**Q fever** is a zoonotic disease caused by the bacteria *Coxiella burnetii*. Human infections occurs after shedding by animals (goats, sheep, cattle, etc.) The disease is observed world-wide The largest outbreak occurred in the Netherlands with an estimated 100,000 human infections.



Datasheet

QDXS-0036

Q-detect<sup>™</sup> is a whole blood interferon-gamma (IFN-y) release assay for Q fever measuring the specific T-cell response to *in-vitro* stimulation with antigen. The assay is a unique test based on the cellular immune response against *Coxiella burnetii*.

Why use Q-detect?

- High sensitivity of > 94% at 5 years after exposure to bacteria
- Supports diagnosis of Q fever and Q fever related Fatigue Syndrome
- Very relevant for screening for exposure to Coxiella
- Specificity of > 92%
- Easy to implement with existing equipment
- Validated in studies with over 1,600 volunteers in the Netherlands

Q-detect requires stimulation with heat-killed *Coxiella* within 12 hours from blood collection, followed by incubation at 37 °C for 24 hours. Read-out is performed by ELISA.

## **Ordering information**

		Product number
Q-detect™	IFN-y release assay for Q fever (36 samples in duplicate) Stimulation plates, ELISA plates, seals, Q- detect <sup>™</sup> antigen Cb2009- 02629, Positive control (PHA), IFN-y standards, block solution, assay buffer, anti-IFN-y coating antibody, anti-IFN-y biotin conjugate, streptavidin-HRP conjugate, TMB ELISA substrate, wash buffer solutions	QDXS-0036

Please contact us for ordering and delivery in your country: <a href="mailto:sales@innatoss.com">sales@innatoss.com</a>

Manufacturer: Innatoss Laboratories B.V. Kloosterstraat 9 - RE21, 5349 AB Oss, the Netherlands

t: +31 412 700507 e: support@innatoss.com i: www.innatoss.com







## **Contents and Storage conditions**

Part	Temperature
96-well stimulation plates	Room temperature
96-well ELISA plates	Room temperature
Seals	Room temperature
Q-detect™ Antigen Cb2009-02629	-20 °C
Positive control (PHA)	-20 °C
Concentrated block solution	-20 °C
Concentrated assay buffer	-20 °C
Anti-IFN-y coating antibody	-20 °C
Anti-IFN-y biotine conjugate	-20 °C
IFN-y standards	-20 °C
Streptavidine-HRP conjugate	-20 °C
TMB ELISA substrate	2 - 8 °C
Wash buffer detergent	Room temperature
Wash buffer concentrate	Room temperature

Not included but required for use		
Biosafety cabinet class I for working with blood		
37 °C incubator (no CO <sub>2</sub> required)		
Calibrated pipettes, variable volume of 1 $\mu$ l to 1000 $\mu$ l and disposable tips with filter		
Calibrated multichannel pipettes, variable volume 50 $\mu$ l to 300 $\mu$ l and disposable tips		
Microtiter plate shaker and washer		
Absorbance reader for microtiter plates with 450 nm filter		
Ultrapure water		
Na <sub>2</sub> CO <sub>3</sub> ( $\geq$ 99.8 %) and NaHCO <sub>3</sub> ( $\geq$ 99.5 %) or ready to use carbonate buffer		
RPMI medium, without phenol red and glutamine (Dutch modification)		
Gentamicin 10 mg/ml		
Sodium pyruvate 100 mM		
Glutamine 200 mM		
Stop solution		

## Q-detect<sup>™</sup> is CE-marked and patented (US9551717/ EP2606361/ AU2011292497)

Q-detect is an *in-vitro* diagnostic aid for the detection of exposure to *C. burnetii*. It is intended to support clinical decisions and should be part of the overall patient assessment. Q-detect should not be use as single reference for clinical decisions and does not distinguish between active and past infection.

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## I a b o r a t o r i e s b v

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