

**In case of emergency, or if you find this card,  
please contact the doctor listed below:**

Doctor's Name/Clinic, Center or Hospital Name:

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Telephone contact:

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**DARATUMUMAB**

**IMPORTANT  
MEDICAL INFORMATION  
INSIDE**

**Daratumumab PATIENTS: Provide this card to healthcare providers BEFORE blood transfusion and carry it for 6 months after treatment has ended. For further information please refer to the Patient Information Leaflet**

## **Patient ID Card for DARATUMUMAB**

Name: \_\_\_\_\_

**I am taking the following medication:**

Daratumumab antibody product for the treatment of  
multiple myeloma or AL Amyloidosis

I stopped taking this medication on \_\_\_\_ / \_\_\_\_ / \_\_\_\_  
DD MM YYYY

### **Dear Healthcare Provider,**

Daratumumab is associated with the risk of interference with blood typing.

The Indirect Coombs test (Indirect antiglobulin test [IAT]) may show positive results in patients taking daratumumab, even in the absence of antibodies to minor blood antigens in the patient's serum which may persist for up to 6 months after the last dose. The determination of a patient's ABO and Rh blood type are not impacted.

If an emergency transfusion is required, non-cross-matched, ABO/RHD-compatible RBCs can be given per local blood bank practices.

*For more information, please contact local medical information service at Janssen (placeholder to be completed with country details) or use this reference as a source of additional information:*

<http://onlinelibrary.wiley.com/doi/10.1111/trf.13069/epdf>

*Additional information on interference with blood compatibility testing can be found on (placeholder for local website, if available, to be completed with country details)*

**Before starting daratumumab my blood test results collected on \_\_\_\_ / \_\_\_\_ / \_\_\_\_ were:**  
DD MM YYYY

Blood type: ☐A ☐B ☐AB ☐O ☐Rh+ ☐Rh-

**Indirect Coombs test (antibody screen) was:**

☐Negative ☐Positive for the following antibodies:

Other: \_\_\_\_\_

Contact details of institution where the blood tests were performed: \_\_\_\_\_

**Guidance on Adverse events Reporting-Janssen pharmacovigilance email:**  
E-mail: [JACEG-PV@its.jnj.com](mailto:JACEG-PV@its.jnj.com)