





# PROCEDURE FOR EQUINE MAJOR CROSSMATCH

Material provided:



10 Gel Tests

1 buffer solution



1 box of Gel Tests



Sample material: Donor packed red blood cells (pRBCs) and Recipient plasma.

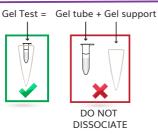
Preferably drawn into EDTA, CPD or ACD. Do not use Heparin.

For reliable results, use of freshly collected blood is indicated (<3 days at 2 - 8 °C).

Further material required: Centrifuge Hettich EBA270; micropipette; test tubes; tips.

### WARNING: Use only the swinging bucket centrifuge Hettich EBA270.

### N°1: Preparation of material provided before use





Allow the buffer solution and Gel Test to reach room temperature before use.

### N°2: Centrifugation of the Gel Test before use

Centrifuge the Gel Test for 2 minutes at 3000 RPM with Hettich EBA270, in order to remove air bubbles or gel drops in the upper part of the Gel Test.



## N°3: Preparation of blood samples for Major XM

# DONOR

Centrifuge blood tube for 5 minutes at 3000 RPM, with Hettich EBA270.
Discard the plasma to collect pRBCs.



Collect blood from blood bag segment.

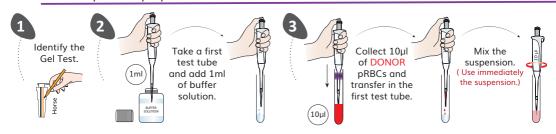


#### RECIPIENT

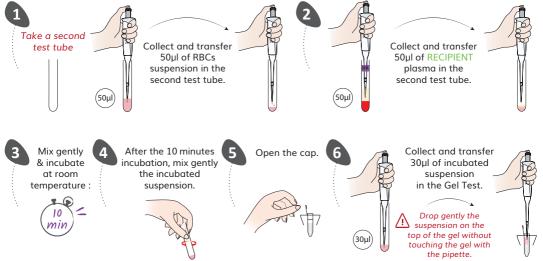
Centrifuge blood tube for 5 minutes at 3000 RPM, with Hettich EBA270, in order to collect plasma.



### N°4: RBCs suspension preparation

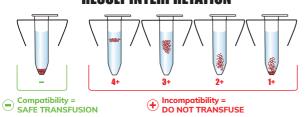


### N°5: Major XM Gel Test procedure









SAFE TRANSFUSION

Troubleshooting Please contact the Scientific Service Laboratory contact@alvedia.com +33(0)478 380 239

Gel Test



## **LIMITATIONS**

- Do not use Gel Test tubes which show signs of drying.
- Gel Test tubes which show air bubbles or gel drops in the upper part of the tubes must be centrifuged before use.
- Strict adherence to the procedures and recommended equipment, especially the Hettich EBA270, is essential for a reliable and validated result.
- A non-specific centrifuge (fixed angle centrifuge) will give you false positive results.
- Fibrin residues in the red blood cell suspension may trap non-agglutinated cells presenting a fine pink line on top of the gel while most of the cells are on the bottom of the microtube after centrifugation.
- Use of suspension solutions other than the provided one may modify the reactions.
- Too diluted or concentrated red blood cell suspensions can cause aberrant results.
- If the blood tube is hemolyzed, wash 3 times in PBS or saline buffer (Nacl 0,9%) to obtain washed pRBCs. Washing RBCs movie procedure: www.alvedia.com



