

Better Test=Better Flow

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2025 Diarrhoea Stats

Row Labels	Count of Organism Desc
Adenovirus type 40/41	50
Aeromonas sp	55
Astrovirus	79
Campylobacter sp.	674
Clost.difficile Toxin AB	206
Cryptosporidium sp	58
Entamoeba sp	22
Enterovirus	75
ETEC	73
Giardia sp	48
Negative	12159
Norovirus(Norwalk-like virus)	731
Rotavirus	138
Salmonella sp	49
Sapovirus	140
Shigella sp/EIEC	54
STEC	66
Vibrio cholerae	1
Yersinia sp	23
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Grand Total	14701

**Why did the lab
change method?**

Testing strategies – Pre- 2022

TARGETED ENTERIC TESTING

Combination of:

- Culture
- ELISA
- Modular PCR

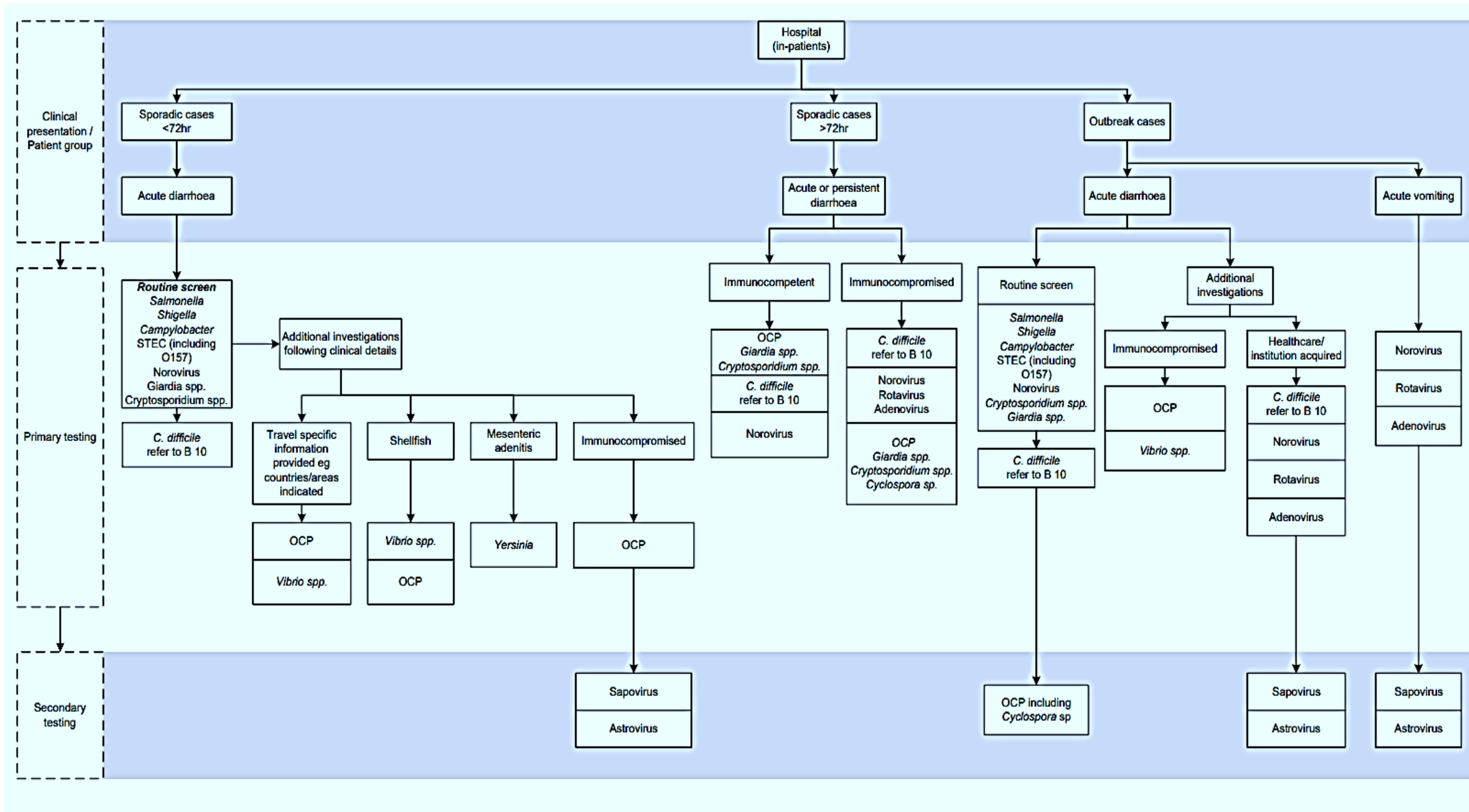
- *previously adopted by BTH*

UK STANDARDS FOR MICROBIOLOGY INVESTIGATIONS – GASTROENTERITIS (S7)

Testing Algorithm: Gastroenteritis in Hospital

S7 | Issue number: 2.2 | Issue date: 08.10.24

Issued by UKHSA



UK STANDARDS FOR MICROBIOLOGY INVESTIGATIONS – GASTROENTERITIS (S7)



>12,000
samples/year

**per 450,000
residents
in catchment area**

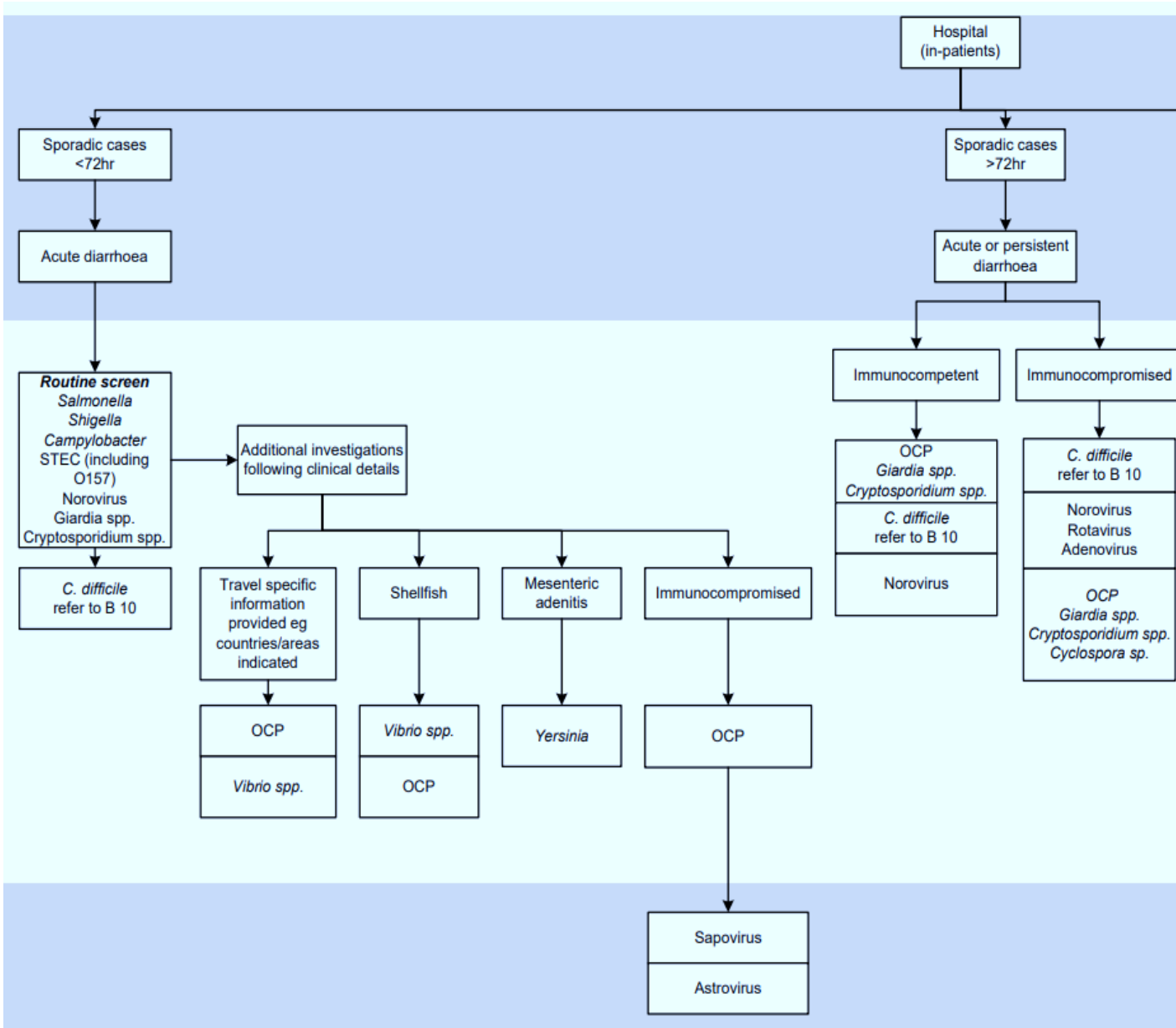
“Stool samples are usually collected and referred for investigation in the following situations:

- *when the clinician requires a microbiological diagnosis
 - when there is **persistent diarrhoea**/malabsorption
 - when there is **blood, mucous** or pus in the stool
 - the patient is **systemically unwell**
 - when there is a history of recent **hospitalisation** or for inpatients as soon as infective diarrhoea is suspected
 - when there is a history of **antibiotic therapy***
- *when a **public health situation** requires sampling to be carried out*
- *when the patient is **immunocompromised***
- *when the patient has **travelled** within 14 days of symptoms onset”*

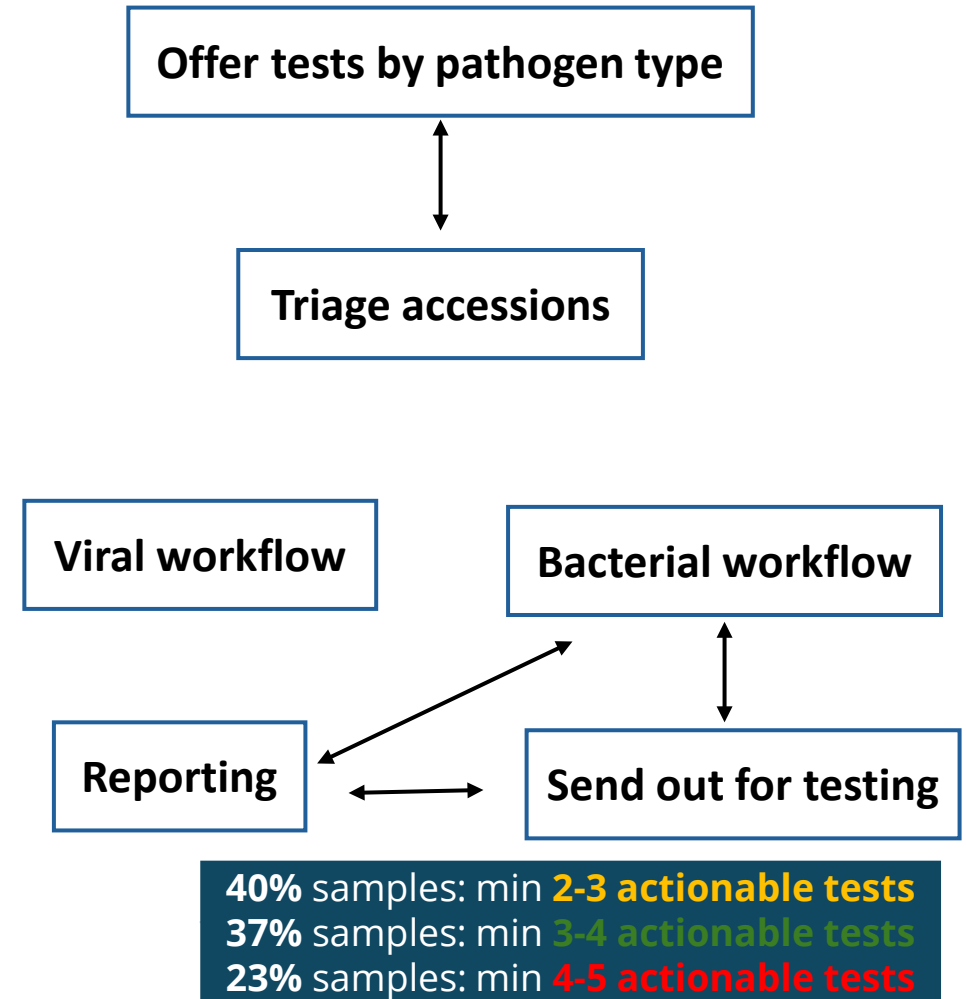
S 7 | Issue number: 2.2 | Issue date: 08.10.24

Issued by the Standards Unit, UK Health Security Agency

UK STANDARDS FOR MICROBIOLOGY INVESTIGATIONS – GASTROENTERITIS (S7)



Conventional testing paradigm



40% samples: min **2-3 tests**
37% samples: min **3-4 tests**
23% samples: min **4-5 tests**

TARGETED ENTERIC TESTING

- Combination of culture, ELISA and Modular PCR - *previously adopted by BTH*
- Modular PCR

CHALLENGE

- ⊗ Complexity of implementing algorithm in laboratory
- ⊗ Reliant on adequate provision of clinical details for supplementary tests
- ⊗ Overall low yield due to low sensitivity and narrow spectrum of pathogen cover
- ⊗ Patient status may change (e.g. community to in-patient)

CONSEQUENCE

- Batching and increased TATs
- Missed or delayed diagnosis
- Low cost-effectiveness
- Duplication and delay in testing

Testing strategies - Post 2022

PAN-ENTERIC PCR TESTING – *implemented by BTH in 2022*

*A single actionable test (single process) for
all samples that tested for the 24 most
relevant pathogens*

Viruses

- > Adenovirus 40/41
- > Astrovirus
- > Enterovirus
- > Norovirus GI
- > Norovirus GII
- > Rotavirus A
- > Sapovirus

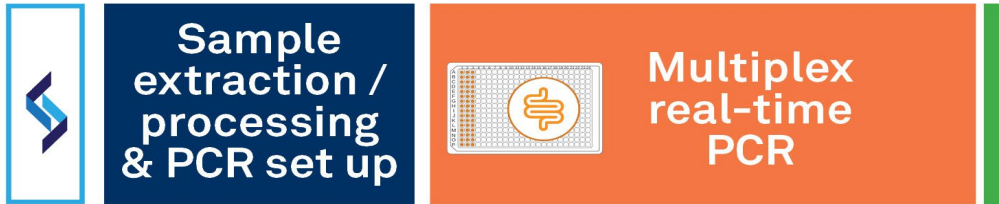
Parasites

- > *Cryptosporidium* spp.
- > *Entamoeba histolytica*
- > *Giardia lamblia/intestinalis*

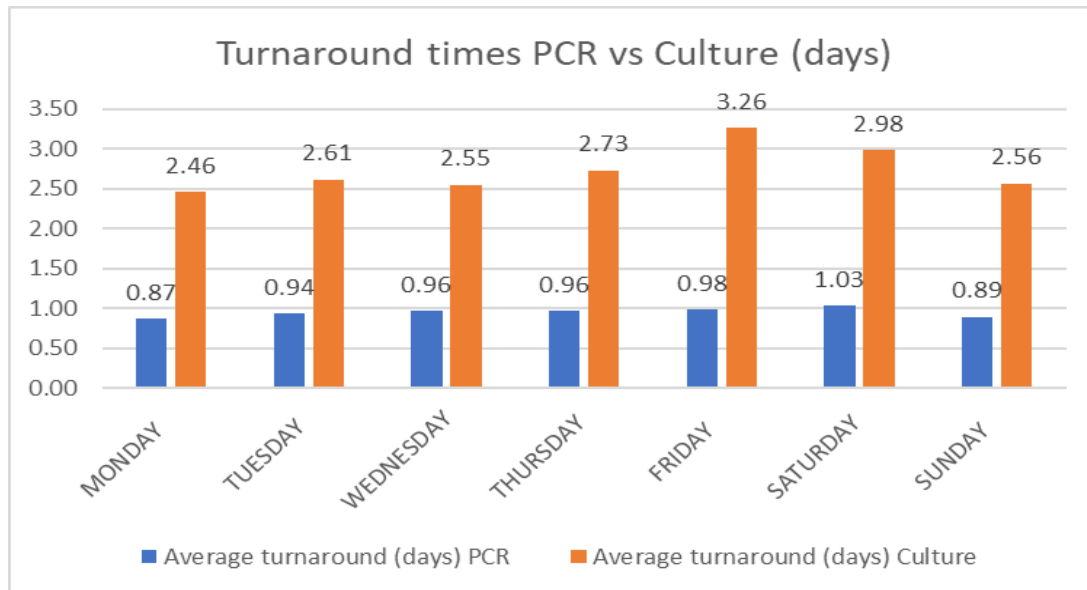
Bacteria

- > *Aeromonas* spp.
- > *Campylobacter* spp.
- > *Clostridioides difficile* (TcdA/B)
- > *E. coli* - eaeA
- > *E. coli* Labile Toxin (LT)
- > *E. coli* Stable Toxin (ST)
- > *E. coli*- stx-1
- > *E. coli*- stx-2
- > *E. coli*- 0157:H7
- > *Salmonella* spp.
- > *Shigella* spp. / EIEC
- > *Vibrio cholerae*
- > *Vibrio* spp.
- > *Yersinia* spp. (excl. *Y. pestis* & *Y. pseudotuberculosis*)

Better Test for the lab and Clinical decision makers



Example: 24 samples / run



Syndromic testing paradigm

Universal sample prep

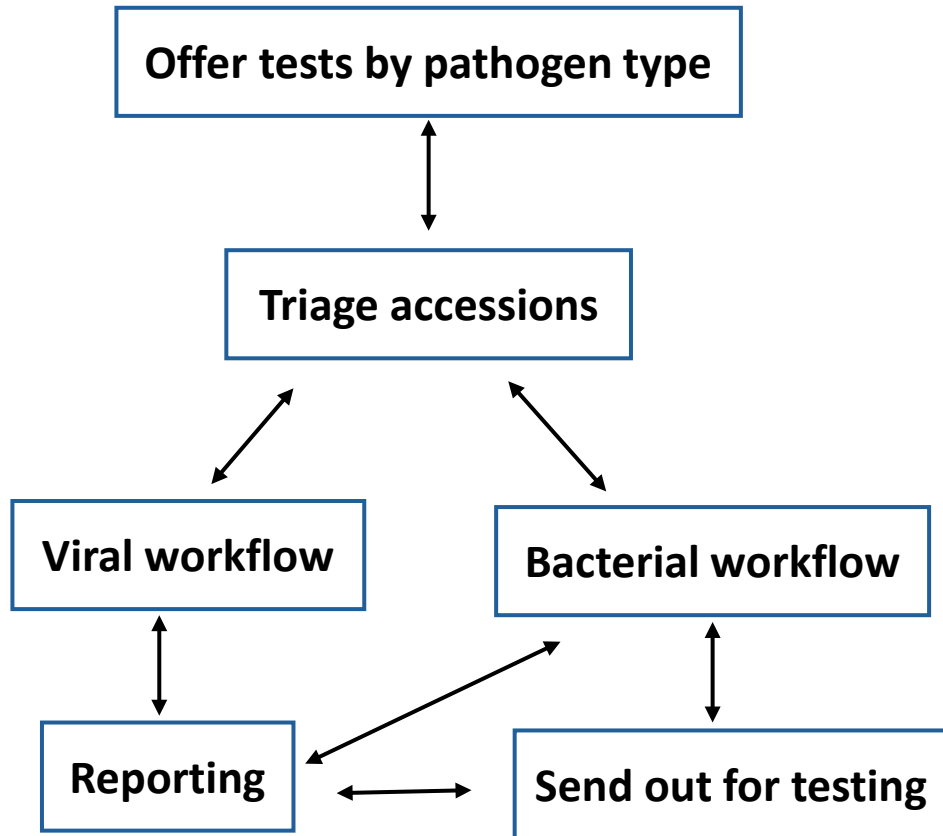
Multiple targets across viral, bacterial and parasitic pathogens (no reflex)

Faster, actionable results across syndromic panels

100% samples: **1 actionable test**

Better Test for the lab and Clinical decision makers

Conventional testing paradigm



40% samples: min **2-3 actionable tests**
37% samples: min **3-4 actionable tests**
23% samples: min **4-5 actionable tests**

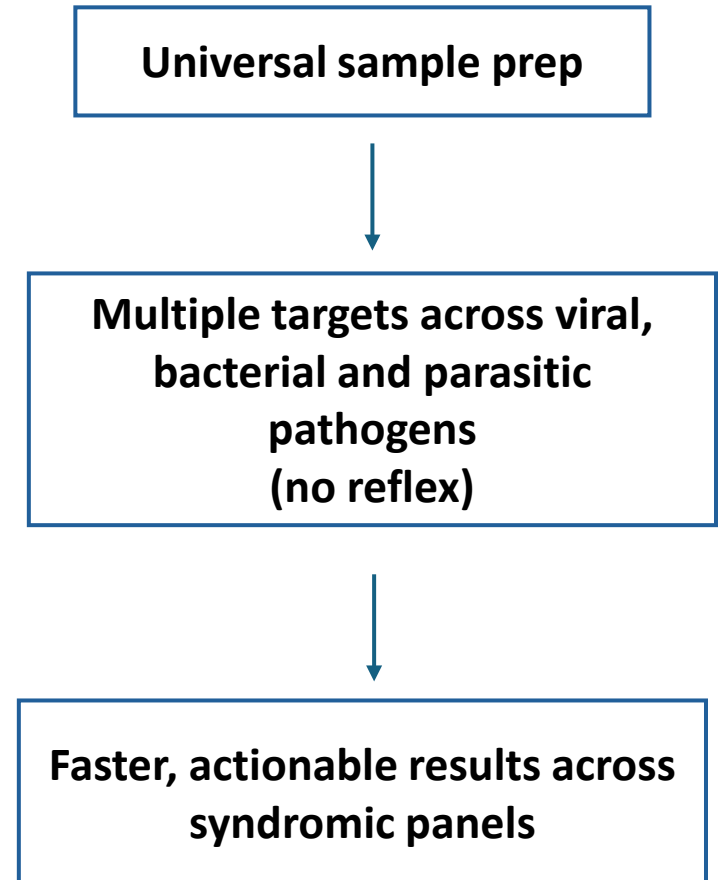
Better Test



Better Flow



Syndromic testing paradigm



100% samples: **1 actionable test**

**BETTER
FLOW**



If patient develops diarrhoea

Proper placement

IPC precaution

Stool sample

Improved Patient Management

Patient Admission Management

- All patients presenting on admission with diarrhoea (Bristol Stool Chart 5-7) stool samples are referred for testing with Pan-Enteric Detection kit.
- At this point symptomatic patients are placed in side-room isolation awaiting result.
- Of the 24 pathogens routinely tested, IPC are interested in the following targets:
 - Toxigenic C-Diff
 - Norovirus

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(blank)	
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C. Difficile Management – Positive Result

All patients presenting on admission with diarrhoea (Bristol Stool Chart 5-7) stool samples are referred for testing with Pan-Enteric Detection kit.

- ✓ 24-48 hours Cdiff ward round with MDT (Microbiologist, IPCN, Antimicrobial Pharmacists,
- ✓ Continue IPC measures until 48 hours free of symptoms

IF 2 or more cases

- ✓ COHORT
- ✓ Enhanced cleaning- high touch areas
- ✓ Daily Enhanced IPC audit by IPNs
 - Hand hygiene
 - Environmental and equipment cleanliness
 - B-B-E
 - I & O documentation
 - Daily check of stool chart
 - Risk assess 1 day prior to completion of treatment

Reinforce IPC Best Practices

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Norovirus Management – Positive Result

All patients presenting on admission with diarrhoea (Bristol Stool Chart 5-7) stool samples are referred for testing with Pan-Enteric Detection kit.

IF 2 or more cases

- ✓ COHORT
- ✓ Enhanced cleaning- high touch areas
- ✓ Daily Enhanced IPC audit by IPNs
 - Hand hygiene
 - Environmental and equipment cleanliness
 - B-B-E
 - I & O documentation
 - Daily check of stool chart
- ✓ Continue IPC measures until 48 hours free of symptoms

Reinforce IPC Best Practices

NEGATIVE
RESULTS

Goodbye!



What's Next



Patient-centred approach

- Clinician further risk assess – Factors that might be the cause of diarrhoea
- Integrate clinical pathway
 - ✓ Refer to Gastro Team
 - ✓ Refer to a Nutritionist
 - ✓ Review Medication
 - ✓ History of travel 14 days prior to symptom onset
- Continue nursing in the SR if available
- Remove from the SR if needed for another patient with infection

Reinforce IPC Best Practices

Summary

Better Test (Multiplex Faecal PCR)



Rapid Turnaround Time



Improved Infection Control



Detection of Co-infections



Reduced Unnecessary Testing and Antibiotic Use

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Better Patient Flow

Thank You

