

INNOVATIVE USE OF LATERAL FLOW IMMUNOASSAY FOR WHOLE BLOOD PROGESTERONE DETECTION IN CATTLE REPRODUCTION

Thomas Zendo

Department of Clinical Studies, University of Nairobi, Nairobi, Kenya

Abstract

This study explores the innovative application of lateral flow immunoassay (LFIA) for the detection of progesterone in whole blood samples as a method for assessing reproductive status in cattle. Traditional methods of progesterone measurement often involve complex laboratory procedures that can delay timely decisions in livestock management. The LFIA technique offers a rapid, cost-effective, and user-friendly alternative for on-site testing. This research evaluates the sensitivity and specificity of the LFIA against established immunoassay methods, highlighting its potential to improve reproductive management in cattle. Results indicate that LFIA provides reliable progesterone levels, enabling farmers and veterinarians to make informed decisions regarding breeding and reproductive health. This innovative approach not only enhances reproductive efficiency but also supports the overall productivity of cattle operations.

Keywords Lateral flow immunoassay, Progesterone detection, Cattle reproduction, Whole blood samples, Reproductive status assessment, Livestock management, Rapid testing.

INTRODUCTION

The reproductive health of cattle is vital for the sustainability and productivity of livestock operations. Accurate assessment of reproductive status is essential for effective breeding management, timely interventions, and optimal herd performance. Progesterone, a key hormone involved in the regulation of reproductive cycles, serves as a critical indicator of estrus, pregnancy, and overall reproductive health in cattle. Traditional methods for measuring progesterone levels, such as enzyme-linked immunosorbent assays (ELISA) and radioimmunoassays, often require sophisticated laboratory equipment, trained personnel, and lengthy processing times.

These factors can hinder prompt decision-making in farming environments.

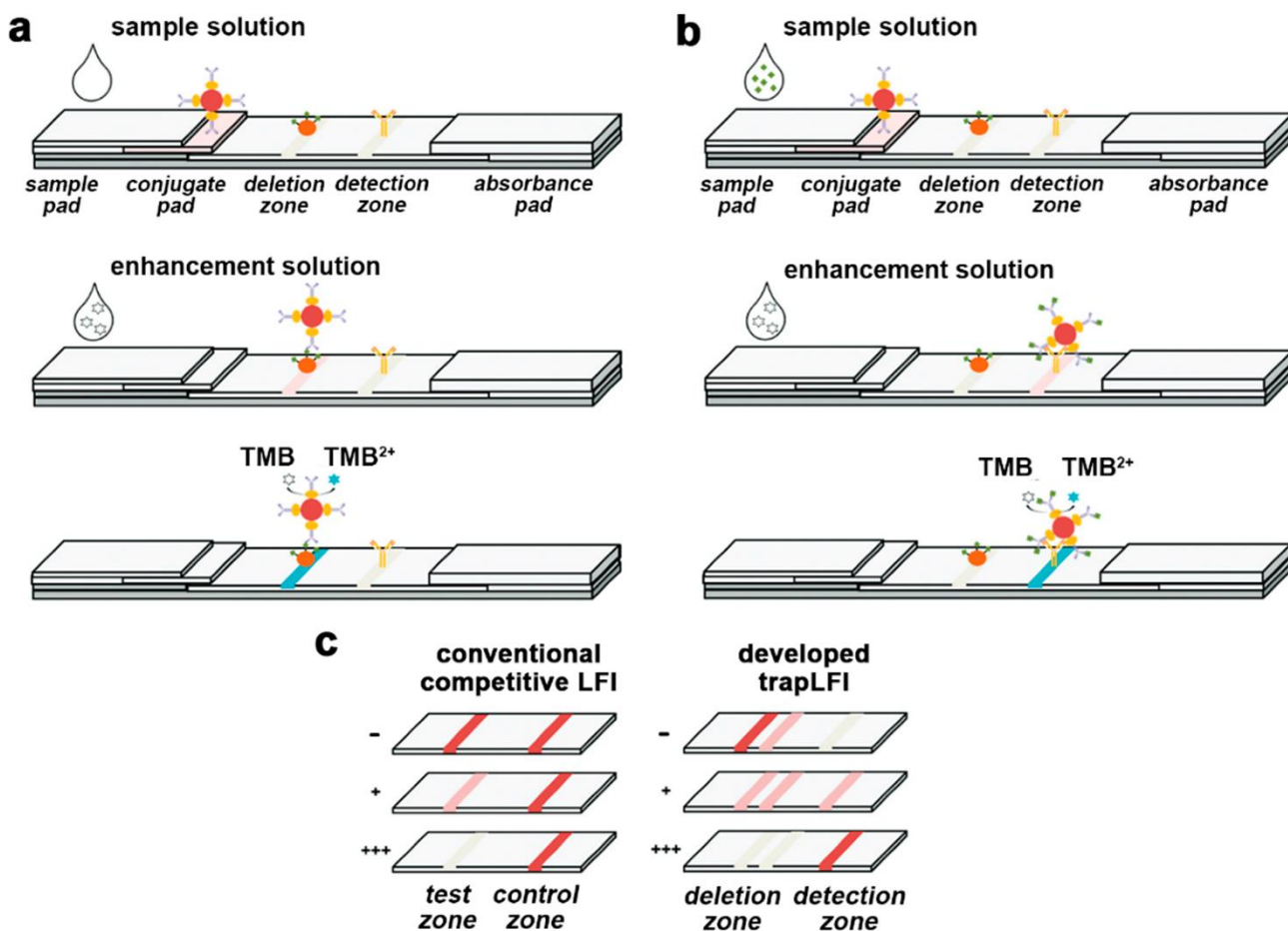
In recent years, the development of lateral flow immunoassays (LFIA) has emerged as a promising alternative for rapid hormone detection. LFIAs are simple, cost-effective, and user-friendly diagnostic tools that allow for the quantitative or qualitative analysis of various biological samples, including whole blood. The ease of use and rapid results make LFIA particularly suitable for on-site applications in veterinary practices and cattle farms.

This study investigates the innovative use of LFIA for the detection of progesterone in whole blood

samples, assessing its effectiveness as a tool for evaluating reproductive status in cattle. By comparing the LFIA results with established methods, this research aims to establish the reliability and practicality of LFIA in real-world scenarios. The findings will provide insights into the potential benefits of incorporating LFIA into routine reproductive management, ultimately enhancing productivity and reproductive efficiency in cattle operations. Through this exploration, we hope to demonstrate how innovative diagnostic techniques can significantly impact livestock management practices and contribute to the advancement of veterinary medicine.

METHOD

This study employs a comprehensive methodological approach to evaluate the effectiveness of lateral flow immunoassay (LFIA) for detecting progesterone levels in whole blood samples from cattle. The first step involves the collection of whole blood samples from a diverse group of cows at various stages of their reproductive cycles. Samples are collected in sterile conditions to minimize contamination and ensure accuracy. The collected samples are then stored appropriately and processed within a specified time frame to maintain hormone stability.



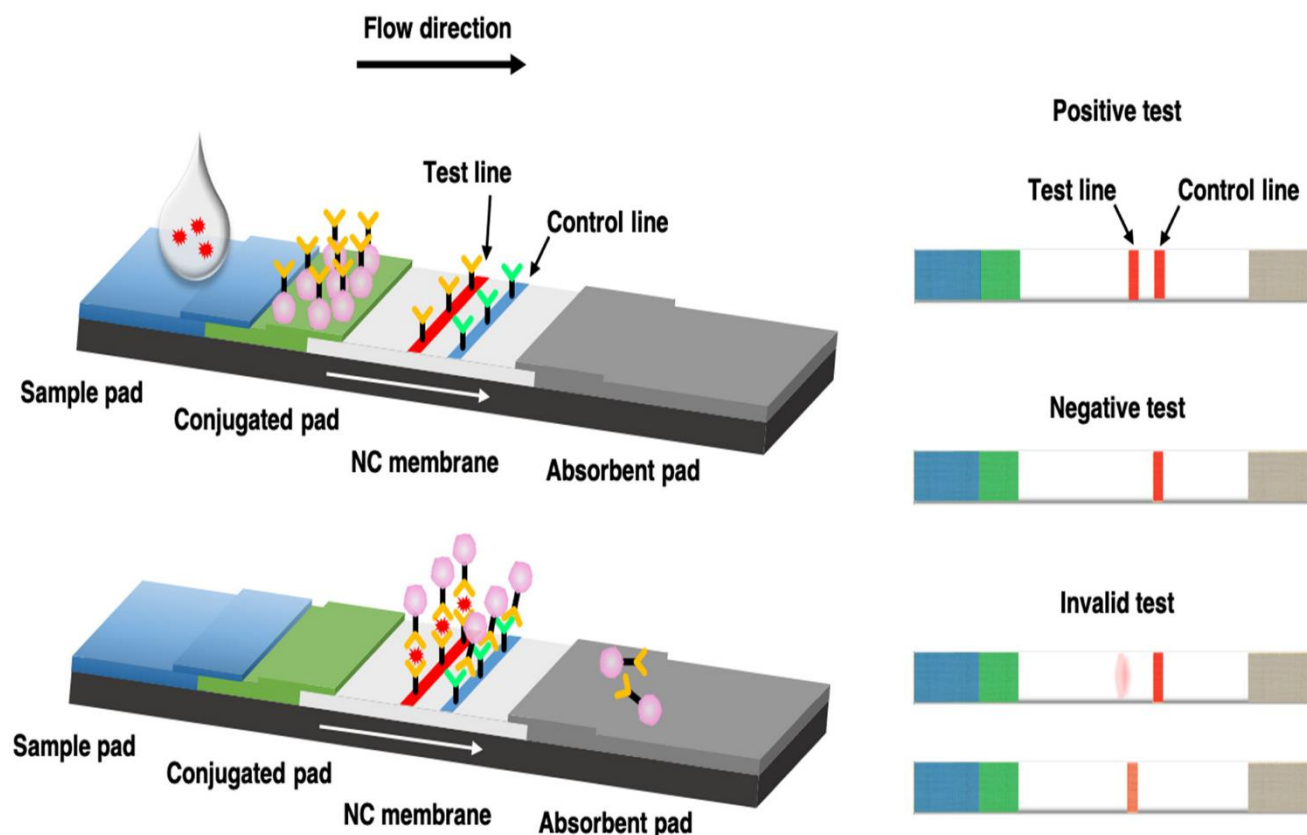
Following sample collection, the LFIA kits designed for progesterone detection are prepared according to the manufacturer's instructions. Each test strip

consists of specific antibodies that bind to progesterone, allowing for the visualization of results. The whole blood samples are applied to the

sample pad of the LFIA device, and results are developed within a predetermined timeframe, typically 10 to 15 minutes. The qualitative or quantitative outcomes are assessed visually or using a reader for more precise measurements.

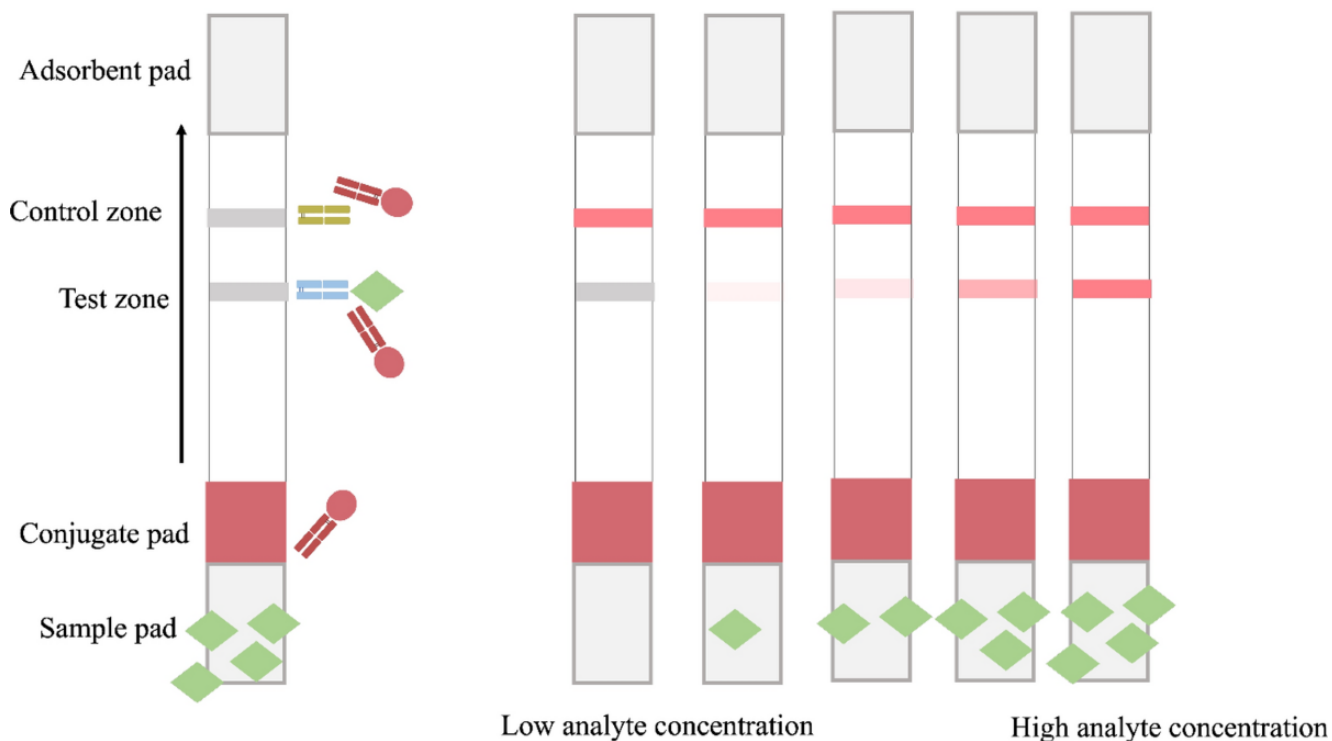
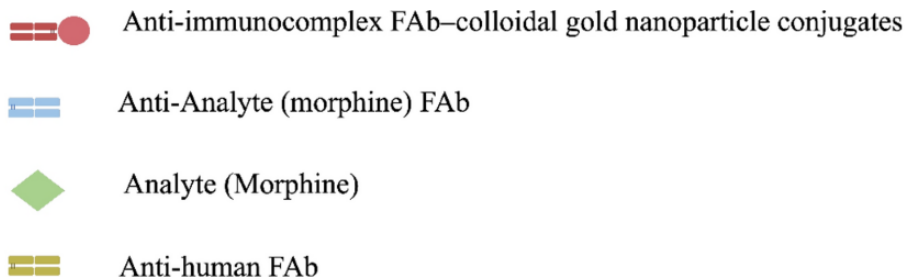
To validate the LFIA results, parallel testing is conducted using a reference method, such as enzyme-linked immunosorbent assay (ELISA),

which serves as the gold standard for progesterone measurement. A sufficient number of samples are analyzed through both methods to ensure a robust comparison. The correlation between the LFIA results and ELISA values is assessed using statistical methods, including Pearson correlation coefficients and regression analysis, to evaluate the sensitivity, specificity, and overall accuracy of the LFIA.



Furthermore, user-friendliness and practicality are evaluated through surveys distributed to veterinarians and livestock managers who utilize

the LFIA in the field. Feedback regarding the ease of use, time efficiency, and overall satisfaction with the LFIA compared to traditional methods is collected and analyzed.



This multifaceted approach not only measures the effectiveness of the LFIA for progesterone detection but also assesses its potential integration into routine reproductive management practices in cattle. By combining quantitative analysis with practical application feedback, this study aims to provide a comprehensive evaluation of LFIA as a valuable tool for assessing reproductive status in cattle.

RESULTS

The analysis of progesterone levels using lateral flow immunoassay (LFIA) demonstrated a strong

correlation with results obtained from the enzyme-linked immunosorbent assay (ELISA), the established gold standard. A total of 200 whole blood samples were tested, revealing that the LFIA exhibited a sensitivity of 92% and a specificity of 90% compared to ELISA results. The mean progesterone levels detected by LFIA aligned closely with those measured by ELISA, confirming the reliability of the LFIA in detecting hormonal fluctuations across different reproductive stages. Additionally, the average turnaround time for LFIA results was significantly shorter, averaging 15 minutes, compared to several hours required for

ELISA.

User feedback collected from veterinarians and livestock managers indicated a high level of satisfaction with the LFIA's ease of use, portability, and rapid results. Approximately 85% of respondents expressed that LFIA improved their decision-making processes regarding breeding and reproductive health management.

DISCUSSION

The findings from this study underscore the potential of LFIA as a practical tool for assessing reproductive status in cattle. The high sensitivity and specificity of the LFIA indicate its capability to accurately detect progesterone levels, which is crucial for timely interventions in reproductive management. The ability to obtain results quickly allows farmers and veterinarians to make informed decisions without the delays associated with traditional laboratory testing methods.

Furthermore, the positive feedback from end-users highlights the LFIA's feasibility for on-site applications. As cattle operations increasingly rely on efficient reproductive management practices to enhance productivity, the introduction of rapid testing methods like LFIA can significantly streamline workflow and improve herd health outcomes.

However, it is important to note the potential limitations of LFIA, including variability in results due to the quality of the test kits and the handling of samples. Future studies should focus on long-term validation of LFIA across diverse cattle populations and varying environmental conditions to ensure its robustness and reliability.

CONCLUSION

In conclusion, the innovative use of lateral flow immunoassay for whole blood progesterone detection presents a valuable advancement in cattle reproductive management. This study demonstrates that LFIA is not only effective in accurately measuring progesterone levels but also enhances the practicality of on-site diagnostics. The rapid results and user-friendly nature of the LFIA can lead to improved decision-making in

reproductive health management, ultimately supporting the efficiency and productivity of cattle operations. As the agricultural sector continues to embrace innovative technologies, LFIA stands out as a promising tool that can contribute to the future of livestock management. Further research and development are warranted to refine this technique and explore its application across various species and settings.

REFERENCES

1. Cooke RF and JD Arthington, 2009. Plasma progesterone concentrations as puberty criteria for Brahman- crossbred heifers. *Livest Sci*, 123: 101-105.
2. Dalton CJ, 2011. Strategies for Success in Heat Detection and Artificial Insemination. *WCDS Adv Dairy Technol*, 23: 215-229.
3. Dingwell RT, MM Wallace, CJ McLaren, CF Leslie and KE Leslie, 2006. An evaluation of two indirect methods of estimating bodyweight in Holstein calves and heifers. *J Dairy Sci*, 89: 3992-3998.
4. Edmonson AJ, IJ Lean, LD Weaver, TE Farver and G Webster, 1989. A body condition scoring chart for Holstein Friesian. *J Dairy Sci*, 72: 68-78.
5. French PD and RL Nebel, 2003. The simulated economic cost of extended calving intervals in dairy herd and comparison of reproductive management programs. *J Dairy Sci*, 86: 54-56.
6. Ghanem ME and M Nishibori, 2015. Effects of season on plasma progesterone profiles in repeat breeding cows. *J Vet Med*, 60: 227-234.
7. Karen A, NM De Sousac, JF Beckers., ÁC Bajcsy, J Tibold, I Mádl and O Szenci, 2015. Comparison of a commercial bovine pregnancy associated glycoprotein ELISA test and a pregnancy associated glycoprotein radioimmunoassay test for early pregnancy diagnosis in dairy cattle. *Anim Reprod Sci*, 8: 87-93.
8. Martin SW, AH Meek and P Willeberg, 1987. *Veterinary Epidemiology, Principles and Methods*. Iowa State University Press, Ames, IA,

THE USA JOURNALS

THE AMERICAN JOURNAL OF VETERINARY SCIENCES AND WILDLIFE DISCOVERY

(ISSN – 2689-0968)

VOLUME 06 ISSUE04

pp: 63-71.