Lab Dept: Chemistry

Test Name: INTERLEUKIN-2 RECEPTOR

**General Information** 

Lab Order Codes: IL2R

**Synonyms:** IL 2 Receptor, Soluble (CD25)

**CPT Codes:** 83520 – Immunoassay for analyte other than infectious agent antibody

or infectious agent antigen; quantitative, not otherwise specified

**Test Includes:** Interleukin-2 Receptor level reported in pg/mL.

Logistics

**Test Indications:** Clinical conditions in which elevated soluble IL-2R levels are detected

include AIDS, autoimmune disease, sarcoidosis, and a variety of leukemias and lymphomas. In HIV positive individuals, IL-2R is elevated during the asymptomatic phase as well as during persistent generalized lymphadenopathy and symptomatic phases. IL-2R detection may be useful in measuring T cell activation and monitoring HIV pathogenesis. Elevated IL-2R levels have clinical and prognostic significance in

patients with malignant lymphoma, non-Hodgkin's lymphoma, B cell and

undifferentiated lymphomas.

**Lab Testing Sections:** Chemistry - Sendouts

Referred to: Mayo Clinic Laboratories forward to ARUP (MML Test: FIL2S), (ARUP

Test: 0051529)

Phone Numbers: MIN Lab: 612-813-6280

STP Lab: 651-220-6550

**Test Availability:** Daily, 24 hours

**Turnaround Time:** 3 - 6 days

**Special Instructions:** Separate specimens must be submitted when multiple tests are

ordered.

Specimen

Specimen Type: Blood

**Container:** SST (Gold, marble or red)

**Draw Volume:** 3 mL (Minimum: 1.2 mL) blood

**Processed Volume:** 1 mL (Minimum: 0.4 mL) serum/plasma

**Collection:** Routine venipuncture

Special Processing: Lab Staff: Centrifuge specimen and separate serum/plasma within 2

hours of collection; aliquot into a screw-capped round bottom plastic vial. Store and ship at frozen temperatures. Forward promptly.

Patient Preparation: None

**Sample Rejection:** Mislabeled or unlabeled specimens; refrigerated specimens;

contaminated or heat-inactivated specimens

Interpretive

**Reference Range:** 175.3 – 858.2 pg/mL

Critical Values: N/A

Limitations: Cytokine levels may demonstrate diurnal variation. For longitudinal

comparison, it is recommended that cytokine levels be determined at

the same time of day.

Results are to be used for research purposes or in attempts to

understand the pathophysiology of immune, infectious or inflammatory

conditions.

Methodology: Quantitative Multiplex Bead Assay

References: Mayo Clinic Laboratories Web Page May 2020

**ARUP Laboratories May 2020** 

**Updates:** 3/18/2004: Test moved from Specialty Laboratories to Mayo Medical

Laboratories forward to Focus Technologies. Note changes in CPT

code, Methodology, and reference range.

9/16/2008: CPT update, previously reported at 83520

12/16/2013: MML now forwards to ARUP. Note change in CPT

(previously 84238), method and reference ranges.

2/19/2014: Ref range update, previously listed as 0-1033 pg/mL.

9/18/2017: Update tube types.

5/18/2020: Updated ref range and minimum volume requirements.