

# MULTIPLES MICRO 5x10 ml

Determination of antibodies associated with salmonella and brucella infections, by coloured suspension in microplate

#### TEST SUMMARY

The Antibodies associated with Salmonella and Brucella infections cause agglutination o.f inactive bacteria present in suspension. The intravital colouring permits an easier reading of agglutination formation.

#### SAMPLES

Serum. Stability 6 days at 4°C.

#### REAGENTS

Coloured intravital inactive bacterial Suspension suspension; conservative stabilizer.

Salmonella

Solution of rabbit antisera that gives Positive control: a clear agglutination with Salmonella

Suspension: conservative stabilizer

Brucella Positive control:

Solution of rabbit antisera that gives a clear agglutination with Brucella Suspension; conservative

stabilizer.

Negative control: Proteic bovine solution that doesn't

react with suspension; conservative

and stabilizer.

#### **REAGENTS PREPARATION**

The bacterial suspension must be resuspended with much care, shaking many times by inversion

The Positive Control be diluted 1:10 with physiologic solution (100 ul + 900 ul).

Stability: the components of this kit will remain stable until the expiration date stated on the label, when stored at 2-8°C protected from direct light. Do not freeze.

# MATERIAL REQUIRED BUT NOT SUPPLIED

Physiologic solution. Automatic micropipette. Normal laboratory equipment.

### **PRECAUTIONS**

Reagent may contain not reactive and conservative components. It is opportune to avoid contacts with the skin and do not swallow.

Perform the test according to the general "Good Laboratory Practice" (GLP) guidelines.

## **SAMPLES PREPARATION**

The serum must be diluted 1:10 with physiologic solution (100 µl of serum with 900 µl of physiologic).

# **PROCEDURE**

In a microplate with "U" wells dilute the serum with physiologic solution as indicated in the following table Using the same pipette (inspiring and discharging many times) mix carefully content of the second well and transfer 100 µl in the following well etc.

Discharge 100 µl from last well (well n°9).

Well	1	2	3	//	9	Susp. Contr.	Contr.	Contr. +
Physiolo.	 100 µl	100 μl 100 μl	100 μl 100 μl	//	100 µl 100 µl	100 μΙ	-	
serum	100 ді	100 ді	from 2	//	from 8	-	-	
Discharge 100 µl from well n°9								
Diluted Positive control	-		-	,		-	-	100 μΙ
Negative control	-	-	-	1		-	100 μΙ	
Bacterial suspens.	100 μΙ	100 μΙ	100 μΙ	//	100 µl	100 μΙ	100 μΙ	100 μΙ
Titre	1:20	1:40	1:80	//	1:5120	-	-	-

Shake the plate by slow rotations for 20-30 sec. Incube at 37  $^{\circ}$ C for 16-18 h or at 22 $^{\circ}$ C for 2 days, to improve bottoms formation it is advisable put the plate in the fridge after the incubation for 2

#### RESULTS INTERPRETATION

A coloured bottom with a clear point shape, on the well bottom, indicates negativity.

An agglutinate that cover all the well bottom indicates a clear positivity, while, a no uniform agglutinate with a bottom in the centre, on the well bottom, indicate a feeble positivity.

The serum titre is given by a high dilution in which there is a feeble positivity.

### **DIAGNOSTIC VALUES**

Titres until 1:40 are considered negative; from 1:80 to 1:160 are suspect, and from 1:320 are positive.

It is a distinctive sign for the infection diagnosis the significant increase of titre between examinated samples after some days.

### NOTE

If the results are incompatible with clinical presentation, they have to be evaluated within a total clinical study

#### CALIBRATION/QUALITY CONTROL

Positive and Negative control sera should be always used to distinguish an eventual background's agglutination of reactive

#### **TEST PERFORMANCE**

#### Sensitivity

In presence of high antibodies titres, phenomenon of prozone can happen, therefore positivity is absent for low dilutions also being present for higher dilutions.

A comparison with an available commercial method gave following results on 50 samples compared, giving a specificity = 100%.

		TYPHI H LTA srl			
		+	-	TOT.	
MPE-	+	17	0	17	
SOL	-	0	33	33	
	тот.	17	33	50	

		TYPHI O LTA srl			
		+	-	TOT.	
OMPE-	+	16	0	16	
SP	-	0	34	34	
•	тот.	16	34	50	

		PARATYPHI A TOTAL LTA srl			
		+	-	TOT.	
COMPETITORS	+ AH	8	0	8	
	+ AO	9	0	9	
	+ AH / AO	4	0	4	
	-	0	29	29	
	тот.	21	29	50	

		PARATYPHI B TOTAL LTA sri			
		+	-	TOT.	
COMPETITORS	+ BH	9	0	9	
	+ BO	12	0	12	
	+ BH / BO	3	0	3	
	-	0	26	26	
	тот.	24	26	50	

		BRUCELLA TOTAL LTA sri			
COMPETITORS		+	-	TOT.	
	+ ABORTUS	12	0	12	
	+ MELITENSIS	7	0	7	
	+ ABORT./MELITEN.	2	0	2	
		0	29	29	
	тот.	21	29	50	

#### WASTE DISPOSAL

Product is intended for professional laboratories. Waste products must be handled as per relevant security cards and local regulations.

### **PACKAGING**

CODE BM01400/P	
Salmonella typhi H	1 x 10 ml
Salmonella typhi O	1 x 10 ml
Salmonella paratyphi A total	1 x 10 ml
Salmonella paratyphi B total	1 x 10 ml
Brucella total	1 x 10 ml
Salmonella Positive Control	1 x 0.5 ml
Brucella Positive Control	1 x 0.5 ml
Negative Control	1 x 0.5 ml
"U" bottom plate with 96 wells	1

Widal F. - Bull. Men. Soc. Med. Hop de Paris - 6; 26 (1986) Bergey's Manual of Determinative Bayteriology 8 Th Ed. Williams and Wilkins Co (1974) Weil E., Felix A .-Wein.Klin.Woch 29; 974 (1916) Gualtney J.B. e coll. -Microagglutination procedures for febrile agglutination tests-Applied microbiology-4; 635-640 Vol.22 (1971) Rose N.R., Friedman H.-Manual of clinical Immunology-American Society for Microbiology, II ed.

### **MANUFACTURER**

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# **SYMBOLS**

Only for IVD use

LOT Lot of manufacturing

REF Code number

Storage temperature interval

Expiration date

Warning, read enclosed documents

Read the directions

Biological risk

Mod. 01.06 (ver. 2.4 - 13/03/2012)

