

1-888-593-5969 biolynx.ca · tech@biolynx.ca

## Purification

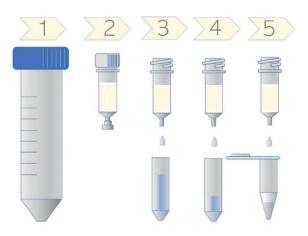
Exo-spin<sup>™</sup> Exosome Purification: Overview



#### A flexible range of bench-top products for quick and easy purification of exosomes from a variety of sources

- Excellent yields and high levels of purity Exosomes with ultralow protein and rRNA contamination.
- No ultracentrifugation required
   Protocol provides consistent results every time.
- Simple and reliable Isolate intact whole exosomes for functional studies.

#### Exo-spin<sup>™</sup> Exosome Isolation Workflow



Step 1: Remove cells and cellular debris.

**Step 2:**If required, use Exo-spin<sup>™</sup> Exosome Precipitation Buffer to precipitate exosomes.

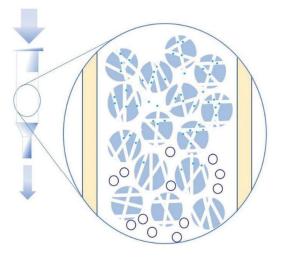
Step 3: Prepare the column by equilibrating with PBS.

**Step 4:** Add the pre-cleared sample or the precipitated exosome pellet resuspended in PBS to the Exo-spin<sup>™</sup> column, elute and discard the flow-through.

**Step 5**: Add PBS to the Exo-spin<sup>™</sup> column and elute your purified exosomes.

Cat code	Product name
EX01	Exo-spin™
EX02	Exo-spin™ blood
EX03	Exo-spin™ mini columns
EX04	Exo-spin™ midi columns
EX05	Exo-spin™ miniHD
EX06	Exo-spin™ buffer
EX07	Exo-spin™ 96
EX10	Exo-rack

#### Size exclusion chromatography (SEC) Technology





Principles of size exclusion chromotography



Exosomes run outside the beads, so elute first



Small particles and free proteins are trapped in beads

### Purification Exo-spin<sup>™</sup> Exosome Purification: Product Range

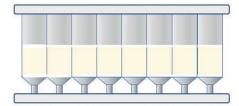
## A comprehensive range of products for isolation of exosomes from various sample types and volumes

#### Exo-spin<sup>™</sup> mini columns



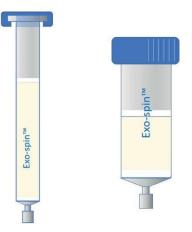
- A simple, twostep protocol
   Allows consistent and reliable
   purification of samples in under
   hours.
- Purify from blood Isolate directly from both sera and plasma samples.
- Process up to 50 ml per column Combine with Exospin™ Exosome Precipitation Buffer to isolate from up to 50 ml of low protein biofluids.

#### Exo-spin™ 96



- 96-well format Incorporates the technology of Exo-spin<sup>™</sup> mini columns into a highthroughput system.
- Process 8-96 samples at once
   Detachable strips of 8 columns
   gives flexibility
   to your isolations.
- **Gravity protocol** Purification achieved without the need for specialist instrumentation.

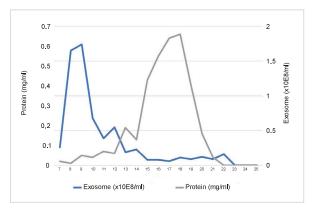
#### Exo-spin<sup>™</sup> mini-HD and midi columns



- High-resolution protocol Optimizes yield and purity and enables finer fraction collections.
- Process large and intermediate volumes Isolate from up to 500 ml of low protein biofluids and 1 ml of blood.

#### **Exosome Extraction Profiles**

Exo-spin<sup>™</sup> Midi Columns allow for gravity assisted fractionation, where exosomes are separated from the vast majority of proteins.



Exosomes were isolated using Exo-spin™ Midi Columns (Cat Code EX04) from 60 ml of conditioned medium generated by a human breast carcinoma cell line. Fractions (each of 500 µl) were collected and analysed to (1) evaluate particle numbers and (2) measure absorbance at 280 nm to evaluate protein concentration.

### **Detection** ExoLISA<sup>TM</sup> Exosome Detection Assay

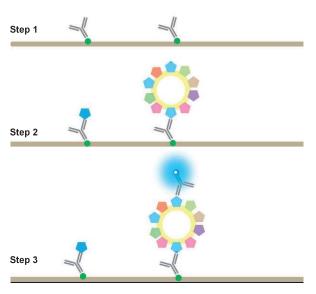


#### An exquisitely sensitive Europium Time-Resolved Immunofluorescence assay for exosome markers

Simplicity
 Assay is clear and
 simple allowing for high
 reproducibility.

• Sensitivity Europium-labelled antibodies are used for detection enabling the use of TRF. • **Specificity** Only antigens displayed in multiple copies are detected.

#### **ExoLISA™ Exosome Detection Assay**



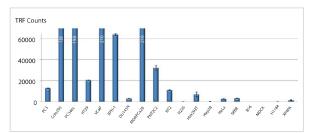
**Step 1:** Biotinylated antibody is bound to streptavidin coated assay plates.

**Step 2:** Biological samples are added. Exosomes and any free antigen are captured by the antibody.

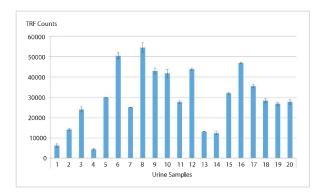
**Step 3:** Europium-labelled antibody of the same clone as the one used in Step 1 is added and binds specifically to exosome antigen. The epitopes of bound monomers are already occupied and not detected. Samples are read on a time-resolved fluorescence plate reader.

Cat code	Product name
EX501	ExoLISA™ CD9 Exosome Assay, 96 wells
EX502	ExoLISA™ CD63 Exosome Assay, 96 wells
EX503	ExoLISA™ CD81 Exosome Assay, 96 wells
EX-P31	Wash Buffer (Concentrate 25x), 20ml

#### Sample profiling



CD9 ExoLISA<sup>™</sup> exosome assay performed for 19 different cell lines. Off-the-scale readings indicated (x1000) on individual bars. Samples generating vastly different signal intensities can be measured due to the broad signal range covered by the assay.



ExoLISA™ exosome assay analysis of 20 urine samples shows great variation in CD9 content between samples.

# **Purified Exosomes**



Instant Exosomes<sup>™</sup> to accelerate your research

#### A range of freeze-dried, characterized exosomes

#### High quality a combination of size

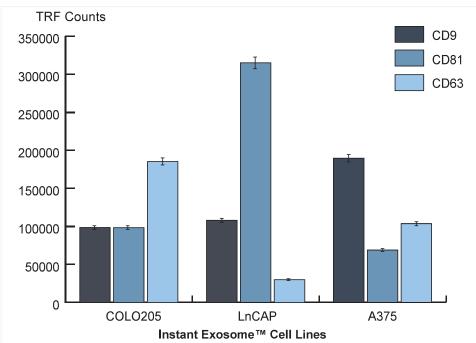
Exosomes are isolated using exclusion chromatography (SEC) and precipitation.

 Stable Instant Exosomes<sup>™</sup> are stable for longterm storage.

#### Characterized Each lot is characterized by ExoLISA<sup>™</sup> exosome detection

assay, nanoparticle tracking analysis (NTA) and bicinchoninic acid (BCA) assay.

#### Each sample is characterized by tetraspanin markers.



Commonly expressed exosomal markers assessed using the ExoLISA<sup>™</sup> assay are shown to be differentially expressed across the Instant Exosome™ samples isolated from human cancer cell lines.

Instant Exosomes<sup>™</sup> should not be used in a mass spectrometer or other instrumentation that may be sensitive to precipitants.

Cat code	Product name
EX301	Instant Exosomes™ from COLO205 cell line (Human colon carcinoma)
EX302	Instant Exosomes™ from LnCAP cell line (Human prostate adenocarcinoma)
EX303	Instant Exosomes™ from A375 cell line (Human malignant melanoma)

Characteristics of Instant Exosomes™					
Protein content per vial	25 µg				
Method of isolation	Precipitation + SEC				
Characterized by	ExoLISA™, NTA + BCA				
Nanoparticles/ml (average)	1 x 10 <sup>9</sup>				
State	Freeze-dried				
Storage temperature	-20°C				
Shelf life	12 months				

# **NTA Profiling**

Nanoparticle Tracking Analysis (NTA) service provided using ZetaView<sup>®</sup> Instrument

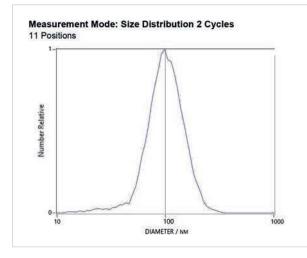
### Exosome characterization service for analysis of particle size and particle concentration

#### High quality

The service is performed in our labs by highly qualified scientists.

- Quick turnaround times Full reports are e-mailed within 5 to 10 business days.
- Competitive price Very low price per sample without the need of purchasing the equipment.

#### Example of an NTA report generated with the Zetaview<sup>®</sup>.

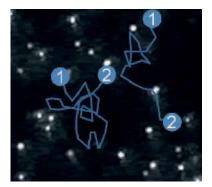


Result					
Concentration:		1.2E	+7 Particle	as / mL	
Dilution Factor:			2000		
Original Concen	tration:	2.4E	+10 Partic	les / cm³	
Quality					
Average Count	ed Partic	les per Fra	ame: 40		
Number of Trac	ed Parti	cles: 368			
Peak Analysis	(Numb	er Absoli	ute)		
Diameter / nm	•	umber Ab	solute	FWHM / nm	Percentage
101.2			28.1	85.6	100.0
X Values					
	Number	Volume			
X10	61.0	89.6			
X50	97.0	138.7			
	150.5	211.6			
X90	130.3				
X90 Mean	106.0	151.2			

#### Zetaview<sup>®</sup> Instrument.



Analysis service set up (purchase one per order)
NTA analysis of a single sample



Individual particle movement is tracked and recorded for characterization of exosome samples.

Image provided by Particle Metrix GmbH.



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Cell Guidance Systems' reagents and services enable control, manipulation and monitoring of the cell, both *in vitro* and *in vivo* 

#### **Growth Factors**

- Conventional (unformulated)
- PODS® Sustained release

#### Exosomes

- Exo-spin<sup>™</sup> Purification
- ExoLISA<sup>™</sup> ELISA-like detection
- Instant Exosomes<sup>™</sup> purified and characterized
- NTA Service
- Freeze drying service

#### **PeptiGel**®

 Tunable self-assembling peptide hydrogels

#### Other products and services

- Small Molecules
- Softwell<sup>™</sup> 2D hydrogel (Europe only)
- Orangu™ Cell counting reagent
- LipoQ<sup>™</sup> Lipid quantification assay
- Primary Hepatocytes

#### Cytogenetics

- Karyotype Analysis
- Array Hybridization

#### Scan for Exosome product page





#### www.cellgs.com

Available in Canada at ...







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