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CALA
Testing
Accreditation No. A 3538

Client:

Date of Sample:

Sample Location:

Type of sample:

Volume of sample (L):

Temp:

pH:

Turbidity:

Telephone:

Fax:

The methodology used to produce this report conforms to USEPA Method 1623. Based on the validation data, the method is fit for its intended use. Hyperion Research Ltd. is accredited for this analysis by CALA under the ISO/IEC 17025 standard.

DETECTION		VIABILITY
Volume Used (uL):		
Sample Equivalent Volume (L):		
Detection Limit:*		cysts or oocysts/ 100 L
# CYSTS/ OOCYSTS	<i>Giardia</i>	<i>Crypto.</i>
Dead		
Viable		
Total	0	0

*Detection Limit is calculated assuming a minimum of 1 cyst or oocyst observed in the Sample Equivalent Volume.

MICROBIOLOGICAL RESULTS

Giardia cysts: /100 L
Cryptosporidium oocysts: /100 L

Notes: 1. A value of "0" really means "below the detection limit".
 2. These results apply to this sample only.

Comments:

Analyst:

LAB NOTES

Date received: _____ Time received: _____

Condition of sample: _____ Lab ID #: _____

Type of filter:

Date concentration complete:

Time concentration complete:

Concentration Analyst:

Pellet volume:

IMS system used:

IMS Lot#:

Monoclonal antibody:

MAb Lot#:

Resuspension Volume (uL):

Control G: _____ Control C: _____

Date staining complete:

Time staining complete:

Staining Analyst:

Microscope Analyst:

Current Quality Control Data

	Lab Water Spike		Matrix Spike	
	% Recovery	RSD	% Recovery	RSD
<i>Giardia</i>	51.1	19.3	47.4	27.2
<i>Crypto.</i>	57.1	23.5	63.8	26.8

These data indicate ongoing precision and recovery from spiked filter samples.