

SEXUAL HEALTH COLLECTION KIT

A urine sample collection kit used for the collection and transportation of samples for laboratory analysis of sexually transmitted infections by nucleic acid technique (NAT)

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

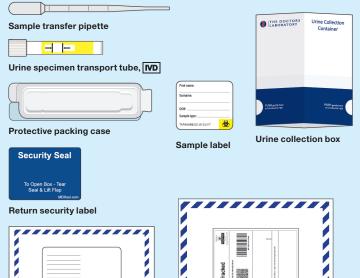


Sample collection instructions (Steps 1–12)

- 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. • You should not have urinated for at least 1 hour before sample taking.

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.

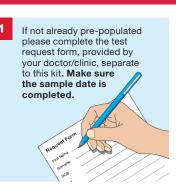


Collection kit outer box

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Return postal envelope

URINE SAMPLE COLLECTION



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

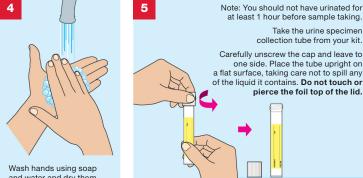
- Your First name
- Your Surname
- Your Date of birth
- · Sample type: Urine

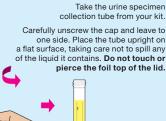
Do not affix it to the tube at this stage.

IMPORTANT SAMPLE TRACKING INFORMATION!

The enclosed postal envelope has a postal tracking label applied. 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







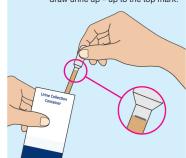
and water and dry them

Squeeze the sides of the urine collection box to expand it. The box has a plastic lining inside to contain the urine. Please take care not to damage

or puncture the lining.

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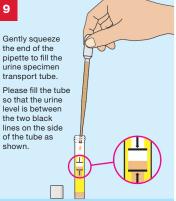
Gently place the end of the pipette into the urine collection box, taking care not to puncture the plastic lining and squeeze the end of the pipette to draw urine up - up to the top mark.

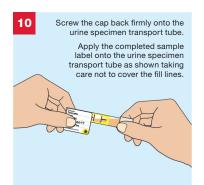


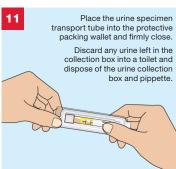


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shown.







12 IMPORTANT CHECKLIST

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 urine specimen transport tube is labelled
- and is 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.

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/SWWruqwKda

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 affix the label to the urine specimen transport tube until after you have collected your sample. You will not be able to see how much urine you have collected if the label covers the tube.
- The Hologic transport buffer contains no substances which at their given concentration, are considered to be hazardous to health. In the case of contact with eyes, skin or mouth please and rinse thoroughly with clean water, if ingested please drink plenty of water and if any symptoms occur please consult a medical professional. In the case of spillages please wipe with absorbent paper and dispose in normal household waste. For further information please refer to safety datasheets which can be found at www.hologic.com or contact MolecularSupport@hologic.com.
- 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 after 30 days (Aptima urine) of sample taking may not be tested.

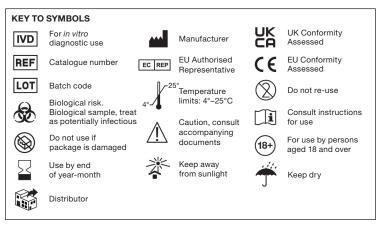
Materials required but not provided

• 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.





REAL Digital International Limited, 2 Queensway, Croydon, Surrey, CRO 4BD, UK Website: www.real-digital.co.uk

EC REP Casus Europe BV, Lange Viestraat 2 B, 3511BK Utrecht, The Netherlands.



The Doctors Laboratory Limited

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

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Revision/Date		Change summary
1	02/2023	Initial RDi product for UK market
2	02/2024	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