

FAECAL COLLECTION KIT

A faecal collection kit used for the collection and transportation of samples for laboratory analysis of biochemical and immunological parameters

This kit contains the materials required for either sample self-collection or collection by a health/social care professional.



For in vitro diagnostic use. sample self-collection









Important: Do not use kit if security seal is broken Keep out of reach of children



Do not re-use

Carefully read the

leaflet before use

enclosed instruction



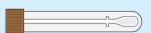
For use by persons aged 18 and over

Sample collection instructions (Steps 1-11)

- Please ensure the kit is within expiry date and read these instructions carefully and completely before attempting to collect the sample.
- If your kit requires online activation, please follow the instructions provided by your healthcare organisation. If you need assistance please contact the healthcare organisation who arranged the test.

Your sample collection kit contents

Please check that the kit contains all of the items outlined below. Do not proceed with sample taking if any items are missing or damaged, contact the healthcare organisation who arranged your test for assistance.



Brown sampling stick and tube for taking your sample



Protective packing case





Return security label

Sample label



Collection kit outer box



Return postal envelope

FAECAL SAMPLE COLLECTION

If not already pre-populated please complete the test request form, provided by your doctor/clinic, separate to this kit. Make sure the sample date is completed.

Please make a note of the unique tracking number displayed under the barcode on the postal tracking label, as shown below. Keep this in a safe place for future reference. You can track delivery of your Royal Mail parcel by entering this number using the following link: http://www.royalmail.com/track-your-item



The test can be done at any time of day or night. Place a bed of toilet paper in the bowl to make it easier to collect your sample. Perform your bowel motion. If possible, try to prevent the sample from touching the water.



Clearly complete the sample collection tube label using a ball point pen with:

- Your First name
- Your Surname
- Your Date of birth
- Sample type: Faecal

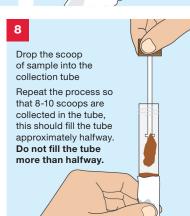
Affix the label to the







Collect your sample by scraping the sampling stick on your bowel motion (stool) until the end of the stick is covered





Place the brown collection tube into the protective packing case and firmly close. Put the case into the outer collection box.



Before you return your sample please do the following: Place your completed request form supplied by your doctor/clinic into the postal envelope Make sure the brown collection tube is labelled and has been placed inside the protective packing case Place the protective packing case into the collection kit outer box Close the collection kit outer box firmly and apply the return security label to seal it Place the collection kit outer box into the return postal envelope Check that you have taken note of your postal tracking number

You are now ready to seal the postal envelope. Please store at room temperature until posted.

IMPORTANT CHECKLIST

Please post your sample to **The Doctors Laboratory** as soon as possible (ideally on same day or within 24 hours of sample collection) from **ANY** Royal Mail post box in the UK. No stamp is required within the UK.

If you need assistance please contact The Doctors Laboratory on **020 7307 7373** or email **samples@tdlpathology.com**.



We would welcome feedback on your experience of using this self-collection kit to help us improve our services. To complete a short online survey please scan the code or visit:

https://forms.office.com/r/EZfeMc4TVs

Warnings and precautions

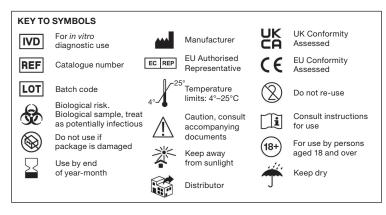
- This kit is designed for use by persons aged 18 and over and upon request of a healthcare professional or healthcare organisation.
- Please consult with a healthcare professional for guidance on sample collection processes for adolescent and younger children.
- The kit should not be used by individuals lacking the physical or mental capacity
 to correctly follow the self-collection instructions. If you have problems or feel
 unwell or lightheaded during the collection process, please pause and if necessary,
 consult with your advising healthcare professional.
- Do not place the sampling spoon directly into your body. The stool sample bottles are designed for collecting faecal samples, no other sample types.
- Do not obtain faecal samples during menstruation (your period) and avoid sample contact with lubricants, bleach and toilet cleaners.
- If your stool sample is excessively watery or hard please discuss this with your healthcare professional to ensure you are collecting a suitable sample for testing.
- The accuracy of your results may be compromised if you do not read and follow the instructions in full.
- Samples arriving at the laboratory which show signs of degradation or general damage or arrive between 3–7 days of sample taking (depending on test requested) may not be tested.

Materials required but not provided

- Test request form. This will be provided by your healthcare professional or healthcare organisation. Please complete the request form with your details as instructed.
- · Clean tissue paper

Laboratory Tests

- The tests and procedures undertaken by The Doctors Laboratory Limited are verified and performed in line with supplier product instructions for use and supported by additional validation data for use with self-collection procedures.
- Test results are provided in line with clinically approved results pathways, agreed between The Doctors Laboratory Limited and the patients designated healthcare professional or healthcare organisation.





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The Doctors Laboratory Limited

Distribution is restricted to regions in which the product is registered by the manufacturer.

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Re	vision/Date	Change summary
1	02/2023	Initial RDi product for UK market
2	09/2023	Steps 8 & 9 updated with clearer instructions
3	01/2024	RDi IVD product CE registration

If any serious incident occurs in relation to the use of this kit, it should be reported to the manufacturer and the competent authority of the Member State in which the user and/or the patient is established.