

Ingredient	Test/Result
Sodium Azide	Chromosome aberration: Chinese hamster ovary cells – Negative
	Unscheduled DNA Synthesis assay: Chinese hamster lung cells – Negative Sister Chromatid exchange assay: Chinese hamster ovary cells – Negative

No data available for other ingredients listed in subsection 3.2.

(f) Carcinogenicity

No data available for ingredients listed in subsection 3.2.

(g) Reproductive toxicity No data available for ingredients listed in subsection 3.2.

(h) STOT-single exposure

No data available for ingredients listed in subsection 3.2.

(i) STOT-repeated exposure

Ingredient	Exposure/Result/Target Organ				
Sodium Azide	Oral - may cause damage to organs through repeated				
	exposure - Brain				

No data available for other ingredients listed in subsection 3.2.

(i) Aspiration hazard

No data available for ingredients listed in subsection 3.2.

11.2 Information on other hazards

(a) Endocrine disrupting properties

The substance/mixture does not contain components considered to have endocrine disrupting properties according to Commission Regulations (EU) 2017/2100 and (EU) 2018/605.

(b) Other information

As the kit components have not been tested for their toxicological effects, other hazardous properties cannot be excluded but are unlikely when the product is handled appropriately.

SECTION 12: Ecological information

The kit components have not been tested for their ecological effects, therefore no information is known for these mixtures. The following ecological data is known for ingredients listed in subsection 3.2:

12.1	Toxicity		*Definitions c	an be found in section 16
	Ingredient	Toxicity to	Measurement*	Value
	Sodium Azide	Fish (Oncorhynchus mykiss	LC ₅₀	2.75 mg/L (96h)
		(rainbow trout) Algae (<i>Psuedokirchneriella</i> subcapita)	ErC ₅₀	0.35 mg/L (96h)

No data available for other ingredients listed in subsection 3.2.

12.2 Persistence and degradability

No data available for ingredients listed in subsection 3.2.

12.3 Bioaccumulative potential

No data available for ingredients listed in subsection 3.2.

12.4 Mobility in soil

No data available ingredients listed in subsection 3.2.

12.5 Results of PBT and vPvB assessment				
	Ingredient Test/Result			
		This substance/mixture contains no components considered to be either persistent, bioaccumulative and toxic (PBT), or very persistent and very bioaccumulative (vPvB) at levels of ≥0.1%.		

No data available for other ingredients listed in subsection 3.2.

12.6 Endocrine disrupting properties

The ingredients listed in subsection 3.2 do not have endocrine disrupting properties with respect to non-target organisms as it does not meet the criteria set out in section B of Regulation (EU) No 2017/2100.

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12.7 Other adverse effects The concentrations of ingredients listed in subsection acceptable limit for hazardous substances; the ecologi However, it is recommended that reagents do not er quantities.	cal risk is minimal.	(listed in subsec P233: Keep cont P260: Do not bre P270: Do not ea	tion 3.2) according tainer tightly closec eathe dust/fume/ga t, drink or smoke w	s/mist/vapours/spray /hen using this product.
SECTION 13: Disposal considerations		P281: Use perso P301 + P330 + F	P331: IF SWALLOV	pment as required. VED: rinse mouth. Do NOT induce vomiting.
13.1 Waste treatment methods Chemical and biological residues are classified as special are covered by regulations which may vary according to lo local waste disposal authority for advice or pass to company. Observe all national and local environmental reg Contaminated packaging should be disposed of using the	ocation. Contact your a licensed disposal gulations.	P302 + P352: IF P304 + P340: IF comfortable for b P305 + P351 + Remove contact P314: Get medic	ON SKIN: Wash w FINHALED: Remove preathing. P338: IF IN EYES lenses, if present a cal advice/attention	vith plenty of soap and water. ve victim to fresh air and keep at rest in a position : Rinse cautiously with water for several minutes. and easy to do so. Continue rinsing.
SECTION 14: Transport information		in accordance with local, regional, national and/or international regulation. H300: Fatal if swallowed.		
This product is not covered by international regulation on the tra goods (IMDG, IATA, ADR/RID). Transport of this product can be carried out at ambient tempera	ture but in the event	H310: Fatal in co H330: Fatal if inf	ontact with skin. naled.	lungs) through prolonged or repeated exposure if
of delays store at 2 – 8°C with all reagents contained within the p 14.1 UN number or ID number Not applicable.	ackaging provided.	 inhaled. H373: May cause damage to organs through prolonged or repeated exposure. H400: Very toxic to aquatic life. H410: Very toxic to aquatic life with long lasting effects. EUH032: Contact with acids liberates toxic gas. Definitions: LD50 = Lethal dose for 50% of the test population. LC50 = The lethal concentration of a substance that kills 50% of the test population within a designated period. ErC50 = The effective concentration of a substance that causes adverse effects in 50% of the test population within a designated period. STEL = Short term exposure limit (15 minute reference period). 		
14.2 UN proper shipping name Not applicable.				
14.3 Transport hazard class(es) Not applicable.				
14.4 Packing group Not applicable.				
14.5 Environmental hazards Not applicable.				
14.6 Special precautions for user See sections 6 to 8.				
14.7 Maritime transport in bulk according to IMO instruments Not applicable.	3			ng term exposure limit (8 hour reference period). d to be correct but does not purport to be all-
SECTION 15: Regulatory information		inclusive and is	provided for guida	ance only. DLD Diagnostika GmbH shall not be
15.1 Safety, health and environmental regulations/legislati substance or mixture. Not applicable	on specific for the	 held liable for any damage or injury resulting from handling or from contact with the above product and assumes no responsibility to the accuracy or completeness of the data contained herein. It is the responsibility of the purchaser to ensure the laboratory workers who use this product are aware of its hazards and take necessary precautions to prevent contact, ingestion, inhalation or any other mode exposure. REVISION INFORMATION 		
15.2 Chemical safety assessment No Chemical Safety Assessment has been carried out for RIA kit by the manufacturer.	the MuSK Antibody			
SECTION 16: Other information		Revision	Effective	
This SDS has been compiled in accordance with Commission		Number	Date	Description of Changes
1907/2006 as amended by Commission Regulation (EU) 2020/8 All information provided on ingredients listed in subsection 3.2 from the appropriate chemical safety data sheets.		1	22 nd May 2023	Revision of SDS to meet (EU) 2020/878 – changes throughout.
Date: <u>22nd May 2023</u> MuSK Antibody RIA (RiaRSR [™] MuSK		sds/30	a Rev. 1 Supersedes R	lev 0 Page 5 of 5