



Patient ID <b>SA00066581</b>	Patient Name <b>SAMPLEREPORT, SLFAT_ABNORMAL</b>	Birth Date <b>1984-05-05</b>	Gender <b>M</b>	Age <b>29</b>
Order Number <b>SA00066581</b>	Client Order Number <b>SA00066581</b>	Ordering Physician <b>Client, Client</b>	Report Notes	
Account Information <b>C7028846 DLMP Rochester</b>		Collected <b>28 Mar 2014 08:55</b>		

**Cryptococcus Ag Titer, LFA, S**



**1:10**

Abn

**SEMI-URGENT 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.

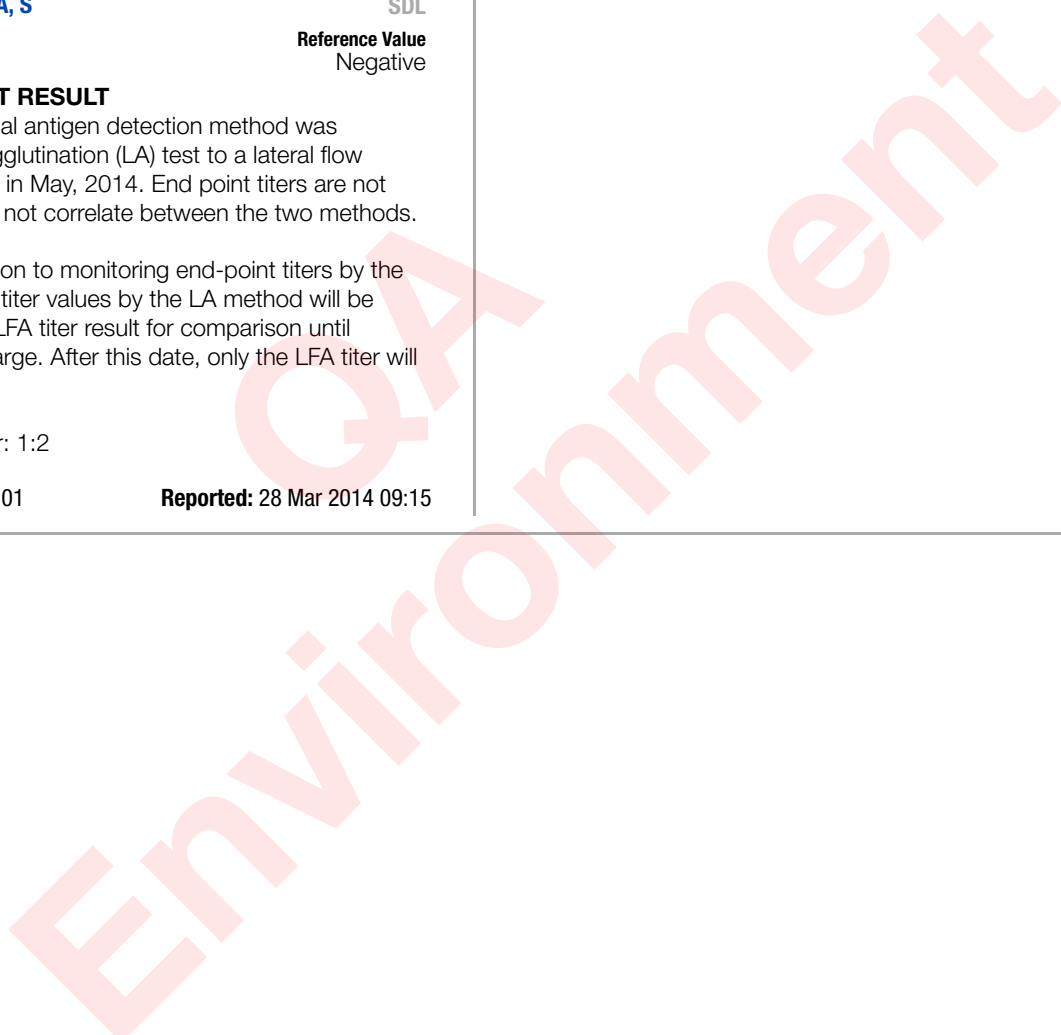
Latex Agglutination Titer: 1:2

**Received:** 28 Mar 2014 09:01

**Reported:** 28 Mar 2014 09:15

SDL

Reference Value  
Negative



**Performing Site Legend**

Code	Laboratory	Address
SDL	Mayo Clinic Laboratories - Rochester Superior Drive	3050 Superior Drive NW, Rochester MN 55901