

SAFETY DATA SHEET

Prekallikrein Activator Assay Ig Kit

Research Use Only. Not for diagnostic procedures.

1. PRODUCT AND COMPANY IDENTIFICATION

- 1.1 Product name:
- 1.1 Code:
- 1.2 Recommended use:
- 1.3 Manufacturer:
- Prekallikrein Activator (PKA) Assay Ig Kit PW302EP In-vitro research test kit Pathway Diagnostics Ltd Eclipse House, Curtis Road Dorking, RH4 1EJ, UK

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2. COMPOSITION INFORMATION

2.1 Kit contents:

- Substrate PW2302, 2x 5ml
- Human Prekallikrein, 2x 2.5 ml
- PKA Standard, 1x1ml 32IU
- PKA Buffer, 1x 6.0ml
- Sample/Standard Buffer, 2x 6 ml
- Immunoglobulin Pre-Treatment Reagent 1x 1ml Buffered solution pH 7.9
- Blank Activity Blocking Reagent 1x 1ml

Chromogenic substrate Human plasma fraction Human plasma Buffered salt solution, pH 7.9. Bovine Albumin Buffered solution pH 7.9 Plant based enzyme Inhibitors

3. HAZARDS IDENTIFICATION

3.1 Human Prekallikrein, PKA standard: Products derived from human plasma. Handle with care, observing the precautions recommended for bio-hazardous material. The human plasma used has tested negative for antibodies to HIV Type 1 and 2, Hepatitis C and Hepatitis B surface antigen. However, since no test method can offer complete assurance that infectious agents are absent these human plasma should always be handled with care, observing the precautions recommended for a bio-hazardous material.

- **3.2 Sample Buffer bovine albumin:** This kit contains material of animal origin and could be potentially capable of transmitting infectious diseases.
- **3.3 Emergency overview:** General rules for handling biological material have to be followed
- **3.4 Health hazard:** Minimal health hazard upon ingestion or risk of combustion.

4. FIRST AID MEASURES

4.1 Skin Contact:	If a component of this kit contacts the skin and causes discomfort, remove any contaminated clothing. Wash affected area with plenty of soap and water. If pain or irritation occurs, obtain medical attention.
4.2 Eye Contact:	If a component of this kit enters the eyes and causes discomfort, wash eyes gently under potable running water for 15 minutes or longer, making sure that the eyelids are held open. If pain or irritation occurs, obtain medical attention.
4.3 Inhalation:	Inhalation of any component in this kit is unlikely. If a component of this kit is inhaled and causes discomfort, move exposed individual to fresh air. Seek medical attention if breathing is difficult or symptoms persist. Seek medical advice immediately.
4.4 Ingestion:	If a component of this kit is ingested, wash mouth out with water. If irritation or discomfort occurs, obtain medical attention.
4.5 Other:	In case of injury and contamination with human plasma material it is strongly recommended to obtain medical attention.



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5. FIRE FIGHTING MEASURES

5.1 Extinguishing media:	No limitation. For small fires, use dry chemical, carbon dioxide, or
	alcohol-resistant foam.
5.2 Flammable properties:	None

6. ACCIDENTAL RELEASE MEASURES

6.1 Personal precautions: This kit contains materials of biological origin. Wear gloves to avoid personal contact. Use Universal Precautions during clean-up procedures
6.2 Clean up procedures: Absorb liquid and place in container suitable for disposal. Dispose of in accordance with applicable local procedures.

7. HANDLING AND STORAGE

- **7.1 Handling:** For Professional use only. Wash thoroughly after handling components of this kit. Do not eat or drink while using this kit. This kit should be handled only by qualified clinical or laboratory staff trained on the use of this kit and who are familiar with the potential hazards. This kit should be handled as though capable of transmitting infectious diseases. Universal Precautions should be followed when using this kit.
- 7.2 Storage:Store at +2°C to +8°C

8. EXPOSURE CONTROLS AND PERSONAL PROTECTION

8.1 Ventilation:	No particular ventilation requirements are necessary.	
8.2 Respiratory protection:	None needed under normal conditions of use.	
8.3 Hand protection:	Impervious gloves should be worn to prevent hand contact.	
8.4 Eye protection:	Safety glasses are recommended to prevent eye contact.	

9. Physical and Chemical Properties

9.1 Appearance:	Solid
9.2 Colour:	Colourless
9.3 Odour:	Odourless
9.4 Solubility:	Miscible with water
9.5 Viscosity:	No data available
9.6 pH-value:	pH 7.4 – pH 8.3

10. STABILITY AND REACTIVITY

10.2. Materials to avoid:	None known
10.2. Hazardous reactions:	None known if handled correctly
10.3. Hazardous	
decomposition:	Thermal decomposition may release irritating fumes or toxic gases

11. TOXICOLOGICAL INFORMATION

11.1. Acute toxicity: Not known



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11.2. Skin effects: 11.3. Eye effects: 11.4. Other:

Not known No cases of intoxication known so far. Human Plasma used for the production of the test kit components is derived exclusively from donors who have been tested negative for antibodies against HIV 1/2, HCV and HBs antigen using approved test procedures. Nevertheless human plasma should be considered as potentially infectious and handled with care. The bovine albumin used is of animal origin and could be potentially capable of transmitting infectious diseases.

12. ECOLOGICAL INFORMATION

12.1. Ecotoxicity:

No adverse effects on the environment are expected from the components of this kit. There is no aquatic toxicity data for any component of this product at this time.

13. DISPOSAL CONSIDERATIONS

13.1. Product disposal:	Incineration according to local and national legislation.
13.2. Packaging disposal:	Dispose of in a manner in accordance with local and national
	regulations.

Not known

14. TRANSPORT INFORMATION

- 14.1. UN. No.:
- 14.2. Road and rail transport (ADR/RID):
- 14.3. Air transport (ICAO/IATA-DGR):
- 14.4. Sea transport (IMO/IMDG):
- 14.5. Inland navigation (AND(ADNR):
- 14.6. Other information:
- Not applicable Not applicable Not applicable Not applicable None

Not applicable

15. REGULATORY INFORMATION

- **15.1:** Not classified as dangerous acc. to European Community Directive 93/21/EEC
- **15.2:** National legislation: For further product information, we recommend consideration of the corresponding appropriate National Legislation.

16. OTHER INFORMATION

- **16.1:** This Safety Data Sheet complies with the European Community Directive 93/112/EC amended 91/155/EC.
- 16.2: All information and instructions provided in this Safety Data Sheet are based on the current state of scientific and technical knowledge at the date indicated on this Safety Data Sheet. Pathway Diagnostics Ltd. shall not be held responsible for any defect in the product covered by this Safety Data Sheet, should the existence of such a defect not be detectable considering the current state of scientific and technical knowledge. It characterises the product with regard to the appropriate safety precautions. It does not represent a guarantee of the properties of the product.

Prepared by:	Pathway Diagnostics Ltd Eclipse House, Curtis Road Dorking, Surrey RH4 1EJ, UK	Date of Preparation: 20 January 2017
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