

REF	90218024
Pack Size	24 Cards

ABO Subgrouping Card with Anti-H

+ 4°C	Manufacturer	IVD In vitro Diagnostic Medical Device	LOT Batch Number / Lot Number	Expiry date	Anti-H
Consult Instructions for use	Date of Manufacture	REF Catalogue Number	This side up	Keep Away from Sunlight	SBIOCAT [™] ABO Subgrouping Card with Anti-H

SUMMARY

On group O red cells, there is no A or B antigen, and the membrane expresses abundant H antigen. The H antigen is a precursor of A and B antigens, A and B persons have less H substance than O persons. The amount of H antigen detected on red cells with the Anti-H Lectin (Ulex europaeus) using classical tube technique is in order of diminishing quantity, O>A.>B>A.B.A.B.A.B.A.B.

Individuals whose red cells and secretions lack H, A, and B antigens and whose plasma/serum contains potent Anti-H, Anti-A, and Anti-B are termed as O_n(Bombay) phenotype.

REAGENTS

 $SBIOCAT^{\text{\tiny{TM}}} \quad ABO \quad Subgrouping \quad card \quad Anti-H \quad contains \quad six \\ microtubes prefilled \quad with a gel in a suitable buffer containing Anti-H \ Lectin.$

STORAGE AND STABILITY

Store SBIOCATTM gel cards in an upright position at 4-25°C. Do not freeze. Avoid exposure of SBIOCATTM gel cards to direct sunlight or any heat source. The shelf life of SBIOCATTM gel cards is as per the expiry date mentioned on the label. Do not use beyond expiry date. Once the aluminium foil is removed from the microtube, it should be used immediately.

ADDITIONAL REAGENTS AND MATERIALS REQUIRED

SBIOCAT™ Diluent -2 LISS for preparation of red cell suspension, Papain solution suitable for serological applications (Refer package insert before use). Gel card centrifuge (85g), Work station, Micropipette capable of delivering 5-50µl of specimen and Bottle top dispenser.

PRINCIPLE

As the SBIOCAT[™] gel card containing red blood cells is centrifuged under specific conditions, the red blood cells possessing the corresponding antigen will agglutinate in presence of the specific antibody and will be trapped in the gel column. The red blood cells, which do not react are not trapped in the gel column and get settled at the bottom of the microtube. The reactions are then read and graded according to their reactivity pattern

SAMPLE COLLECTION

No special preparation of the patient is required prior to sample collection by approved techniques. For optimal results, freshly

collected sample should be used. Serum or plasma sample can be used. Anticoagulants like EDTA, CPD-A and Citrate can be used. Samples should be centrifuged at 1500g for 10 minutes to avoid fibrin residue which may interfere with results.

SAMPLE PREPARATION

Prepare 0.8% red blood cell suspension in SBIOCATTM Diluent-2 LISS as follows:

- Bring the SBIOCAT[™] Diluent- 2 LISS to room temperature before testing.
- Dispense 1.0 ml of SBIOCAT™ Diluent- 2 LISS into a clean test tube.
- Add 10µI of packed red cells to SBIOCAT[™] Diluent-2 LISS collected in test tube and mix gently.
- Red blood cell suspension so obtained should be used for testing.

TEST PROCEDURE

- Label the SBIOCAT™ ABO Subgrouping Card Anti-H with patient's name / identification number. Remove the aluminium foil of required number of microtubes carefully by pulling it backwards.
- Pipette 50 µl of 0.8% patient's red cell suspension to appropriate microtube.
- 3. Add 25 µl of Enzyme (Papain) to the above microtube.
- 4. Allow the card to incubate for 10 minutes at room temperature.
- Centrifuge the SBIOCAT™ gel card for 10 minutes in the gel card centrifuge.
- Retrieve the card from centrifuge, read and record the results.

Note: For applications on SBIOCATTM HEXA ,50 μ l of 0.8-1.0% red cell suspension can be used instead of 10 μ l of 5% red cell suspension.

INTERPRETATION OF RESULTS

Positive reaction: Agglutinated red blood cells forming a clear line on the surface of gel column or agglutinates dispersed in the gel column.

Negative reaction: Non agglutinated red blood cells settle at the bottom of the microtube forming a compact button.

The reaction strength may be recorded as follows:

Strength of reaction	Comments		
4+	Agglutinated red blood cells form a line on the surface of the gel microtube.		
3+	Most agglutinated red blood cells remain in the upper half of the gel microtube.		
2+	Agglutinated red blood cells are observed throughout the length of the gel microtube. A small button of red blood cells may also be visible at the bottom of the gel microtube.		
1+	Most agglutinated red blood cells remain in the lower half of the gel microtube. A button of cells may also be visible at the bottom of the gel microtube.		
±	Most agglutinated red blood cells are in the lower third part of the gel microtube.		
Negative	All the red blood cells pass through and form a compact button at the bottom of the gel microtube.		
Mixed field agglutination	Agglutinated red blood cells form a line on the surface of the gel and nonagglutinated red blood cells form a compact button at the bottom of the gel microtube.		
Н	Hemolysis of red blood cells		

Note: Visual reading of reactions in a card may differ from the reactions read by any automated software through image processing. However this may not change the final result

Reactions for different blood groups with Anti-H Lectin.

	Anti-A	Anti-AB	Anti-A₁	Anti-H
A ₁	++++	++++	++ to ++++	+ to +++
A ₂	+++ to ++++	++++	Neg to +	++ to ++++
A₁B	++++	++++	++ to ++++	+ to +++
A_2B	+++ to ++++	++++	Neg to +	+ to +++
В	Neg	++++	Neg	++ to +++
0	Neg	Neg	Neg	+++ to ++++
O _h	Neg	Neg	Neg	Neg

NOTE

- In vitro diagnostic reagent for laboratory and professional use only. Not for medicinal use.
 The SBIOCAT™ gel cards contains sodium azide
- < 0.1% as preservative. Avoid contact with skin and
- mucosa. On disposal flush with large quantity of water.
 All SBIOCAT™ gel cards should be centrifuged for one complete cycle (10 minutes) in gel card centrifuge before

- Visually inspect the SBIOCAT $^{\text{TM}}$ gel cards before use. SBIOCAT $^{\text{TM}}$ gel cards having bubble(s) entrapped within the gel can be centrifuged for two complete cycles in gel card centrifuge to remove the bubble, if
- bubbles are not removed the card should not be used.

 SBIOCAT™ gel cards that exhibit any signs of drying 6. (i.e. absence or reduced level of reagent buffer above the gel column), decreased volume of gel, cracked gel
- should not be used.

 SBIOCAT™ gel cards with damaged aluminium foil seal should not be used.

 Freezing of SBIOCAT™ gel cards or evaporation of gel
- or reagent buffer due to exposure to heat may lead to erroneous results.
- Fibrin or particulate matter if present in the sample may lead to erroneous results.
- Fibrin if present in the sample may trap red blood cells on the surface of the gel column presenting a pink line. To avoid, samples should be well centrifuged at 1500g for 10 minutes before taking serum or plasma and RBCs should be washed if not collected properly in an anticoagulant
- Use of red blood cells concentration/ volume and reagents other than those described may lead to erroneous results. Follow the instructions carefully.
- Aged or stored red blood cells may exhibit weaker reactivity than freshly collected cells.
- Do not use hemolysed, lipemic or icteric samples.
- Extreme turbidity or discoloration may indicate microbial contamination or denaturation of protein due to thermal damage. Such SBIOCAT™ gel cards should be discarded. Contamination of reagents during usage may cause false
- 15. positive or negative results.
- Red cell aggregation in the red cell suspension may
- interfere the passage.

 Aluminium foil seal of SBIOCAT[™] gel cards should be removed gently and carefully by pulling the foil seal backwards to avoid contamination of reagents from one microtube to another.
- To avoid contamination always use fresh tips before dispensing into each microtube.

REMARKS

- Known positive and negative control should be tested as per Good Laboratory Practices.
- SBIOCAT™ Red Cell Preserving Solution (Cat. No.90262020) can be used as red blood cell preservative solution for preservation of known cells.

BIBLIOGRAPHY

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- HMSO, Guidelines for the Blood Transfusion Services, 2nd Edition, 1993.
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