

KOS Diagnostic Lab





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NAME : B/O SULEKHA

AGE/ GENDER : 7 DAYS(S)/Male PATIENT ID : 1656941

COLLECTED BY : REG. NO./LAB NO. : 012410290056

 REFERRED BY
 : 29/Oct/2024 07:17 PM

 BARCODE NO.
 : 01519783
 COLLECTION DATE
 : 29/Oct/2024 07:18 PM

 CLIENT CODE.
 : KOS DIAGNOSTIC LAB
 REPORTING DATE
 : 29/Oct/2024 08:18 PM

CLIENT ADDRESS : 6349/1, NICHOLSON ROAD, AMBALA CANTT

Test Name Value Unit Biological Reference interval

CLINICAL CHEMISTRY/BIOCHEMISTRY G-6-PD (QUANTITATIVE KINECTICS)

G6PD (QUANTITATIVE KINECTICS) by SPECTROPHOTOMETRY

16.51^H

U/gHb

4.6 - 13.5

INTERPRETATION:

- 1.G-6 PD deficiency is a sex/X-linked recessive genetically inherited RBC enzyme disorder making the cells vulnerable to oxidative denaturation of haemoglobin characterized by abnormally low levels of glucose-6-phosphate dehydrogenase.
- 2. G6PD deficiency is the most common human enzyme defect.
- 3. G-6 PD levels are highest in young cells and decrease as cells age, hence in cases of G-6 PD deficiency, the older cells are preferentially destroyed.
- 5.G6PD helps body process carbohydrates and turn them into energy.
- 6. Hemolytic susceptibility in affected persons can increase greatly during intercurrent illness or upon exposure to various drugs that have oxidant properties like Primaquin, Nalidixic acid, Nitrofurantoin etc., Marked genetic heterogeneity has been reported in G-6 PD deficiency cases and > 300 variants have been defined. This heterogeneity causes variability in the degree of deficiency, types of cells affected, types of drugs causing hemolysis and susceptibility to chronic hemolysis and neonatal jaundice.

COMMON DRUGS THAT CAN INDUCE HEMOLYSIS IN G6PD DEFICIENT INDIVIDUALS INCLUDE:

- 1. Anti Malarial drugs (like primaquine, pamaquine, and chloroquine).
- 2. Sulfonamides (such as sulfanilamide, sulfamethoxazole, and mafenide).
- 3. Thiazolesulfone, methylene blue and naphthalene.
- 4. Certain analgesics (such as aspirin, phenazopyridine, and acetanilide)
- 5. Few non-sulfa antibiotics (nalidixic acid, nitrofurantoin, isoniazid, dapsone, and furazolidone).

*** End Of Report ***



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