



Ludger Document # LC-EXO-A6-Guide v1.0

Ludger Ltd

Culham Science Centre
Oxford OX14 3EB
United Kingdom

Tel: +44 1865 408 554

Fax: +44 870 163 4620

Email: info@ludger.com

www.ludger.com



Contents

	Page
Contents	2
Ludger Post-Exoglycosidase Clean-up Spin Columns	3
Kit Contents	4
Additional Reagents and Equipment Required	4
Safety and Handling	4
Clean-up Procedure	4
Time Line for Procedure	4
Method	5
Apply the samples onto the clean-up column	5
Wash the wells of the clean-up plate	5
Dry the glycans if necessary	5
Warranties and Liabilities	5
Document Revision Number	5
Appendix 1: Troubleshooting Guide	6
Appendix 2: Material Safety Data Sheet	7



Ludger Post-Exoglycosidase Clean-up Spin Columns

Applications Post-exoglycosidase Clean-up

For removal of exoglycosidase enzymes and other protein material following glycan enzymatic digestion/ sequencing. This will prevent contamination of HPLC columns during subsequent chromatographic analysis. The spin columns can also be used to remove exoglycosidases or other proteins before mass spectrometry analysis of glycans.

Description The LC-EXO-A6 spin columns contain a specialized modified polyethersulfone

membrane with a molecular weight cut-off of approximately 30 kDa. This product is compatible with any centrifuge equipped with a rotor suitable for Eppendorf tubes. Glycans pass through the membrane whilst proteins are retained on the membrane

allowing separation of these two components.

Number of Samples Each pack contains 6 spin columns

Amount of Sample Up to 500 µL per column

Centrifugal Force Up to 14,000 x g

Suitable Samples Unlabelled and fluorophore labelled (e.g. 2-AB, 2-AA or procainamide labelled) glycans

released from glycoproteins or other sources and treated with exoglycosidase

enzymes.

Storage Store at room temperature. Protect from sources of heat, light, and moisture. When

stored correctly, the products should be stable for 36 months from date of purchase.

Shipping The product should be shipped at ambient temperature.

For research use only. Not for human or drug use



Kit Contents



The kit contains the following materials:

Cat. # Item Quantity

LC-EXO-A6 LudgerClean Post-Exoglycosidase Clean-up Spin Columns

With Sample Collection Vials 6

Additional Reagents and Equipment Required

- Pure water: resistivity above 18 MΩ-cm, particle free (>0.22 μm), TOC <10 ppb
- Centrifuge equipped with a rotor suitable for Eppendorf tubes, operating speed up to 14,000 x g

Safety and Handling

- Ensure that any glass, plasticware or solvents used with this item are free of environmental carbohydrates. Use powder-free gloves for all sample handling procedures and avoid contamination with environmental carbohydrate.
- Once used, the plate should be discarded according to local safety rules.

Clean-up Procedure

Time Line for Procedure

ProcedureTimeSample loading5 minutesElution of glycans10 minutesDrying glycansas requiredTotal time15 minutes plus drying time



Method

Apply the samples onto the spin column

- Make sure that the post-exoglycosidase clean-up spin column is securely assembled inside of the sample collection vial provided.
- Pipette the glycan sample onto the spin column membrane. Wash out each sample vial with 100 µL of water and add this to the spin column membrane. Close the vial and centrifuge for 3 minutes at 10,000 x g or until the liquid has all gone through the membrane.

The maximum centrifugal force used should not exceed 14,000 x g.

Wash the membrane of the spin column

• Pipette 100 μL of water directly onto the spin column membrane to wash through any remaining sample. Close the vial and centrifuge for 3 minutes at 10,000 x g or until the liquid has all gone through the membrane.

Dry the glycans if necessary

• At this stage glycan samples may be at sufficient concentration for their intended use. Alternatively you additionally concentrate the glycans in a vacuum centrifuge. We do not recommend applying heat at this stage. Long drying times under elevated temperature may lead to glycan desialylation.

Warranties and Liabilities

Ludger warrants that the above product conforms to the attached analytical documents. Should the product fail for reasons other than through misuse Ludger will, at its option, replace free of charge or refund the purchase price. This warranty is exclusive and Ludger makes no other warrants, expressed or implied, including any implied conditions or warranties of merchantability or fitness for any particular purpose. Ludger shall not be liable for any incidental, consequential or contingent damages.

This product is intended for *in vitro* research only.

Document Revision Number

Document # LC-EXO-A6-guide, version v1.0



Appendix 1: Troubleshooting Guide

1. Liquid does not flow.

- Insufficient centrifugation time: We recommend using extended centrifugation time for samples where high protein concentration/ high sample density is expected. Do not use higher than recommended centrifugation speed as this may disrupt the membrane structural integrity.
- Too much protein material: High concentration of protein or other high molecular weight material
 present in the sample can disrupt the liquid flow through the membrane. If liquid does not continue to
 pass despite extended centrifugation time, we recommend splitting the sample into two or more
 replicates. Maximum amount of protein per column should not exceed 20 µg.



Appendix 2: Material Safety Data Sheet

Manufacturer Ludger Ltd

Culham Science Centre, Oxford OX14 3EB, UK
Tel: +44 870 085 7011, Fax: +44 870 163 4620
Email: info@ludger.com, Website: www.ludger.com

Identification of the substance

LudgerClean Post-Exoglycosidase Clean-up Spin Columns

Composition

Polypropylene spin columns containing modified polyethersulfone

membrane.

Hazard indentification Non hazardous.

surrounding fire conditions.

Accidental release measures Not applicable.

Handling and storage Store at room temperature. Handle in accordance with Good

Laboratory Practice.

Exposure Controls / Wear appropriate protective clothing (safety spectacles, gloves,

laboratory coat) in accordance with Good Laboratory Practice.

Constructed of solid plastic and polymeric materials.

Physical and chemical properties

Not combustible.

Stability and reactivity

Toxilogical information

Toxicological, carcinogenic and mutagenic properties have not been

investigated.

Ecological information Data not available.

Disposal considerationsNo special requirements. Dispose of according to local requirements.

Transport informationContact Ludger Ltd for transportation information.

Regulatory information Data not available.

Other information The advice offered is derived from the currently available

information on the hazardous materials in this product or component. Consideration has been made regarding the quantities offered in the pre-dispensed container. The advice offered is, therefore, not all inclusive nor should it be taken as

descriptive of the compound generally.