

**POINT OF CARE TESTING USING FEBRIDX ® TO  
IMPROVE ANTIBIOTIC USE FOR RESPIRATOR  
TRACT INFECTIONS IN PRIMARY CARE:  
  
A MIXED METHODS FEASIBILITY STUDIY PROTOCOL  
(PREFIX)**

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## POINT OF CARE TESTING

- There is a drive to safely reducing unnecessary antibiotic prescriptions for lower respiratory tract infections (LRTIs) in primary care
- Differentiating viral vs bacterial LRTIs can be challenging
- A promising area is 'point-of-care' testing (POCT) using rapid diagnostic tests
- Current devices mostly limited to single biomarkers (e.g. CRP) or specific pathogens (e.g. influenza) and require large desktop analyzers

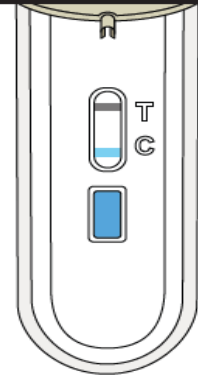
## FebriDx®

- FebriDx ® is a single-use, dual-marker, lateral flow test that tests for viral and bacterial host response using a fingerpick of blood and gives a result in 10 minutes
- It identifies immune responses associated with both bacterial infection (C-Reactive Protein - CRP) and viral infection (Myxovirus resistance protein A - MxA)
- Costs around £12.75 per test

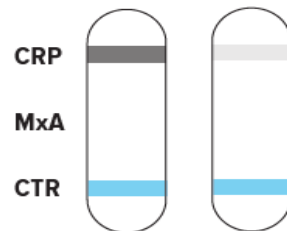


# FebriDx®

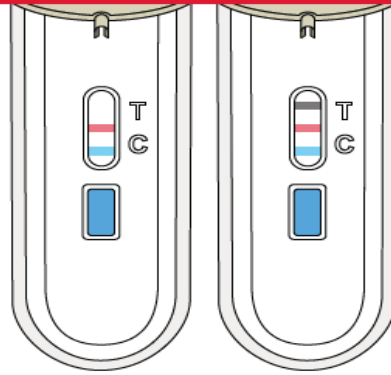
## Bacterial Infection



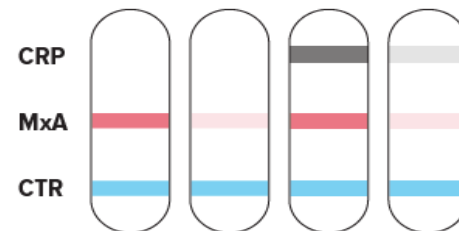
### Black Only Result Line



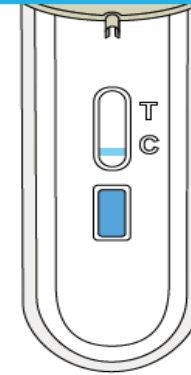
## Viral Infection



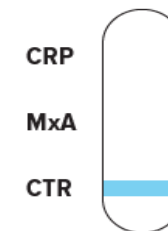
### Any Red Result Line



## Negative Result



### No Result Line



## STUDIES OF FebriDx®

- Nearly all studies based in secondary care
- For bacterial detection, agreement between FebriDx ® and gold-standard PCR was **91.7%** (sensitivity 80%, specificity 93%) and for viral detection agreement was **84%** (sensitivity 87%, specificity 83%).
- Has been rolled out in some centres internationally (including Southampton General Hospital ED)
- To-date only one small study in primary care showing good ease-of-use and cost-effectiveness

## AIMS OF OUR FEASIBILITY STUDY

- 1) Explore the potential clinical utility and facilitators/barriers to using FebriDx ® to safely reduce the use of antibiotics for LRTI in primary care.
- 2) Explore the feasibility of conducting a future trial assessing the clinical impact, safety and cost-effectiveness of FebriDx ® to guide antibiotic use for LRTIs in UK primary care

# STUDY DESIGN

- Mixed-methods, multi-centre, feasibility study of up to 300 patients at GP practices in South England

## STAGE ONE (quantitative data collection)

- **P:** Patients aged >1 year, with symptoms of a LRTI (<28 days), deemed likely to be prescribed antibiotics (immediate or delayed)
- **I:** FebriDx
- **C:** N/A
- **O:** Outcomes include FebriDx results and success rate, time-to-result, antibiotic use, and subsequent re-consultation/antibiotic use and complications

## STAGE TWO (qualitative semi-structured interviews)

- 10-20 patients and 10-20 staff

# INITIAL RESULTS FROM FIRST 38 PATIENTS

## FebriDx results

- 60% were negative (i.e. no detection of CRP or MxA)
  - 10% both CRP and MxA positive
  - 25% positive CRP alone
  - 5% positive MxA alone
- 
- Approximate reduction in antibiotic use of around 50% (not including any re-consultation data)
- 
- Clinicians felt more confident that:
    - Antibiotics were not necessary in 50% of cases
    - Antibiotics were necessary in 42%
    - No difference in 8%



## INITIAL RESULTS – QUALITATIVE

### Three interviews with patients

- Overall positive about the test – found it painless and happy to wait for results
- One patient (had a negative test result) felt the results weren't well explained and left feeling confused and unsatisfied

*I think it's definitely a way forward. I come from the business world and I can see speed and efficiency and time saving and cost saving, so I think it's fantastic. Patient 001-004*

### Early feedback from GP practice staff

- Enthusiastic about the test
- There is definitely a learning curve
- Some confusion about the interpretation of a negative result