



Metropolis Healthcare Ltd., EQAS

KRSNAA DIAGNOSTIC LTD UNA (1236)

Outlier And Analyte Summary Report

Outlier Details For Cycle No 240104 and Sample No 01

Report Date : 10/07/2024

Analyte	Instrument	Result Value	Standard Unit
X Glucose (urine)	Manual	1+	-
X Blood (urine)	Manual	Negative	-
X Urine Pregnancy Test	-	Negative	-

Legend @ : Acceptable
X : Unacceptable
: Not Evaluated
⌚ Delayed Result Entry

Problem Classification: SAMPLE NOT MAINTAIN COOL PACK CHAIN.

Corrective Action: SPLIT TESTING DONE.

Reviewed by: [Signature]

Dated: 12/7/24.

Instrument : Manual

Analyte	Standard Unit	Result Value	Accepted Value
X Glucose (urine)	-	1+	+2,+3,+4
X Blood (urine)	-	Negative	+1,+2,+3
@ pH	-	5	5,5.5,6
@ Specific Gravity	-	1.025	1.020,1.025,1.030
@ Protein (Urine)	-	1+	+1,+2,+3
@ Ketone (urine)	-	Negative	+1,Neg,Tr

Instrument : -

Analyte	Standard Unit	Result Value	Accepted Value
X Urine Pregnancy Test	-	Negative	Pos

Legend @ : Acceptable
 @ : Acceptable
 X : Unacceptable
 # : Not Evaluated
 ⌚ : Delayed Result Entry

Total Parameters	7
Not Evaluated Parameters	0
Evaluated Parameters	7
Outlier Parameters (X)	3
EQAS Score Clinical Path - Urine Routine Chemistry	57.14 %

Dr Puneet Kumar Nigam

Dr Puneet Kumar Nigam
 PT coordinator & Technical Manager, MHL EQAS
 Unit No. 409-416.
 Commercial Building - 1A
 Kohinoor Mall, Kirod Road, Kurla (W),
 Mumbai - 400070

CORRECTIVE ACTION & PREVENTIVE ACTION (CAPA)

Name of the Department: *Clinical Pathology*

Sr. No	Date	Non Conformity Observed	Root Cause Analysis	Corrective Action & Preventive Action	Date of Closure	Signature
	<i>2-8-24</i>	<i>UPT</i>	<i>Random error</i>	<i>ILC DONE</i>	<i>2-8-24</i>	<i>mj</i>

Issued by : Vinod Lonkar <i>Vinod</i>			Issued on : 01.03.2024		
Approved by : Dr Amol Patil <i>Amol</i>					
Doc No. GEN-69		Version No : 2		Amend No :	
				Amend Date :	



Krsnaa Diagnostics Ltd.
Regional Hospital UNA
Himachal Pradesh 174303

INTER LABORATORY COMPARISON RESULTS

Sr. No	Date	Analyte Name	Barcode ID	Lab Result	Unit	ILC lab Result	Unit	Acceptability Criteria	Observed CV %	Acceptability range	Result Acceptable Yes / No	Done by	Reviewed by
	2-8-24	UPT	116313615	Negative	NA	Negative	NA	NA	NA	NA	Yes	<i>[Signature]</i>	<i>[Signature]</i>

Qualitative: Clinically comparable results shall be Concordance.
Clinically not comparable results shall be Disconcordance.

Quantitative: Acceptability: 10 %. (As per NABL 112)

Signature of Doctor *[Signature]*

Issued by : Vinod Lonkar <i>[Signature]</i>		Issue Date : 01.09.2023	
Approved by : Dr. Manish Karekar <i>[Signature]</i>		Amend No. --	
Doc No. GEN-01		Version No : I	
		Amend Date .	

SPLIT TESTING RECORD

Sr.No.	Date	Analyte Name	Barcode ID	Initial Result	Lab BRI	Initial test done by	Split test result	Split test done by	Acceptability criteria applied	Acceptability Range	Split test result acceptable (Yes / No)	Comment if not applicable
1	12-7-24	Sugar	20060	Absent	NA	NIHITA	Absent	POOJA	NIA	NA	Yes	-
2	" "	RBC	" "	Absent	NA	NIHITA	Absent	POOJA	NIA	NA	Yes	-
3	" "	WBC	" "	4-5	NA	NIHITA	5-6	POOJA	NA	NA	Yes	
4	" "	E-cell	" "	2-4	NA	NIHITA	3-4	POOJA	NA	NA	Yes	
5	" "	CRYSTAL	" "	Absent	NA	NIHITA	Absent	POOJA	NA	NA	Yes	
6	" "	OTHERS	" "	Absent	NA	NIHITA	Absent	POOJA	NA	NA	Yes	

NIHITA
Entered by

Reviewed by

Approved by

Issued by : Vinod Lonkar <i>Vinod</i>	Issue Date : 01.09.2023
Approved by : Dr. Manish Karekar <i>Manish</i>	Amend No. --
Doc No. GEN-14	Version No : 1
	Amend Date .

SPLIT TESTING RECORD

Sr.No.	Date	Analyte Name	Barcode ID	Initial Result	Lab BRI	Initial test done by	Split test result	Split test done by	Acceptability criteria applied	Acceptability Range	Split test result acceptable (Yes / No)	Comment if not applicable
1	12-7-24	Colony	20060	PIY	NA	NILITA	PIY	POOJA	NA	NA	Yes	
2	"	Approach	"	CLCQX	NA	NILITA	UcaD	POOJA	NA	NA	Yes	
3	"	PIH	"	G.S	NA	NILITA	G.G	POOJA	NA	NA	Yes.	
4	"	SGT	"	1.020	NA	NILITA	1.015	POOJA	NA	NA	Yes.	
5	"	Dotter	"	Absent	NA	NILITA	Absent	POOJA	NA	NA	Yes.	
6	"	Blood	"	Absent	NA	NILITA	Absent	POOJA	NA	NA	Yes.	

NILITA
Entered by

Reviewed by

Approved by

Issued by : Vinod Lonkar <i>Vinod</i>	Issue Date : 01.09 2023
Approved by : Dr. Manish Karekar <i>Manish</i>	
Doc No. GEN-14	Version No : 1
Amend No. --	Amend Date .



मुख्यमंत्री निःशुल्क निदान योजना


स्वास्थ्य सुविधाएँ घर-द्वार, कृत संकल्पित है हिमाचल सरकार



UNA

Medical Laboratory Report

Patient Name : Mrs NAMWATI
Age and Gender : 19 Years / Female
Category : OPD - HP UNA LAB
Referring Doctor : Self
Sample Processed at : HP UNA LAB

Patient UID No : UNAP240800092394
PRN No : 4073**6--
Registered On : 01 Aug 2024 18:37
Sample UID No. 
110313615

CLINICAL PATHOLOGY

Test Done	Observed Value	Units	Biological Reference Interval
Pregnancy Test Rapid Immunochromatography	Negative		Negative



Roopam

DR .ROOPAM (MD.PATHOLOGY)
CONSULTANT PATHOLOGIST

--- END OF REPORT ---



मुख्यमंत्री निःशुल्क निदान योजना

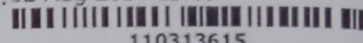
स्वास्थ्य सुविधाएँ घर-द्वार, कृत संकल्पित है हिमाचल सरकार



HAMIRPUR.

Medical Laboratory Report

Patient Name : Mrs ILC NAMWATI
Age and Gender : 19 Years / Female
Category : OPD - HP UNA LAB
Referring Doctor : Self
Sample Processed at : HP HAMIRPUR LAB

Patient UID No : UNAP240800092507
PRN No : UNAP240800092507
Registered On : 02 Aug 2024 13:09
Sample UID No. 
110313615.

CLINICAL PATHOLOGY

Test Done	Observed Value	Units	Biological Reference Interval
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Pregnancy Test

Negative

Negative

Sample Type- Urine
Rapid Immunochromatographic Immunoassay

Test Description:

The Card Pregnancy Test is a rapid test that qualitatively detects the presences of hCG in urine specimen at the sensitivity of 25 mIU/mL. The test utilizes a combination of monoclonal and polyclonal antibodies to selectively detect elevated levels of hCG in urine. At the level of claimed sensitivity, the hCG Card Pregnancy test shows no cross-reactivity interference from the structurally related glycoprotein hormones hFSH, hLH and hTSH at high physiological levels.

Test Interpretation: -

POSITIVE: * Two distinct red lines appear. One line Should be in the control region (C) and another line should be in the test region (T)

*NOTE:

The intensity of the red color in the test line region (T) will vary depending on the concentration of hCG present in the specimen. Therefore, any shade of red in the test region (T) should be considered positive.

NEGATIVE: One red line appears in the control region (C). No apparent red or pink line in the test region (T)

INVALID: Control line fails to appear. Insufficient specimen volume or incorrect procedural techniques are the most likely reasons for control line failure. Review the procedure and repeat the test with a new test device. If the problem persists, discontinue Using the test kit immediately and contact your local distributor.

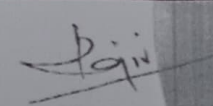
Test Limitations:-

1. The hCG Card Pregnancy Test is a qualitative test, therefore, neither the quantitative value nor the rate of increase in hCG can be determined by this test.
2. Very dilute Urine specimens, as indicated by a low specific gravity, representative levels of hCG. If pregnancy is still suspected, a first morning urine specimen should be collected 48 hours later and tested.
3. False negative results may occur when the levels of hCG are below the sensitivity level of the test. When pregnancy is still suspected, a first morning urine specimen should be collected 48 hours later and tested.
4. Very low levels of hCG (less than 50 mIU/mL) are present in urine specimen Shortly after implantation however, because a significant number of first trimester pregnancies terminate for natural reasons, a test result that is weakly positive should be confirmed by retesting with a first morning urine specimen collected 48 hours later.
5. A number of conditions other than pregnancy, including trophoblastic disease and certain non-trophoblastic neoplasms including testicular tumors, prostate cancer, breast cancer, and lung cancer, cause elevated levels of hCG. Therefore, the presence of hCG in urine specimen should not be used to diagnose pregnancy unless these conditions have been ruled out.
6. This test provides a presumptive diagnosis for pregnancy. A confirmed pregnancy diagnosis should only be made by a physician after all clinical and laboratory findings have been evaluated.

Reflex Test: Serum Beta-HCG

References: Kit Insert Rapid Test.




Dr. Rajiv Dogra
(MDPhysician), DCP Pathologist

--- END OF REPORT ---