

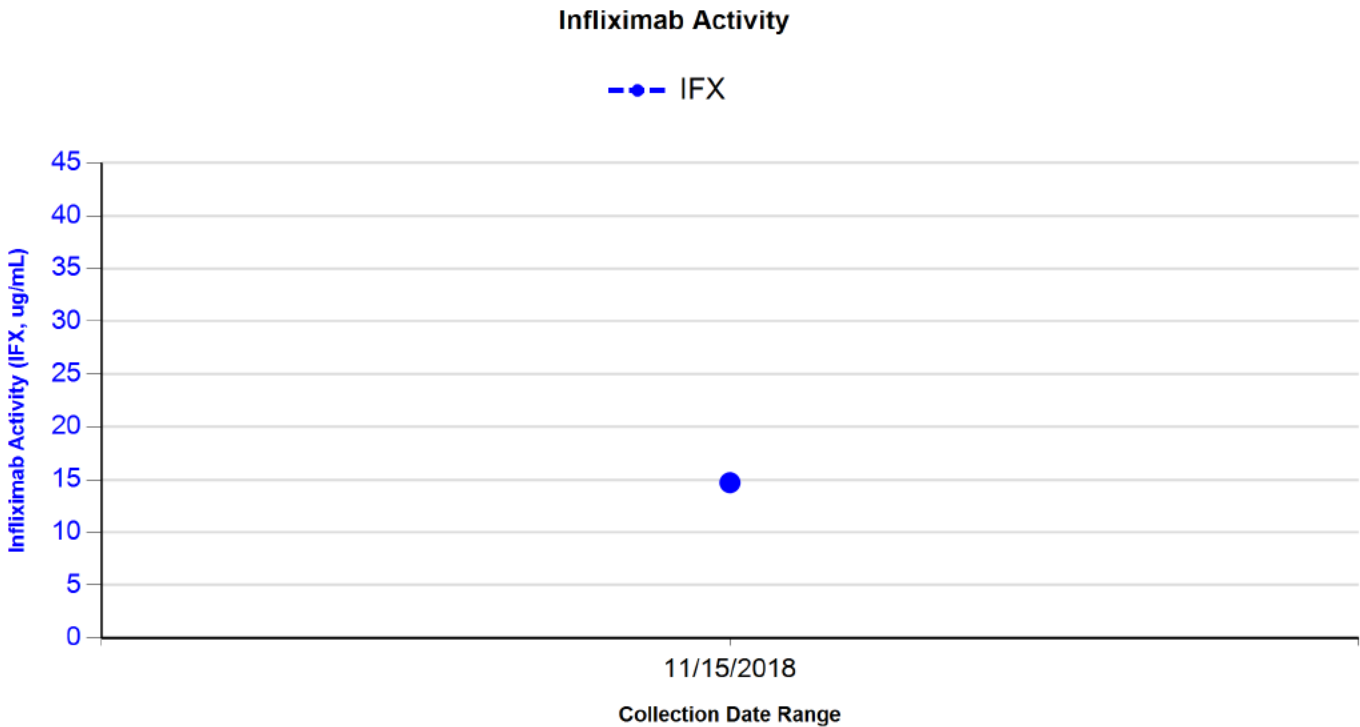
Patient: [REDACTED]
 DOB: [REDACTED] Age: 17 Gender: F
 Patient Identifiers: [REDACTED]
 Visit Number (FIN): [REDACTED]

Client: [REDACTED]
 Physician: [REDACTED]

ARUP Test Code: 2013612
 Collection Date: 11/15/2018
 Received in lab: 11/16/2018
 Completion Date: 11/17/2018

Patient History *

| <u>Collection Date</u> | <u>Infliximab Activity (IFX) ug/mL</u> | <u>Accession Number</u> |
|------------------------|--|-------------------------|
| 11/15/2018 | 14.71 | 18-319-102289 |



Cutoff values are 0.65 ug/mL for infliximab.

*Consecutive test results are displayed on this chart; however, this result set may be incomplete due to variations in the demographic information submitted for prior tests. If the information on this chart appears incomplete, please consult this patient's prior charts.



Patient [REDACTED]
 ARUP Accession: 18-319-102289

Infliximab or Biosimilar Activity with Reflex to Antibody

Patient: [REDACTED] | Date of Birth: [REDACTED] | Gender: F | Physician: [REDACTED]
Patient Identifiers: [REDACTED] | Visit Number (FIN): [REDACTED]

Interpretive Comments

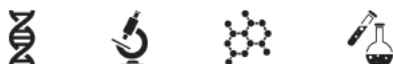
Interpretive Data: This test measures the capacity of infliximab to neutralize TNF-alpha activity. If infliximab is not detected, testing for neutralizing antibodies (NAb) will be performed. Infliximab NAb titer is obtained by identifying the minimal serum dilution at which blocking of infliximab activity is no longer observed.

This test is used to evaluate secondary response failures to infliximab therapy. Secondary response failure is defined as loss of clinical response after initial improvement of clinical signs and symptoms. Therapeutic decision should rest on both the clinical response and the knowledge of the fate of the drug including the emergence of immunogenicity in individual patients.

Circulating infliximab levels have been shown to vary considerably between patients. These differences relate to route and frequency of administration and patient-related features such as age, gender, weight, drug metabolism, and concomitant medications such as methotrexate and other immunosuppressants.

| IF Infliximab Activity is... | AND Infliximab Neutralizing Antibody Titer is... | THEN... |
|------------------------------|--|--|
| Not Detected | Not Detected | A higher dosage of infliximab or shortening the dosing interval may be appropriate. |
| Not Detected | 1:20 or greater | A change to another anti-TNF-alpha drug may be appropriate. |
| 0.65 ug/mL or greater | Not Applicable | A change to another type of therapy (not targeting TNF-alpha) may be appropriate, if the patient did not respond adequately to infliximab therapy. |

Test developed and characteristics determined by ARUP Laboratories. See Compliance Statement B: www.aruplab.com/CS.



Patient: [REDACTED]
ARUP Accession: 18-319-102289
[REDACTED]