





PROCEDURE FOR CANINE MAJOR/MINOR CROSSMATCH

Material provided:



1 buffer solution



1 box of Gel Tests



Sample material: Patient's packed red blood cells (pRBCs).

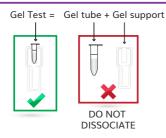
Preferably drawn into EDTA, CPD or ACD. Do not use Heparin.

For reliable results, use of freshly collected blood is indicated (<3 days at 2 - 8 °C). If blood sample > 3 days OR hemolyzed : wash 1 time the blood.

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

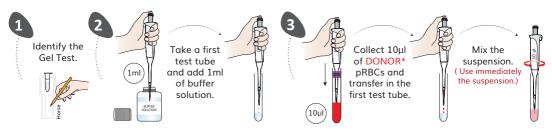
Centrifuge the Gel Test for 2 minutes at 3000 RPM with Hettich EBA270, in order to remove gir bubbles or gel drops in the upper part of the Gel Test.



Plasma

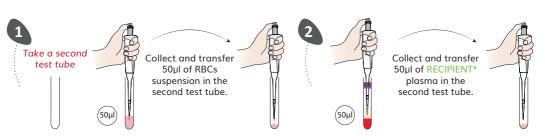
N°3: Preparation of blood samples for MAJOR XM*

RECIPIENT **DONOR** Centrifuge blood tube for 5 minutes OR Collect blood from Centrifuge blood tube for 5 minutes at 3000 RPM, with Hettich EBA270. blood bag segment. at 3000 RPM, with Hettich EBA270, Discard the plasma to collect pRBCs. in order to collect plasma. pRBCs pRBCs



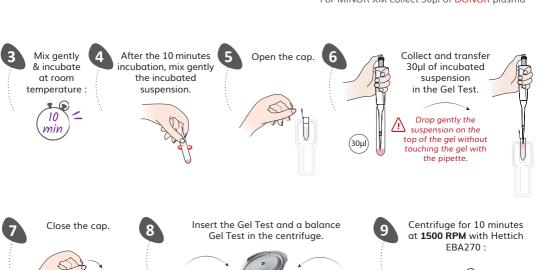
*For MINOR XM collect 10µl of RECIPIENT pRBCs

N°5 : Major XM Gel Test procedure



*For MINOR XM collect 50µl of DONOR plasma

min



Balance Gel Test

Gel Test

RESULT INTERPRETATION

Compatibility = SAFE TRANSFUSION (=)





Incompatibility = DO NOT TRANSFUSE (+)

































LIMITATIONS

- If the blood tube is hemolyzed OR more than 72 hours: wash 1 time in PBS or saline buffer (Nacl 0,9%) to obtain washed pRBCs.
- · 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 tube 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 (fixed angle centrifuge) will give you false positive results.
- Debris, fibrin residues or other artefacts may cause a few unagglutinated cells to trap on top of gel, but these should be interpreted as negative.
- · Use of suspension solutions others than the provided one may modify the reactions.
- Too diluted or concentrated red blood cell suspensions can cause aberrant results.



