



OPEN ACCESS

EDITED BY

Mark D. Tarn,
The University of Leeds, United Kingdom

REVIEWED BY

Zibusiso Ndlovu,
Médecins Sans Frontières, South Africa

*CORRESPONDENCE

Collins Otieno Odhiambo,
✉ cotieno@aslm.org

RECEIVED 27 August 2025

ACCEPTED 25 September 2025

PUBLISHED 08 October 2025

CITATION

Odhiambo CO, Van Puije B, Krisiunas E and
Mataka A (2025) The environmental price of
diagnostic scale-up—addressing a toxic trade-
off we cannot afford.
Front. Lab Chip Technol. 4:1693514.
doi: 10.3389/frlct.2025.1693514

COPYRIGHT

© 2025 Odhiambo, Van Puije, Krisiunas and
Mataka. This is an open-access article
distributed under the terms of the [Creative
Commons Attribution License \(CC BY\)](#). The use,
distribution or reproduction in other forums is
permitted, provided the original author(s) and
the copyright owner(s) are credited and that the
original publication in this journal is cited, in
accordance with accepted academic practice.
No use, distribution or reproduction is
permitted which does not comply with these
terms.

The environmental price of diagnostic scale-up—addressing a toxic trade-off we cannot afford

Collins Otieno Odhiambo^{1*}, Beatrice Van Puije¹,
Edward Krisiunas² and Anafi Mataka¹

¹African Society for Laboratory Medicine, Addis Ababa, Ethiopia, ²WNWN International, Inc., Burlington, CT, United States

The scale up of diagnostic service in many Low-Income Countries (LIC) has greatly improved health outcomes of the populations being served by 21st century laboratory platforms like those seen in High Income Countries (HIC). A challenge because of this scale up is the contribution of increased volume of plastic and hazardous chemical waste that needs to be properly managed with limited options as compared to what is available in HIC. Guanidinium thiocyanate (GTC), as an example, a widely used component of extraction reagents used in Polymerase Chain Reaction (PCR) testing, when inappropriately disposed of can cause harm to animals and the environment because it contains a toxic cyanide compound. While environmental short term disposal methods exist that would improve the practices and offer alternative waste treatment options, challenges in availability of required infrastructure and sustainability remain, thereby limiting their impact. While testing generates revenue, waste associated with testing is a cost that is not adequately funded hence unsustainable in the current environment. The availability of alternative compounds that are less toxic yet can achieve the required actions may provide a lasting solution. Developing platforms geared towards microchemistry as well as the recyclability of the materials used for testing would steer waste/material management in a new direction. We call on diagnostic manufacturers to consider these options upstream of the product lifecycle in the long term that are in line with their internal corporate commitment to sustainability beyond the manufacturing process.

KEYWORDS

waste management, waste disposal, GTC, guanidinium thiocyanate, hazardous waste and management

Introduction

Every diagnostic test conducted has the potential to save lives- but improper disposal of the waste streams (personal protective equipment, blood drawing needles, tubes of blood, pipette tips, platform analyzer tubes, plastic pasture pipettes, Point of Care (POC) cartridges, chemical used in diagnostics platforms of varying hazards) emanating from these tests can result in environmental pollution (Poor air emissions from inappropriately operated incinerators, contaminated/toxic waste water, solid residues from poorly or partially treated wastes) posing a danger to both humans and animals. The scale up in diagnostics while intended to reduce the gap in diagnostic access, must be paralleled with implementation of fit for purpose (versus fit for budget) and sustainable waste management innovations and interventions such as bioremediation and retrofitting incinerators with



FIGURE 1
Viral Load lab waste waiting to be treated and destroyed.

fluid injectors to manage fluid treatment. The scale up of molecular tests to address the diagnostic gaps has the potential to improve health outcomes especially in sub-Saharan Africa, where the unmet need accounts for up to 81% (Fleming et al., 2021). However, we must not ignore that this scale up is accompanied by an increase in the generation of infectious waste, toxic plastics components, and chemicals from the testing platforms (Figure 1). For example, guanidinium thiocyanate (GTC), a strong protein denaturant and a chaotropic agent is an essential component in the functionality of molecular viral load, COVID-19 and other molecular diagnostics. Yet when improperly disposed of, especially when mixed with bleach, it releases cyanide gas potentially posing serious health and environmental hazards (Paik and Wu, 2005; Welch et al., 2020). Options for treatment and disposal are discussed below.

Waste burden and inappropriate practices

Deducing from the estimates by Sleeman et al., over 36 million HIV viral load (VL) tests were conducted globally in 2024, generating approximately 1.1 million litres of effluent chemical waste and 2.5 million kilograms of solid waste annually (Sleeman et al., 2018). The COVID-19 testing surge magnified this issue, with over 140 million supplied kits producing around 2.6 million kilograms of plastic waste and 731,000 L of chemical waste (WHO, 2022). This risk is not theoretical, as demonstrated in a 2020 survey of liquid waste management practices conducted by the African Society for Laboratory Medicine (ASLM) across 11 African countries participating in the laboratory systems strengthening community of practice. The survey reported disposal of GTC-containing liquid chemical waste into the sewer in most viral load testing laboratories thereby creating substantial environmental risks to both humans and animals. Additionally, the 11 countries self-assessed their HIV viral load testing cascades

(Odhiambo et al., 2021). Waste management emerged as the weakest domain due to systemic gaps related to lack of national waste management policies and guidelines, insufficient biosafety infrastructure, and the absence of sustainable waste solutions. Only 2 out of 11 countries had the requisite national waste management and biosafety policies and guidelines document (Odhiambo et al., 2021). There is a lack of understanding of the principles of healthcare waste management along with a lack of understanding of the principles of operating and maintaining treatment technologies for the waste streams to be processed. The issue is particularly acute in laboratories using molecular platforms, where chemical handling is more complex and often overlooked. Most lacking is addressing the financial needs associated with healthcare waste management with encompasses maintenance of the appropriately selected equipment as well the cost operating the equipment (staff salaries, fund spare parts, and required utilities). Waste management is a cost center and not a revenue generator like laboratory testing. Opportunities for cost avoidance do exist that need to be explored on a case-by-case basis in collaboration with platform vendors.

Initiative to support short term solution

To urgently address this challenge, the U.S. Centers for Disease Control and Prevention and the ASLM launched a targeted initiative: a waste management sub-community of Practice. This initiative brought together subject matter experts, country teams and technical consultants from Waste Not, Want Not (WNWN) to co-develop solutions. Together, we raised awareness about GTC toxicity, assessed disposal practices, and explored sustainable alternatives to incineration, in the short term. During this initiative, several promising alternatives to the widespread practice of discharging to sewer or incinerating GTC-containing waste were identified (ASLM, 2025). These include: a)

mixing liquid waste with sawdust or charcoal dust to create material more suitable for thermal combustion, (Potentially low cost based on sawdust or charcoal dust availability) b) encapsulation of liquid waste with cement, (Low cost for small volumes) c) Use of chemicals to precipitate the guanidine thiocyanate as cuprous thiocyanate thereby greatly reducing the liquid fraction to a non-hazardous waste, (Medium cost) d) Use of cement kilns that are amenable to liquid and solid fractions of the waste streams, e) Use of UV light and hydrogen peroxide to break down the thiocyanate fraction of GTC (High cost initially for setting up equipment/most appropriate for large volumes of liquid (>100 L per day), f) bioremediation using bacteria capable of breaking down thiocyanate, (Moderate cost to set up but self-sustaining once established) and g) incinerator fluid injection systems that sprays/mists the liquid waste into the primary chamber of an incinerator (High cost for injector/suitable for large incinerators (>100 kg/h processing capacity). Additionally, many facilities used incinerators that were sub-optimally functional as well as not designed for liquid waste disposal, and in some instances, basic burning chambers mistakenly identified as incinerators. These findings demonstrated there are a range of treatment options available. It also pointed to the need to budget for these as well as other waste management practices. Without appropriate funding, none of these options are sustainable.

Overall, there has been an improvement in waste management practices across the 11 countries exemplified by the increased number of countries with at least a draft of the national waste management and biosafety policies and guidance documents (n = 9) from the last round of self-assessment exercises conducted in October 2024. Despite the availability of policies and guidance documents and the improvement of waste management practices in molecular testing laboratories, there are still lingering questions on the sustainability of these initiatives.

Innovations for long term sustainability

While the waste management alternatives mentioned above can apply in the short term to reduce environmental pollution across sub-Saharan African countries, there is need for more sustainable solutions particularly targeted upstream in diagnostic manufacturing. Some successful innovations have been demonstrated, e.g., in eliminating the use of GTC for nucleic acid extraction in molecular testing. For example, Abbott and Hologic diagnostic manufacturers use Guanidine Hydrochloride and sodium hypochlorite for their mPima and Panther VL tests, respectively as the nucleic acid extraction agent (WHO, 2017; WHO, 2019; WHO, 2021). The use of environmentally friendly material components could allow for less reliance on virgin petrochemical materials that contribute to increasing the carbon footprint through green house gas emissions. Ongaro et al. provided a review of alternative environmental-friendly materials that can be used to replace commonly used materials in the manufacturing of point-of-care tests (Ongaro et al., 2022). These include the use of recyclable plastics, bio-derived and bio-degradable plastics, natural fibrous materials, among others. Availability of these alternatives should inform advocacy efforts to

manufacturers to consider re-designing existing tests to include these environmentally friendly options in the production process. Global laboratory stakeholders like the Global Fund need to enforce the requirement for environmentally friendly alternatives in newer products in addition to stronger legislations and regulations at country level, all contributing to actualizing the 'polluter pays' principle (Gorun, 2018). A holistic approach must be considered by all stakeholders for these waste streams—the application of 10 Rs of waste management would be a good place to begin from most to least effective, Refuse, Rethink, Reduce, Reuse, Repair, Refurbish, Remanufacture, Repurpose, Recycle, and Recover (Munoz et al., 2024).

Call to action

In conclusion, we call upon the waste management stakeholders, including ministries of health, donors, and diagnostic manufacturers to act now, before the sector is flooded with a large numbers of kits produced without environmental considerations. National procurement guidelines must include the requirement for safe disposal of diagnostic products. Target Product Profiles should be put to good use by policymakers and must incorporate quantitative limits to the packaging proportion in testing kits, as well as minimize plastic use, especially in standard kits such as lateral flow tests (Wöhrle et al., 2025). Annual procurement forecasting exercises should include anticipated waste volumes for accurate cost of disposal. Infrastructure investments should prioritize biosafety and meet the requirements for safe waste disposal—not just throughput. Diagnostic manufacturers must take greater responsibility for end-of-life management of their products and reagents including consideration for sustainable, less toxic, and potentially recycled, alternative raw materials.

While championing equitable access to diagnostics to meet the needs of the African continent, we must also champion the right to safety—for healthcare workers, communities, and the environment. Diagnostic access without safe, effective treatment and disposal of these waste streams is a toxic trade-off we can no longer afford.

Data availability statement

The original contributions presented in the study are included in the article/supplementary material, further inquiries can be directed to the corresponding author.

Author contributions

CO: Conceptualization, Writing – original draft, Writing – review and editing. BV: Writing – review and editing. EK: Writing – review and editing. AM: Writing – review and editing.

Funding

The author(s) declare that no financial support was received for the research and/or publication of this article.

Conflict of interest

Author EK was employed by Waste Not Want Not inc.

The remaining authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

The reviewer ZN is currently organizing a Research Topic with the author CO.

Generative AI statement

The author(s) declare that no Generative AI was used in the creation of this manuscript.

References

- ASLM. (2025). ASLM LabCoP waste management ECHO sessions. Available online at: https://www.youtube.com/playlist?list=PLw-gn36_-CCK8VIFCQFHSRpHDB5bG5uDV (Accessed 18 August 2025).
- Fleming, K. A., Horton, S., Wilson, M. L., Atun, R., DeStigter, K., Flanigan, J., et al. (2021). The lancet commission on diagnostics: transforming access to diagnostics. *Lancet* 398 (10315), 1997–2050. doi:10.1016/s0140-6736(21)00673-5
- Gorun, D. (2018). Theoretical and practical aspects regarding the applying of the principle “Polluter Pays”. *WSEAS Trans. Environ. Dev.* 14, 481–494.
- Munoz, S., Hosseini, M. R., and Crawford, R. H. (2024). Towards a holistic assessment of circular economy strategies: the 9R circularity index. *Sustain. Prod. Consum.* 47, 400–412. doi:10.1016/j.spc.2024.04.015
- Odhiambo, C. O., Mataka, A., Kassa, G., and Ondoa, P. (2021). Managing laboratory waste from HIV-related molecular testing: lessons learned from African countries. *J. Hazard. Mater. Lett.* 2, 100030. doi:10.1016/j.jhazl.2021.100030
- Ongaro, A. E., Ndlovu, Z., Sollier, E., Otieno, C., Ondoa, P., Street, A., et al. (2022). Engineering a sustainable future for point-of-care diagnostics and single-use microfluidic devices. *Lab a Chip* 22 (17), 3122–3137. doi:10.1039/d2lc00380e
- Paik, S., and Wu, X. (2005). Measuring toxic gases generated from reaction of guanidine isothiocyanate-containing reagents with bleach. *Chem. Health and Saf.* 12 (4), 33–38. doi:10.1016/j.chs.2004.12.002
- Sleeman, K., Hurlston, C. M., Diallo, K., Zeh, C., Riley, P., and Alexander, H. (2018). “Strengthening waste management during HIV viral load scale-up,” in ASLM 2018 International Conference, Abuja, Nigeria, 10–14.
- Welch, S. R., Davies, K. A., Buczkowski, H., Hettiarachchi, N., Green, N., Arnold, U., et al. (2020). Analysis of inactivation of SARS-CoV-2 by specimen transport media, nucleic acid extraction reagents, detergents, and fixatives. *J. Clin. Microbiol.* 58 (11), e01713–e01720. doi:10.1128/jcm.01713-20
- WHO (2017). *Prequalification of diagnostics: Abbott mPIMA HIV-1/2 v5.0 and hologic aptima HIV-1 quant Dx assay*. Geneva: WHO, 2019.
- WHO (2019). *Prequalification of diagnostics: Abbott mPIMA HIV-1/2 v5.0 and hologic aptima HIV-1 quant Dx assay*. Geneva: WHO.
- WHO (2021). *Prequalification of diagnostics: Aptima HIV-1 quant Dx assay*. Geneva: WHO.
- WHO (2022). *Global analysis of healthcare waste in the context of COVID-19: status, impacts and recommendations*. World Health Organization. Available online at: <https://www.who.int/publications/i/item/9789240039612>.
- Wöhrle, M. L., Street, A., and Kersaudy-Kerhoas, M. (2025). Mass of components and material distribution in lateral flow assay kits. *Bull. World Health Organ.* 103 (4), 236–246. doi:10.2471/blt.24.292167

Any alternative text (alt text) provided alongside figures in this article has been generated by Frontiers with the support of artificial intelligence and reasonable efforts have been made to ensure accuracy, including review by the authors wherever possible. If you identify any issues, please contact us.

Publisher’s note

All claims expressed in this article are solely those of the authors and do not necessarily represent those of their affiliated organizations, or those of the publisher, the editors and the reviewers. Any product that may be evaluated in this article, or claim that may be made by its manufacturer, is not guaranteed or endorsed by the publisher.