MAYO CLINIC
Mayo Medical Laboratories

Laboratory Service Report

1-800-533-1710

Patient Name	Patient ID	Age	Gender	Order #
SAMPLEREPORT,LFACX_ABNORMAL	SA00066584	29	M	SA00066584
Ordering Phys CLIENT,CLIENT				**DOB** 05/05/1984
Client Order # SA00066584	**Account Information**			**Report Notes**
Collected 03/28/2014 08:56	C7028846-DLMP Rochester SDSC 2 - Client Support Rochester, MN 55901			
Printed 03/28/2014 14:10				

Test	Flag	Results	Unit	Reference Value	Perform Site*
Cryptococcus Ag Titer, LFA, CSF	C	1:80		Negative	SDL

RECEIVED: 03/28/2014 09:01 **REPORTED:** 03/28/2014 09:15

CRITICAL RESULT
The cryptococcal antigen detection method was changed from a latex agglutination (LA) test to a lateral flow immunoassay (LFA) test in May, 2014. End point titers are not interchangeable and do not correlate between the two methods.

Providers should transition to monitoring end-point titers by the LFA method. End-point titer values by the LA method will be reported alongside the LFA titer result for comparison until August, 2014, at no charge. After this date, only the LFA titer will be performed.

Latex Agglutination Titer: 1:8

Fungal Culture, CSF	AB				MCR

RECEIVED: 03/28/2014 09:23 **REPORTED:** 03/28/2014 09:24

SOURCE: CEREBROSPINAL FLUID
FUNGAL CULTURE, CSF FINAL
CRYPTOCOCCUS NEOFORMANS
Critical Result.

Cryptococcus Ag w/Reflex, LFA, CSF

RECEIVED: 03/28/2014 09:01 **REPORTED:** 03/28/2014 09:05

Cryptococcus Ag Screen w/Titer, CSF	AB	Positive		Negative	SDL

Reflex testing has been ordered to establish a cryptococcal antigen titer.

* Performing Site:

SDL	Mayo Clinic Laboratories - Rochester Superior Drive 3050 Superior Dr. NW Rochester, MN 55901	Lab Director:
MCR	Mayo Clinic Laboratories - Rochester Main Campus 200 First St SW Rochester, MN 55905	Lab Director:

* Report times for Mayo performed tests are CST/CDT