Determination of antibodies associated to salmonella paraTyphi CO by means of coloured bacterial suspension on slide

TEST SUMMARY
Slide and tube agglutination test for the qualitative and semi-quantitative detection of antibodies associated to Salmonella paraTyphi CO infections. Samples containing the specific antibody cause the agglutination of inactivate bacteria present in suspension. The intravitral coloring allows an easier reading of the formation of the agglutinates.

High levels of agglutinating antibodies are indicative of infection by these microorganisms.

SAMPLES
Fresh clear serum. Stability 7 days at 2-8°C or 3 months at -30°C.

Do not freeze repeatedly.

The samples with presence of fibrin should be centrifuged before testing. Do not use highly hemozized or lipemic samples.

Bring to room temperature before analysis.

REAGENTS
Suspension: Inactivated and intravitral colored bacterial suspension in glycine buffer pH 8.2; preservatives.

MATERIALS REQUIRED BUT NOT SUPPLIED
Saline Solution NaCl 9 g/L; Positive Control serum and Negative Control serum. Slide and stirrer. Automatically microplatte. Mechanical stirrer at 100 r.p.m. Incubator 37°C. Current laboratory instrumentation.

PRECAUTIONS
The reagent may contain non-reactive components and preservatives of various kinds. For precautionary purposes, however, contact with skin and ingestion should be avoided. Use the normal precautions for behavior in the laboratory.

REAGENTS PREPARATION
Reagents are ready to use.

Bacterial suspension has to be carefully resuspended shaking it more times for inversion.

Bring to room temperature before analysis Stability: until expiration date on label stored at 2-8°C. Do not freeze.

PROCEDURE
SLIDE AGGLUTINATION (QUALITATIVE)

Reagents Sample Positive Control Negativ Control
Sample 50 µl 50 µl 50 µl
Positive control -- 50 µl 50 µl
Negative control Suspension 50 µl (1 gtt) 50 µl (1 gtt) 50 µl (1 gtt)

Mix using a disposable stirrer, spread homogeneously over the entire area enclosed by the ring and shake it with a rotary motion or with a mechanical stirrer at 80-100 rpm. for 1 minute.

SLIDE AGGLUTINATION (TITRATION)

Approximate Titre

Titre 1/20 1/40 1/80 1/160 1/320
NaCl 9 g/L 1.9 ml 1 ml 1 ml 1 ml 1 ml
Sample 100 µl 1 ml 1 ml 1 ml 1 ml

2. Prepare 1 tube for Positive Control and 1 tube for Negative Control with 0.1 ml of control + 0.9 ml of NaCl 9 g/L each.

3. Add 50 µl (1 gtt) of suspension to each tube.

4. Mix thoroughly and incubate tube test at 37°C for 24 h.

RESULTS INTERPRETATION
SLIDE AGGLUTINATION
Examine macroscopically the absence or presence of agglutination after 1 minute by comparing the results with the Positive and Negative control.

Agglutination into time established means positivity. Homogeneous suspension with no visible agglutination is negative.

For each positive result it is advisable to confirm the titre with the test-tube titration.

The results obtained with slide titration method are roughly equivalent to those which would occur in tube test with serum dilutions. Respectively: 1/20 – 1/40 – 1/160 – 1/320 – 1/640 – 1/1280.

TUBE AGGLUTINATION
Examine macroscopically the absence or presence of agglutination by comparing the results with the tubes of Positive and Negative control.

Somatic reaction (O) is characterized by coarse, compact agglutination, which tends to be difficult to disperse.

Partial agglutination is a sign of positive reaction.

The title of the serum examined is due to the most higher dilution in which is showed a feeble positivity.

REFERENCE VALUES
For Somatic Antigen (O) Titre ≥ 1/80 indicate a recent infection.

In case of a positive result with a low titre, it is significant for the diagnosis verify the increase of titre between samples taken at a distance of days.

If the titre remains unchanged it may be a previous contact or previous vaccination.

A single positive result has less significance than the demonstration of a rising or falling antibodies titre as evidence of infection.

The level of “normal” agglutinins to these organisms varies in different countries and different communities. It is recommended that each laboratory establish its own reference range.

NOTE
• In some geographical areas with a high prevalence of febrile antibodies, it is recommended to dilute the sample 1:4 with NaCl 9 g/L before to perform the assay.

• As with any diagnostic procedure, if the results are incompatible with the clinical presentation, the physician should evaluate the data obtained using this test by comparing them with other clinical information.

• For in vitro diagnostic use only.

CALIBRATION/QUALITY CONTROL
There is not any International Reference for the sensitivity standardization of these reagents. For this reason, LTA uses an internal control that contains animal serum with antibodies anti-Salmonellas, and titred with commercial reagents of certified quality.

Use of control sera is recommended as reference; the positive control ought to show a partial or complete agglutination, instead the negative control ought to show no agglutination.

Controls should be ever used to distinguish an eventual agglutination of the bottom of reagent.

Controls should be used as described in procedures or even to be treated as samples (dilution, etc.).

The following controls are available on request:

BS00011 3 x 0.5 ml Febrile Positive Control
(0.5 ml Salmonella, 0.5 ml Brucella, 0.5 ml Proteus)

BS00020 1 x 1 ml Febrile Negative Control

TEST PERFORMANCE
Sensibility
The method sensibility decrease at low temperature. Better results will be obtained at higher temperature up to 10°C.

Interference
No interference was observed by the presence of:

- hemoglobin ≤ 1000 mg/dl
- bilirubin ≤ 20 mg/dl
- lipids ≤ 1000 mg/dl
- rheumatic factor ≤ 300 UI/ml

Recent infection, immunodepression or antibiotic treatment can do false negativity.

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

PACKAGING
CODE BS00810 Suspension S. paratyphi CO slide 1 x 5 ml

REFERENCES

MANUFACTURER
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SYMBOLS
IVD Only for IVD use
LOT No of manufacturing
REF Code number
ST Storage temperature interval
EXP Expiration date (year, month)
WARNING Read enclosed documents
Read the directions
BIOL Biological risk
Mod. 01.06 (ver. 3.0 - 02/05/2018)