



analyzed samples with one day delayed frozen samples to identify the reliability on finding and difference in further treatment regimens **FIGURE 1** [9-11].

Material and Methods

Urine samples were collected in a sterile urine sample bottles and were analyzed within 4 hours of collection. 190 samples were analyzed in duplicated and 213 samples were analyzed in a single time run after collection and were repeated after 24 hours at 4°C for solutes concentration like glucose and proteins etc. by the automated analyzer equipment Supertron (Hitachi-Mannheim, Germany). Following this procedure, we tested 190 samples to calculate accuracy and all 403 (190+213) to confirm steadiness of samples [12,13]. Each component assertion of urinalysis was decided for distinctive degrees of analysis. Critical contrasts between precision and stability was estimated.

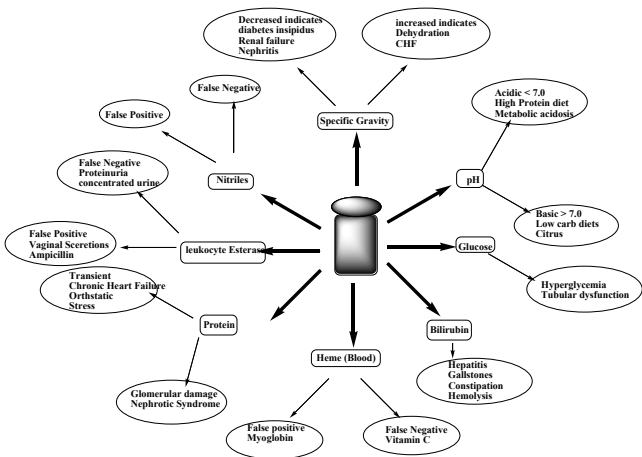
Comparison between the test samples and estimated standards was done to identify the correctness of results.

Results

Highly accurate leukocyte esterase results are shown in **TABLE 1**, which became varied with the passage of time and displayed false positive results due to deterioration. The inadmissible changes in the final readings were >75% whereas the test analyzed as negative went a minor alteration in permanency. In addition to this, 25% of results for leukocyte esterase overdue for twenty four hours were read as negative false, tests for nitrates were together specific and steady as shown in **TABLE 1 AND FIGURE 1**.

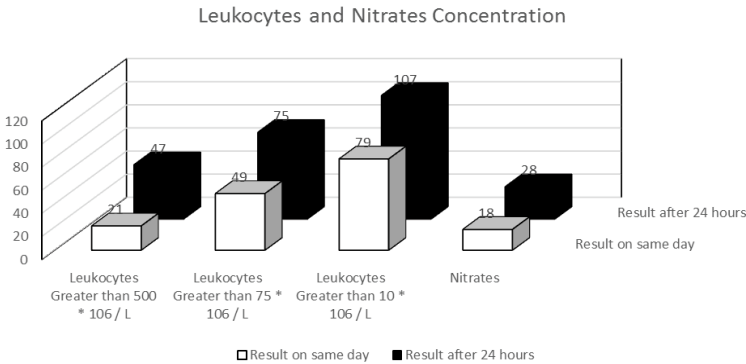
When urine samples were analyzed for proteinuria, **FIGURE 2** indicates the false positive results for greater than 5000 mg/L, 1000 mg/L and 300 mg/L when samples were

FIGURE 1. Showing different parameters of urine analysis.



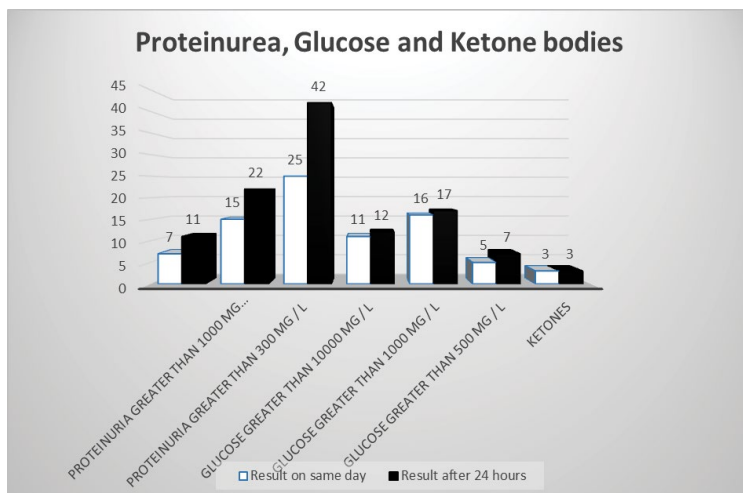
| TABLE 1. Indicating the tests to be analyzed and results. |                    |                       |                |
|-----------------------------------------------------------|--------------------|-----------------------|----------------|
| Test                                                      | Result on same day | Result after 24 hours | Conclusion     |
| Leukocytes Greater than 500 10 <sup>6</sup> /L            | 21                 | 47                    | False positive |
| Leukocytes Greater than 75 10 <sup>6</sup> /L             | 49                 | 75                    | False positive |
| Leukocytes Greater than 10 10 <sup>6</sup> /L             | 79                 | 107                   | False positive |
| Nitrates                                                  | 18                 | 28                    | False positive |

FIGURE 2. Showing comparison of sample before and after 24 hours.



**TABLE 2. Indicating the concentration of testing specimen.**

| Test                               | Result on same day | Result after 24 hours | Conclusion        |
|------------------------------------|--------------------|-----------------------|-------------------|
| Proteinuria Greater than 5000 mg/L | 7                  | 11                    | False positive    |
| Proteinuria Greater than 1000 mg/L | 15                 | 22                    | False positive    |
| Proteinuria Greater than 300 mg/L  | 25                 | 42                    | False positive    |
| Glucose greater than 10000 mg/L    | 11                 | 12                    | No big difference |
| Glucose greater than 1000 mg/L     | 16                 | 17                    | No big difference |
| Glucose greater than 500 mg/L      | 5                  | 7                     | No big difference |
| Ketones                            | 3                  | 3                     | No big difference |

**FIGURE 3. Expressing the concentration of different entities in Urine specimen**

allowed to retain for one day. While in case of Glucose and ketone bodies, not much difference was observed when compared with immediate performed results **TABLE 2 AND FIGURE 3**.

## Discussion

The results of this study identify that urine analysis for RBCs and WBCs shows variation if tested after one day. Results may become unsatisfactory as false positive or false negative during this time. On the other hand, No prominent differences among the precision and steadiness for urine ketone bodies, glucose concentration and nitrates were observed. Instant assessment-retest precision was objectionable for protein concentration, displaying higher values (false positive) when refrigerated for 24 hours.

Reflectance interpretations of urinalysis dip-sticks that are semi-automated are extra detailed than pictorial analyses and trainings of exactness through Miditron Junior to check the dipsticks bring into being that it was specific. Detailed examination justified that value of protein, glucose concentration and nitrates compounds in insincerely synthesized specimens also have great reproductivity, but results for haemoglobin level in urine were less accurate.

High concentration of glucose i.e. 1000 mg/L was good to analyze but below deadline values it was very difficult to measure. The absence of correctness of glucose concentration in commercially obtainable dipsticks has shown good results. When comparing the measurable hexokinase technique, individually the Chem strip could distinguish glucose urine concentration at 0.29 g/L and 0.59 g/L.

Supertron dipsticks need to be preserved cautiously as their exposure to oxygen in open air leads to change of color and variation in results of constituents present and ultimately perception of loss of specificity. In this way, it can show glucose false positive results during first seven days and inaccurate and false negative results for blood in urine during one month time period.

Healthcare professionals and other first line warriors for treatment of diseases relay on the results of laboratories to completely diagnose and treat the ailment. The laboratory technicians also undergoes many circumstances to increase efficacy of results without adding more cost to the patient so that the precise information could be delivered at the right time to physicians and they would have a good decision about patient health.

Urinalysis involves a series of tests and results that are sometimes performed manually and automatically to quantify the constituents present in the urine. The main focus of this investigation is to apply different screening parameters to find the patients renal system problems and other metabolizing diseases. Urine analysis through dipsticks can be compare with other test strategies but this would be probably more beneficial than others due to high efficacy and precision than others but this could more justified by more protocols.

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## Conclusion

The present research comes out with the conclusion that performing urine test within time frame is one of the important area which the most laboratories and physicians lapse. The result of which suffers the innocent people and they may be treated for the disease which they do not have, due to false positive results. Same for those people which have the disease but the result comes false negative and they are not given treatment on time leading towards lethal.

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