

Antigen

Clone



International Blood Group Reference Laboratory

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BRIC 222

CD44

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Product Code 9406

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Immunoglobulin Class Mouse IgG1, kappa light chain

Antigen Description and Distribution

CD44 (also known as Pgp-1, ECMR III, Hermes antigen, p80¹, H- CAM) is a cell membrane glycoprotein of apparent molecular weight 80 kDa. The full amino acid sequence has been deduced from cDNA. It is heavily glycosylated with both N- and O- glycans. The extracellular part of CD44 comprises an N- terminal disulphide bonded domain and an O- glycosylated domain. CD44 carries the In² and In² blood group antigens². There is a strong association between CD44 and the cytoskeleton. CD44 is thought to be involved in mediating cell:cell adhesion particularly lymphocyte-endothelial cell interactions important for lymphocyte migration from blood to lymph nodes and mucosal associated lymph organs. CD44 is a member of the hyaladherin family of hyaluronan-binding proteins, with a structure similar to selectins, and is the principal cell surface receptor for Hyaluronate³. Antibodies in CD44 may facilitate haemopoietic engraftment⁴. CD44 also functions as an adhesion, hyaluronan, fibronectin, osteopontin and MIP-1β receptor and as a co-stimulatory molecule. CD44 is found on a broad range of haemopoietic cells such as lymphoid cells, myeloid cells, fibroblasts, endothelial cells, epithelial cells, erythroid cells and the nervous system, but not platelets⁵. It is found on brain, heart, liver, thymus, kidney and colon epithelium. CD44 has been mapped to chromosome 11p13. There are approximately 10,000 CD44 molecules per erythrocyte.

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BRIC 222 was made in response to human erythrocytes. BRIC 222 reacts by immunoblotting with a component of Mr 80kDa in nonreduced erythrocyte membranes. Epitope mapping correlates BRIC 222 with the Hermes 2 group of CD44 antibodies which is equivalent to epitope 1⁶, as defined by the Vth Leucocyte workshop⁷. This epitope appears to be associated with the N-terminal region of CD44. It inhibits T cell:erythrocyte rosette formation. It has a functional binding affinity to erythrocytes of 3.8 x 10⁸M⁻¹. BRIC 222 is a direct haemagglutinin. The erythrocyte antigen is pronase, trypsin, chymotrypsin and AET (2-aminoisothiouronium bromide) sensitive. BRIC 222 has been used to investigate the key membrane protein changes during *in vitro* erythropoesis of Protein 4.2 cells⁸.

References

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- 7. Spring *et al.* (1993) Proceedings of the fifth workshop and conference on white cell differentiation antigens, Boston, vol. **2** p 1738-1740.
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