

# BEOTECH SUPPORT GROUP

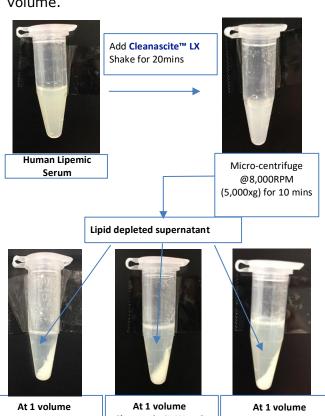
# Cleanascite™ LX

### For Lipemic Serum/Plasma Clarification

- Effectively replaces chlorinated/fluorinated hydrocarbons (eq. freon)
- Based on solid-phase referenced in over 70 publications in varied applications
- Under investigation as alternative to LipoClear for lipemic serum/plasma

#### **Protocol**

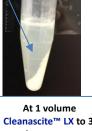
Cleanascite™ LX is supplied as an aqueous suspension of non-ionic adsorbent in DI water, pH 8.0. After centrifugation, the pellet is 1/2 of the total volume and the supernatant is 1/2 of the total volume.



Cleanascite™ LX to 1 volume serum (1:1 v:v ratio) 85-95 % turbidity removed (based on OD600)



OD600)



Cleanascite™ LX to 3 volumes serum (1:3 v:v ratio) 30-40 % turbidity removed (based on OD600)

All non-lipemic samples measured on Beckman Coulter AU680, Average from 3 independent serum samples

Volume ratio used was 1 volume Cleanascite™ LX to 2 volumes of serum

Due to competition on the beads, total analyte recoveries from lipemic serum are expected

% Variance from neat
serum
-2%

	serum
ALB	-2%
ALP	-2%
ALT	0%
AST	-5%
CO2	-5%
TBIL	0%
CALA	-8%
CRE	-11%
GLU	-2%
TP	-8%
BUN	-3%
NA	1%
K	-2%
CL	2%

For Research Use Only. Not for use in diagnostic procedures.



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Volume Ratio, Cleanascite□ LX : Sample	Dilution Factor
1:1	2.0
1:2	1.5
1:3	1.33

Cleanascite LX Item #	Size
LX155-5	5 ml
LX155-10	10 ml

Supplied as an aqueous suspension of non-ionic adsorbent in DI water, pH 8.0. When not in use, keep sealed. For best results store at 4°C. Do not freeze. Cleanascite<sup>TM</sup> retains full activity when stored as directed for at least 6 months.

#### Protocol

Lipid types and amounts can vary greatly, so the ratios shown are only intended to provide general quidance.

- 1. Resuspend **Cleanascite<sup>™</sup> LX** by gentle shaking. Excessive shaking may cause foaming. It should be completely resuspended prior to use.
- 2. Add Cleanascite<sup>TM</sup> LX to the sample at minimum 1:3 (or alternative higher, up to 1:1) ratio. Mix the sample by gentle shaking for 20 minutes.
- 3. Micro-centrifuge sample at 8,000 rpm's (5,000xg) for 10 minutes.
- 4. Carefully aspirate supernatant for analysis.

Optimization. Different sample volumes are easily scaled. Volume ratio can be adjusted up or down as required to remove the amount of impurities present.

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#### **CONTACT US**

We welcome your questions and comments regarding our products.

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