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Tools for change: Quantifying and Understanding Diagnostic Plastic Waste

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ASLM Waste Sub-CoP session
24th March 2026



Photograph: Edward Krisiunas / ASLM



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Outline

- POCT Waste: Scale of the problem
- Material usage in POCTs
- Measuring single-use laboratory waste
 - *Survey*
- Introducing Sustainable materials in POCTs
- Conclusions

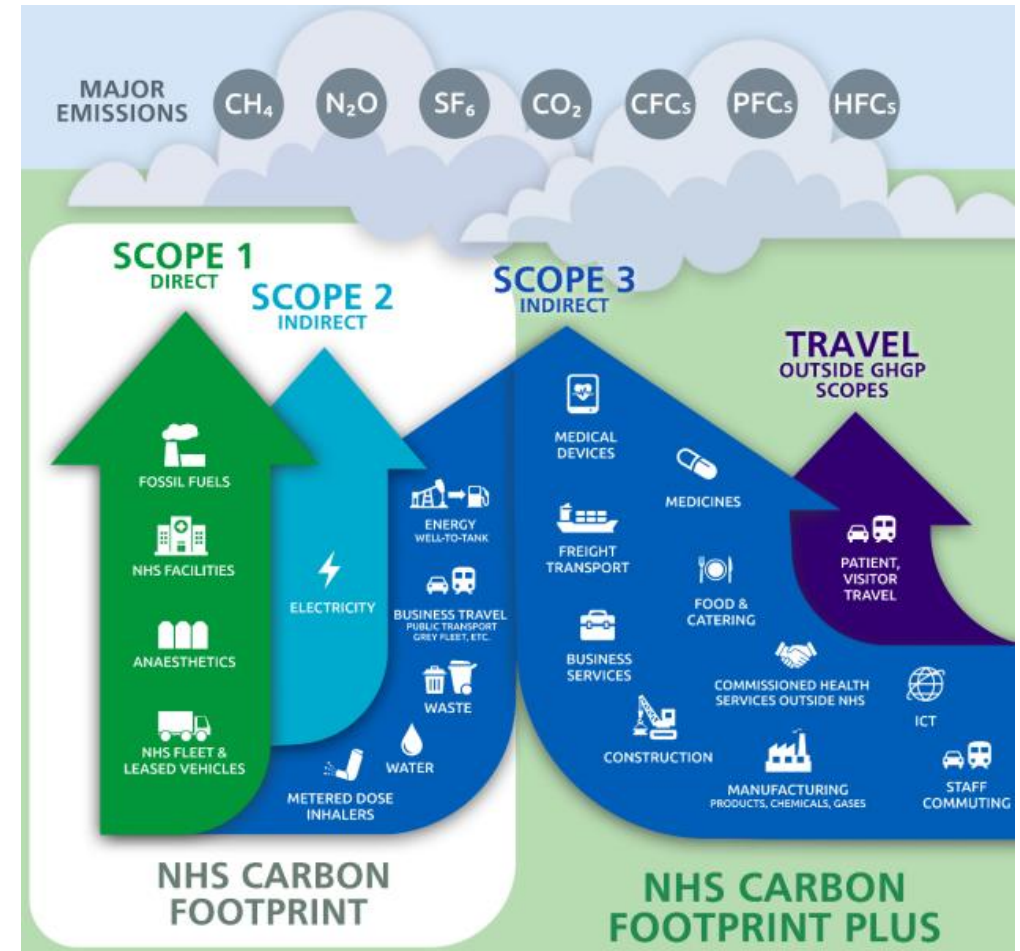


Photograph: Edward Krisiunas / ASLM



Why are we interested in medical waste?

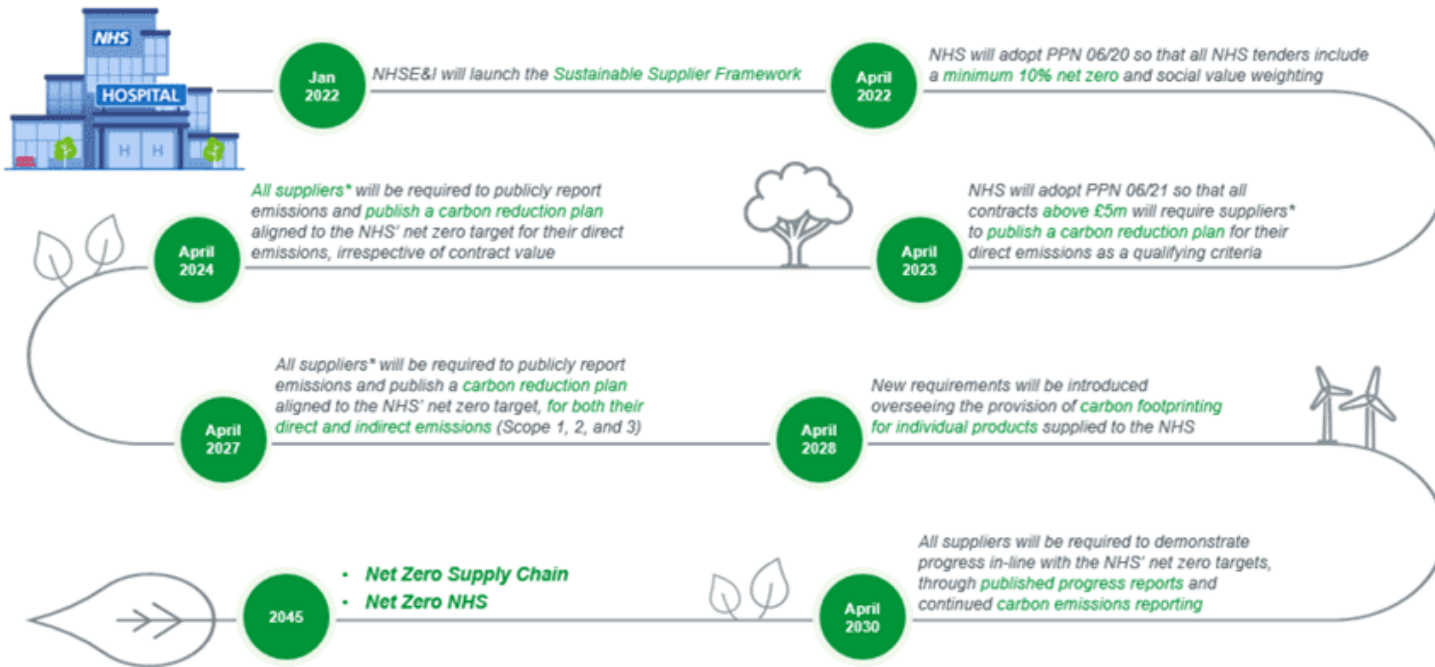
- The healthcare sector is responsible for ~4% of global carbon emissions
- Approximately 6–10% of national health system emissions come from medical devices
- In the UK, strong emphasis on decarbonization
- Most carbon emissions embedded in “scope 3” emissions





Why are we interested in medical waste?

Building net zero into NHS procurement



- From 1 April 2022, all NHS procurements will include a minimum 10% net zero and social value weighting.
- From April 2028, new requirements will be introduced **overseeing the provision of carbon footprinting for individual products** supplied to the NHS. The NHS will work with suppliers and regulators to determine the scope and methodology.

*To account for the specific barriers that Small & Medium Enterprises and Voluntary, Community & Social Enterprises encounter, a two-year grace period on the requirements leading up to the 2030 deadline, by which point we expect all suppliers to have matched or exceeded our ambition for net zero.



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DiaDev
Investigating Diagnostics in Global Health

Research: Diagnostic waste flows Sierra Leone



Qualitative research: Mapping the life-cycle of diagnostic devices



Surveys: Waste management infrastructure at point of testing



Key findings

- LFAs used at every level of health system, laboratories and clinics
- 47% of Community Health Centres had functioning incinerators
- Segregated health care waste was re-mixed at point of discard
- Mixed health care waste was openly burned, buried, or sent to landfill.



Street, A., Vernooij, E., & Rogers, M. H. (2022). Diagnostic waste: whose responsibility?. *Globalization and Health*, 18(1), 30.

Street, A., Vernooij, E., Koker, F., Baxter, M. S., Bah, F., Rogers, J., ... & Ansumana, R. (2023). The “ready-to-hand” test: Diagnostic availability and usability in primary health care settings in Sierra Leone. *PLOS global public health*, 3(2), e0000604.



Key Lessons

- Global health system prioritises single-use devices and deprioritizes waste infrastructure
- Costs of diagnostic waste are ‘externalized’ to users
- Focus on ‘speed’ and ‘access’ removes incentives for manufacturers, regulators, policy makers to consider impact of waste.

COMMENTARY

Open Access

Diagnostic waste: whose responsibility?

Alice Street^{1*}, Eva Vernooij¹ and Mohamed Hashim Rogers²

Abstract

Waste management is notably absent from current discussions about efforts to improve access to diagnostics in low- and middle-income Countries (LMICs). Yet an increase in testing will inevitably lead to an increase in diagnostic waste, especially since many of the diagnostic tests designed for use in LMICs are single-use point-of-care tests. Diagnostic waste poses a threat to both human and environmental health. In this commentary we draw on our experience of diagnostic waste management in Sierra Leone and review current evidence on: the volume and impact of diagnostic waste in LMICs, existing health-care waste management capacity in LMICs, established national and international policies for improving health-care waste management, and opportunities for strengthening policy in this area. We argue that questions of safe disposal for diagnostics should not be an afterthought, only posed once questions of access have already been addressed. Moreover, responsibility for safe disposal of diagnostic waste should not fall solely on national health systems by default. Instead, consideration of the end-life of diagnostic products must be fully integrated into the diagnostic access agenda and greater pressure should be placed on manufacturers to take responsibility for the full life-cycle of their products.

Keywords: Diagnostics, Waste management, Sustainability, Responsibility, Product development

Background

‘Discard sample and assay waste according to your local safety regulations’. So states the standard small print found at the bottom of countless manufacturer labels for single-use diagnostic products. But what if no local or national guidelines or regulations exist? Or what if those guidelines exist on paper, but there is no infrastructure, systems, or workforce in place to enable health facilities to adhere to them? Since the start of the COVID-19 pandemic, the diagnostics pillar of the Access to COVID-19 Tools (ACT) Accelerator partnership has procured over 116.9 million tests for use in low-and-middle-income countries (LMIC), of which 43.2 million were polymerase chain reaction (PCR) tests and 73.7 million antigen-detecting rapid diagnostic tests (Ag-RDT) [20]. This achievement represents valuable progress in efforts to improve global access to life-saving diagnostic tools in

the pandemic. But with the global health community’s attention focused on questions of diagnostic access, far less consideration has been given to the end-life of diagnostic products [13, 15].

Cartridge-based polymerase chain reaction (PCR) tests and Ag-RDT tests [lateral flows] used to detect the SARS-CoV-2 virus are single-use devices. Their self-contained, easy-to-use format makes them especially promising for the extension of testing to places with limited laboratory infrastructure, and for the decentralisation of community testing. Yet, each one of those 116.9 million tests has also generated plastic, infectious, and potentially toxic waste that must be disposed of safely after use. Diagnostic waste poses multiple threats to human and environmental health [21, 24]. Ag-RDT cassettes are made from petroleum-derived plastics, which have a large CO₂ footprint and are non-biodegradable. A recent study estimated that PCR testing for COVID-19 generated 15,000 tons of plastic waste globally to August 2020 [5]. A World Health Organization (WHO) report on COVID-19 related healthcare waste calculated that test kits procured solely through the

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Research: Material components of LFAs



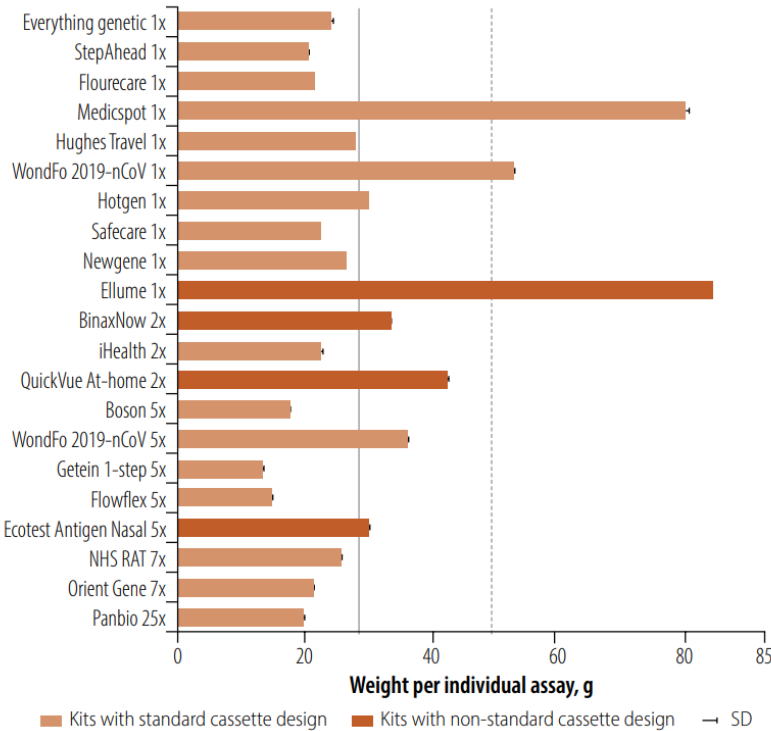
- 22 Covid-19 LFAs
- Manually deconstruct
- Weigh components

Wöhrle, M.-L., Street, A., & Kersaudy-Kerhoas, M. (2025). Mass of components and material distribution in lateral flow assay kits, *WHO Bulletin* 103 : 236–246. <http://dx.doi.org/10.2471/BLT.24.292167>

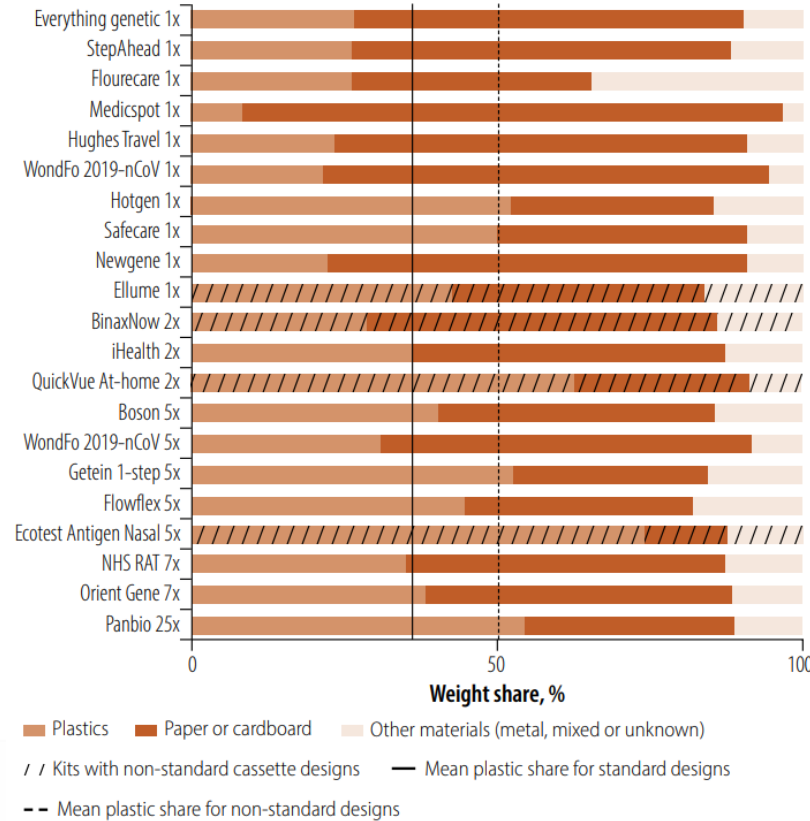


Key findings

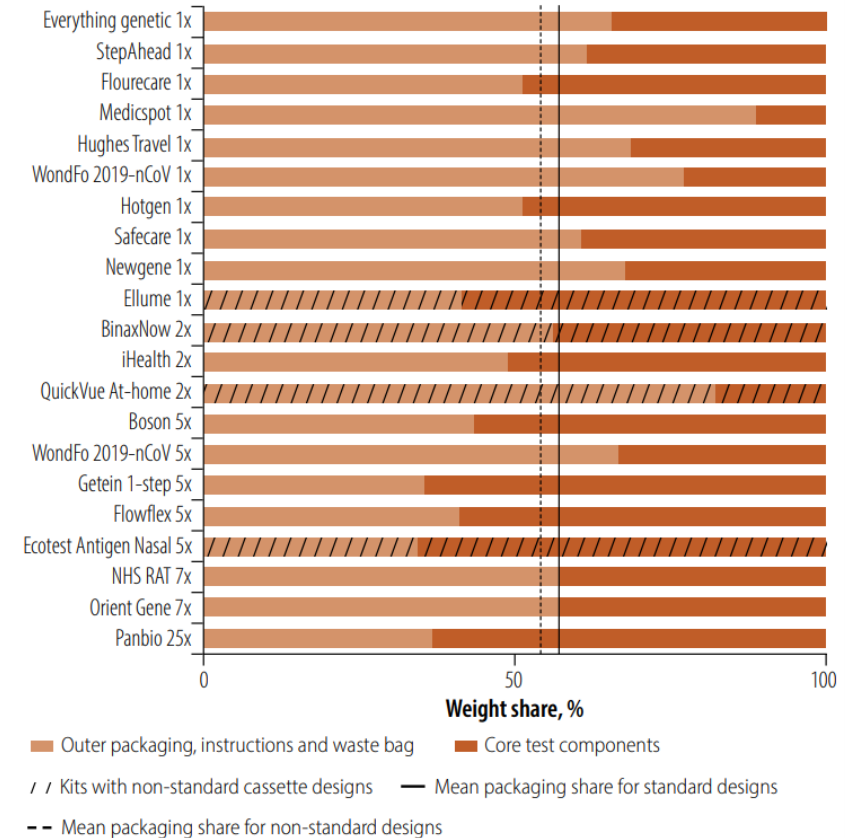
Wöhrle, M.-L., Street, A., & Kersaudy-Kerhoas, M. (2025). Mass of components and material distribution in lateral flow assay kits, *WHO Bulletin* 103 : 236–246.
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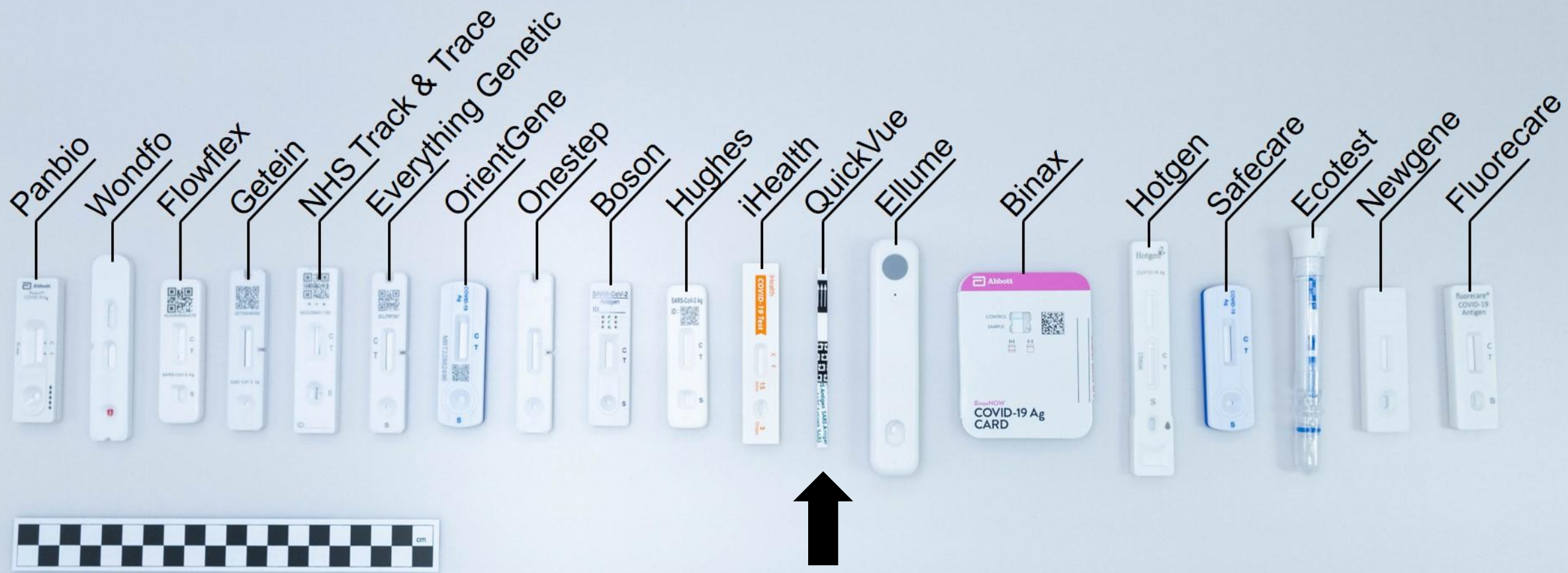
Large weight variations between kits



Large variations in plastic 'weight share'



Packaging made up on average 57% of the whole test kit weight



Wöhrle, M.-L., Street, A., & Kersaudy-Kerhoas, M. (2025). Mass of components and material distribution in lateral flow assay kits, *WHO Bulletin* 103 : 236–246. <http://dx.doi.org/10.2471/BLT.24.292167>



Key lessons

- Lower waste options are available on market
- Easy wins available in packaging
- Case for including environmental criteria in TPPs
- Shift responsibility from the user to the regulator and manufacturer
- Role for procurement





Research: Materials analysis



Materials data not publicly available



Deconstruct devices into individual material components



Use **Fourier-Transform Infrared spectroscopy (FTIR)** to identify materials



Key findings



- 20 components and 6 different plastics
- Presence of harmful materials and additives (PVC, Bisphenols, Polystyrene)
- Complex harmful packaging

Research carried out with Millie Marriott Webb and Alexandra Grosse

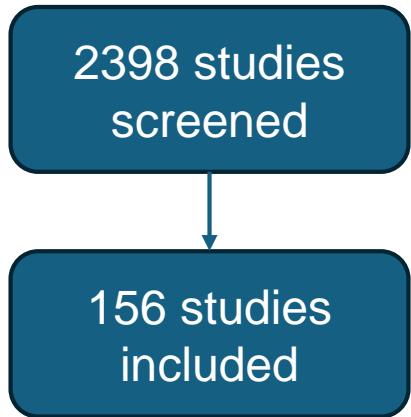


Key lessons

- Materials data is not publicly available
- Harmful materials – an issue beyond wet chemistry (e.g. GTC) and not confined to point of disposal.
- Rationale for materials not always self-evident
- Data can be fed into Life Cycle Analysis
- Need to expand analysis from ‘device’ to ‘procedure’

Measuring Laboratory waste

A meta-analysis by Slutzman et al, 2023



- Studies predominantly done in high-income countries (50%), followed by lower-middle-income (26%), upper-middle-income (20%), and low-income (5%) (Slutzman, 2023)
- Objectives for conducting the audit (all but 21 publications cited more than one reason):
 - To improve local waste sorting policies or practices (74%)
 - To reduce waste generation (45%)
 - To inform regulatory policy development (42%)
 - To increase or implement waste diversion (34%)
 - To save money on waste costs (31%)
 - Over 30 additional reasons identified including: quantifying greenhouse gas emissions, gathering inputs for LCA, creating waste prediction model, and quantifying total amounts of wastes
- Quantifying waste generation can provide the robust data that is currently lacking to enable effective implementation of waste reduction strategies at the financial, logistical, and legal levels.

Waste audits




	Use-based
Access	Difficult
Safety risk	High
Costs	High
Time duration	High
Accuracy	High
Disruption to services	High

Public data-base waste audit

Starting point: UKCAS document

Schedule of Accreditation
Issued by
United Kingdom Accreditation Service
2 Pine Trees, Chertsey Lane, Staines-Upon-Thames TW18 3HR


 8610 Accredited to ISO 15189:2022	NHS Tayside Issue No: 016 Issue date: 02 April 2025	
	Medical Microbiology Ninewells Hospital Ninewells Avenue Dundee DD1 9SY	Contact: Alison McPherson Tel: +44 (0)1382 660111 ext. 36718 E-Mail: alison.mcpherson@nhs.scot Website: http://www.nhstayside.scot.nhs.uk

Testing performed by the Organisation at the locations specified below

Locations covered by the organisation and their relevant activities

Laboratory locations:

Location details		Activity	Location code
Address Microbiology Department Ninewells Hospital Ninewells Avenue Dundee DD1 9SY	Local contact As above	Bacteriology Mycology Enteric Parasitology Virology Serology Molecular Microbiology	A
Address Sexual and Reproductive Health Clinic Ninewells Hospital Ninewells Avenue Dundee DD1 9SY	Local contact As above	Plate inoculation and microscopy	B

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- Public document
- Listing all assays in any UKCAS accredited lab
- Updated every year

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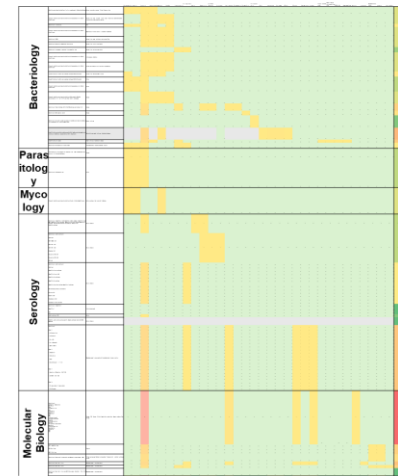
UKCAS document

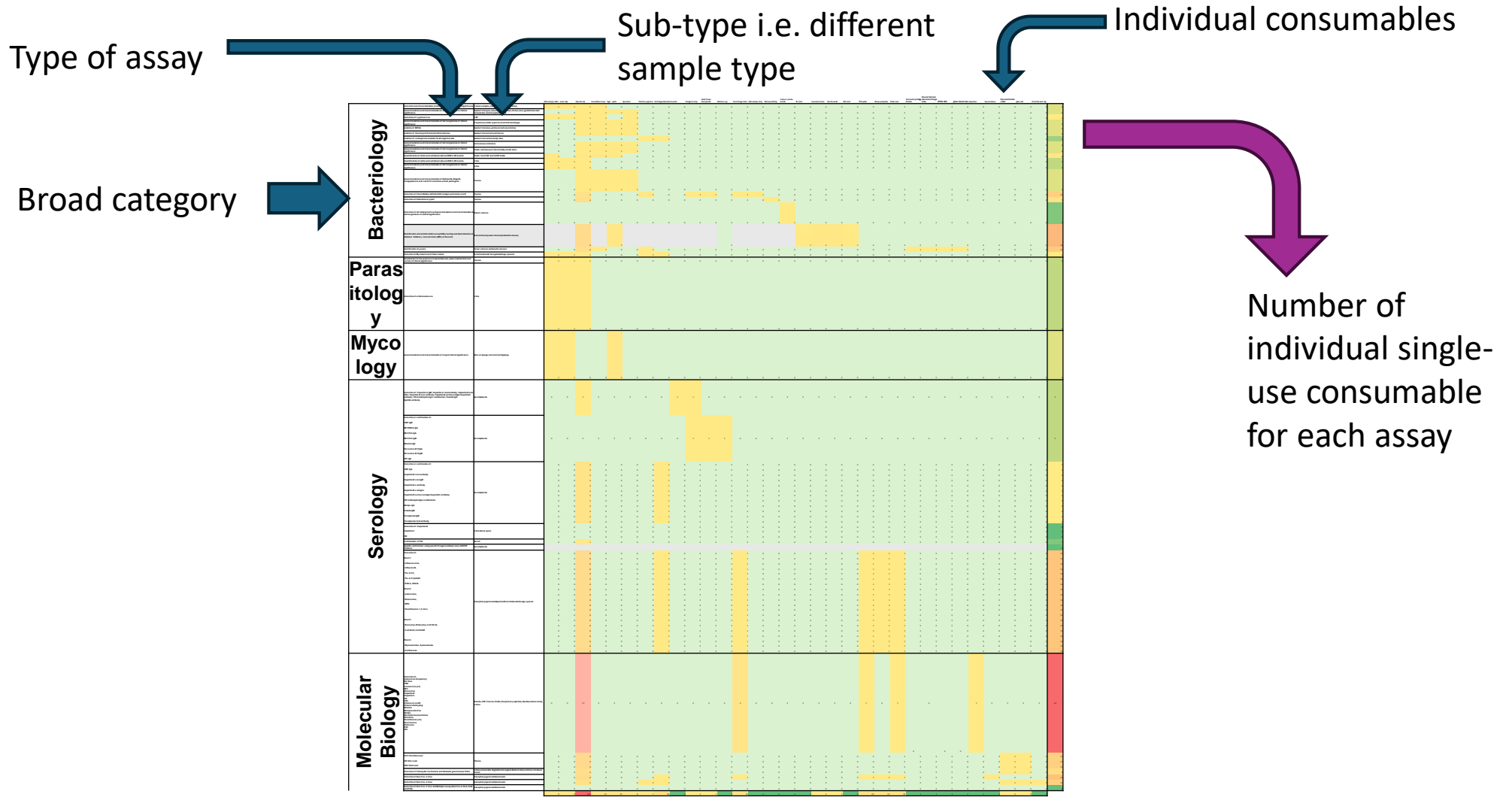
1. Extract all individual assays

2. Consult SOPs and manufacturer instructions

3. List single-use items for all assays

4. Produce heat map of single-use diagnostic consumable for the lab



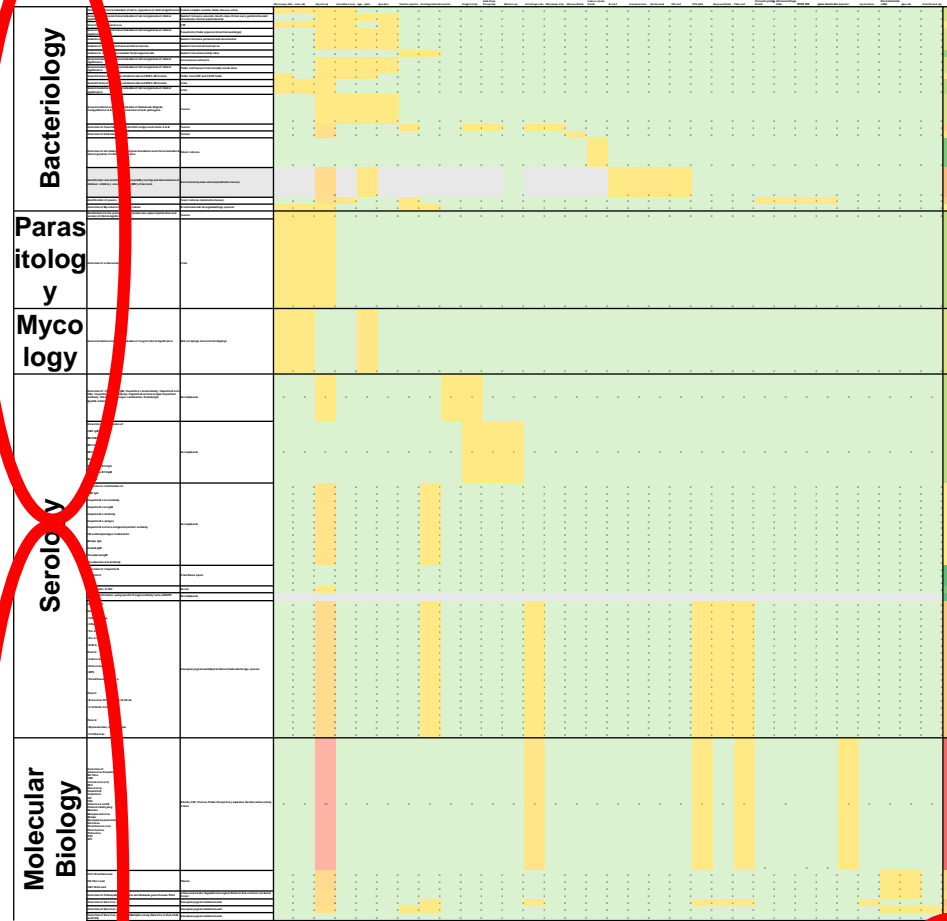


How to read the heat map table

Number of single-use consumable across assay menu

Bacteriology represents 24% of consumable use

Virology represents 75% of consumable use



Detection of viruses contain the most individual single-use consumables



Pipette tips are the single most present consumable across all

Example findings

Key lessons (1)

- Accessing physical waste is dangerous and time consuming
- Procurement approach has no single information points and can be politically fraught
- This audit format via public documentation enables a non-disruptive assessment of consumables used in any labs, but:
 - Is not suitable for waste management costing
 - does not cover ancillary equipment
 - relies on assay statistics for completeness
- Performs a different goal to Waste Cost Analysis Tool

Key lessons (2)

Purpose & scope

- Develop to enable understanding of waste generation by diagnostic tools
- Comparison of diagnostic tools
- Manufacturer agnostic
- Not for financial costing, although could be paired

Key features

- *Currently not operational*
- Concept tested on one lab in Scotland only
- Enable researchers to understand waste generated by assays

Potential Impact

- Support research in diagnostic waste
- Support lobbying manufacturers and policy makers for more sustainable products and sustainable targets

Future work

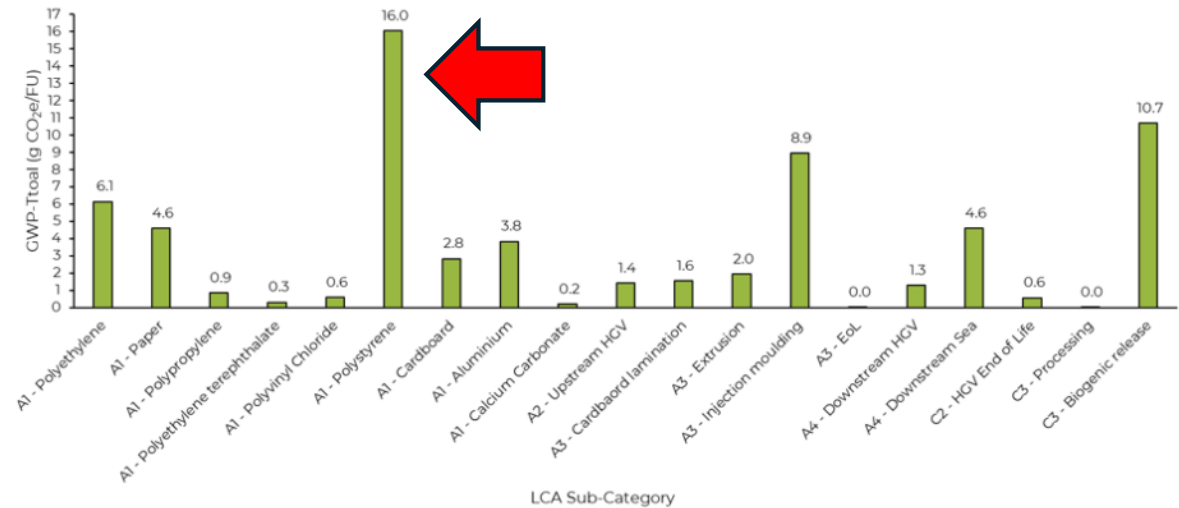
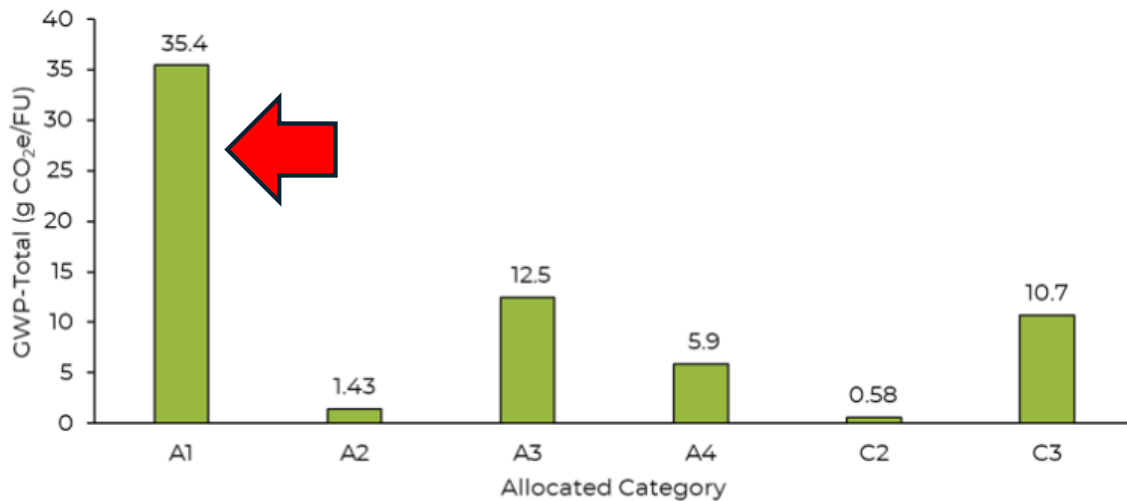
- Web-based platform development
- Test in different countries
- Test against 'use-base' audits

Take our short survey on
our proposed 'public
data' waste audit for
diagnostic lab



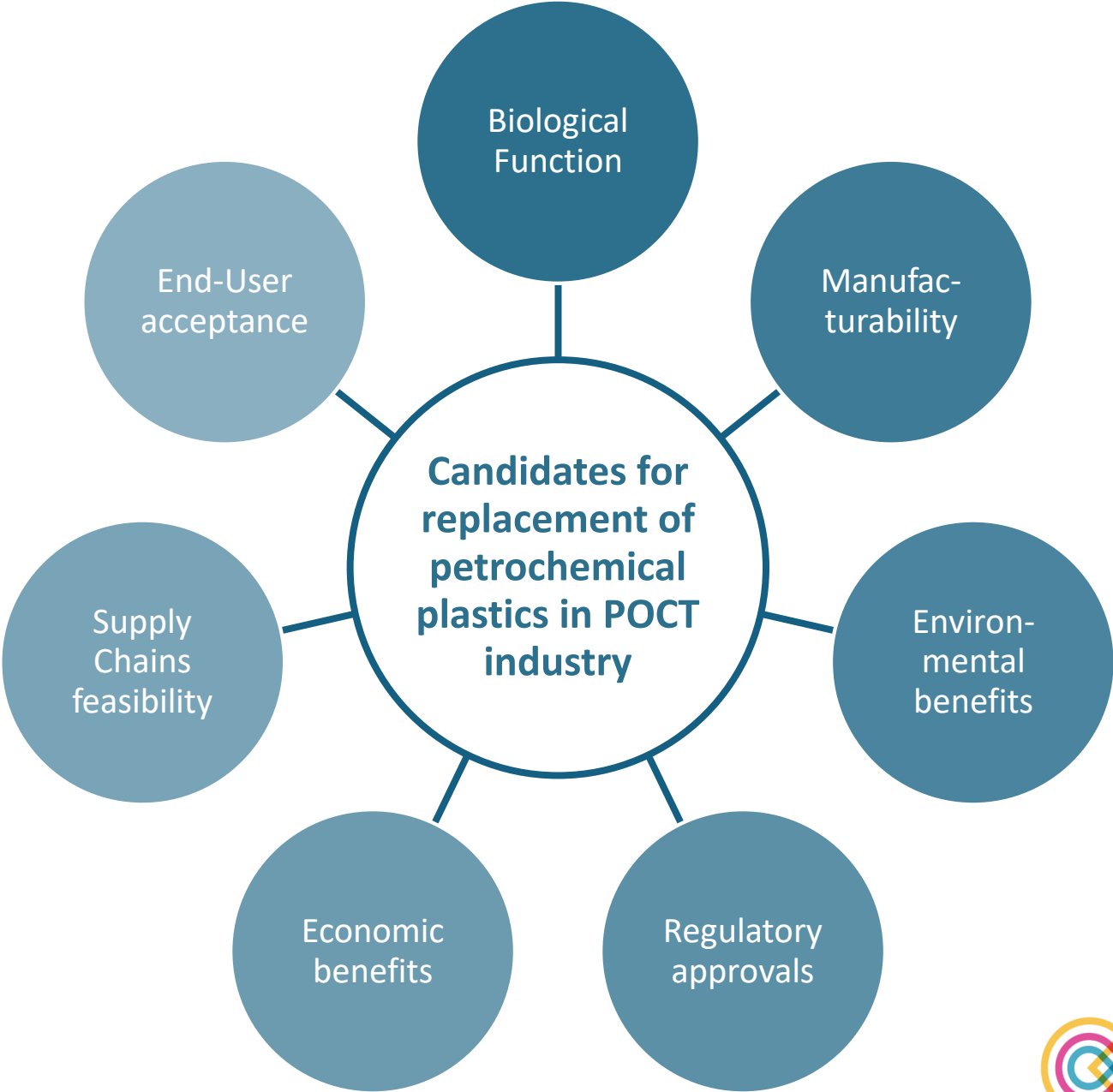
Introducing sustainable materials in diagnostic testing

- Purpose: Lower environmental impact of POCTs



- Solution: Replacing virgin petrochemical plastics, starting by the cassette

Introducing sustainable materials in diagnostic testing



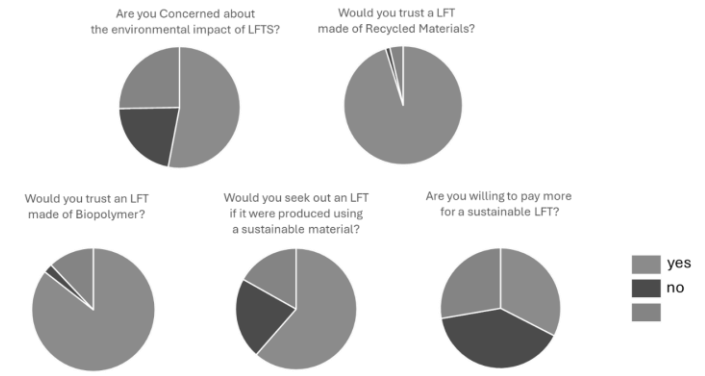
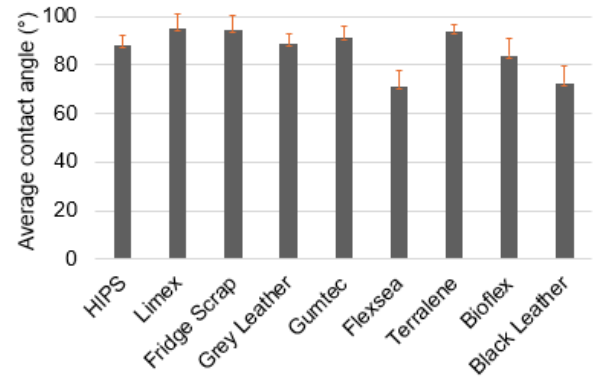
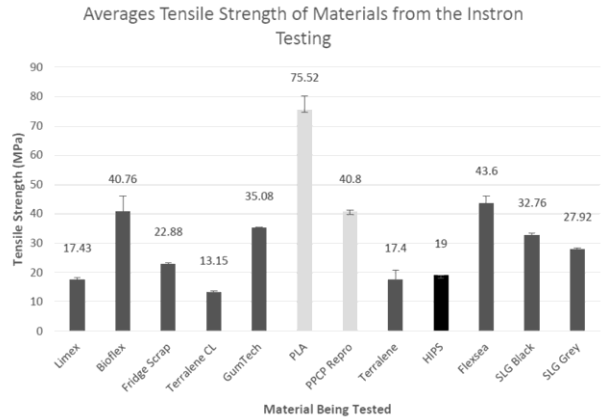
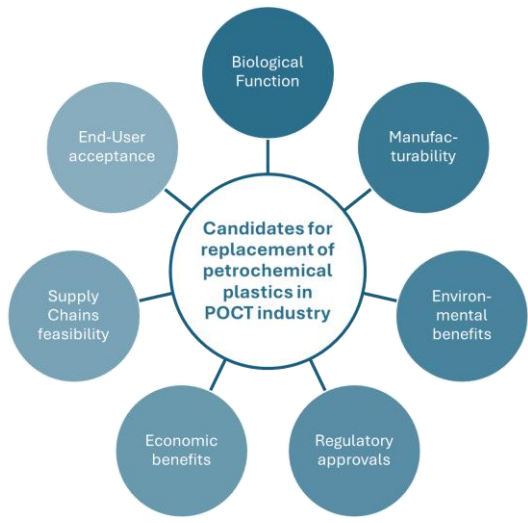
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After the Single Use
Rethinking Plastics in Healthcare

'Conversation starters'





	Manufacturing		Environmental Impact	Engineering		Functional	Supply chain	Economics	Regulatory	End users
	Moulding	Printing		Mechanical testing		Biological testing				Social Opinions
				Tensile strength	Contact angle					
Re-HIPS	Green	Yellow	Grey	Green	Green	Green	Green	Green	White	Green
Bioflex	Green	Red	Grey	Green	Green	Green	Green	Red	White	Green
Gumtec	Green	Green	Grey	Green	Green	Green	Green	Yellow	White	Red
Limex	Green	Green	Grey	Green	Green	Green	Green	Yellow	White	Green
Terralene	Green	Red	Grey	Green	Green	Green	Green	Red	White	Green
Black leather	Green	White	Grey	Green	Green	Green	Red	White	White	Yellow
Grey leather	Green	White	Grey	Green	Green	Green	Red	White	White	Yellow
Flexsea	Green	White	Grey	Yellow	Yellow	Green	Red	Red	White	Green



Introducing sustainable materials in diagnostic testing : Key lessons

Purpose & scope

- Lower environmental footprint of POCTs

Key features

- Comparison of various 'alternative' materials
- Prototypes/ 'Conversations Starters'

Potential Impact

- Lower POCT Sector Carbon Emissions and environmental impact
- Support lobbying manufacturers and policy makers for more sustainable products and sustainable targets

Future work

- Full parametric LCA
- Full manufacturing tests
- Full regulatory approval



Conclusions

- Reducing waste is an environmental priority
- Reducing waste needs to be tackled holistically, by coalition of manufacturers, policy-makers, approved bodies in consultation with end-users
- Collaborations welcomed



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Tools for change: Quantifying and Understanding Diagnostic Plastic Waste

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Photograph: Courtesy Collins Otieno, ASLM

MATERIALS, DESIGN & REAGENTS

Alternative to virgin plastics:

Recycled plastics, natural products: paper, dry cellulose, cardboard, wood pulp

0% toxic gases during waste disposal

Control over plastic additives and reagents

Eco-design

Single plastic solution for greater recyclability, where possible

Upcycled/re-usable components

0% toxic reagents

Novel DNA extraction and non-chemical lysis will enable a new generation of sustainable POCTs

PROCESSES

Local manufacturing

LCA at design stage

End-user involvement

Manufacturer take-back schemes

REGULATIONS

Recycling signage on medical devices

Environmental sustainability requirements in TPPs

PROCUREMENT

Guidance on carbon footprint measurements

Environmental criteria in procurement

ECONOMICS

Cost of material or processes changes

Cost of regulatory approvals for new materials

