

**International Blood Group  
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<b>Antigen</b>	Wr <sup>b</sup> (ISBT No.211002)
<b>Clone</b>	BRIC 14
<b>Product Code</b>	9413
<b>Immunoglobulin Class</b>	Mouse IgG2a, kappa light chain

**Protein Development  
and Production Unit**

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The Wr<sup>b</sup> antigen is defined as the amino acid sequence at residue 658 (GLU) of the major erythrocyte anion transport protein band 3<sup>1,2,3</sup>. The antigen is stabilised by the association between band 3 and Glycophorin A (GPA, CD 235a). It is found only on erythroid cells. There are approximately 10<sup>6</sup> antigen sites per erythrocyte<sup>4</sup>. The antigen is found on all human erythrocytes except for the very rare Wr(b-) and Glycophorin A deficient types.

**Clone**

BRIC 14 was made in response to erythrocytes. It is unreactive by immunoblotting. BRIC 14 is an indirect haemagglutinin which has a functional binding affinity for erythrocytes of 3.2 x 10<sup>7</sup> M<sup>-1</sup>. Incubation of erythrocytes with BRIC 14 causes a dramatic reduction in erythrocyte deformability and inhibits invasion by malarial parasites (*Plasmodium falciparum* and *P. knowlesi*)<sup>5</sup>. BRIC 14 has been used to investigate the key membrane protein changes during *in vitro* erythropoiesis of Protein 4.2 cells<sup>6</sup>.

**References**

1. Paulitschke *et al.* Blood 1995, **86**: 342-348.
2. Bruce *et al.* Blood 1995, **85**: 541-547.
3. Anstee D.J. (1990) Vox Sang. **58**: 1-20 (Review).
4. Gardner *et al* (1989) Immunol. **68**, 283-289.
5. Pasvol G. *et al* (1989) Blood **74**, 1836-1843.
6. Van den Akker E *et al* (2010). Investigating the key membrane protein changes during *in vitro* erythropoiesis of protein 4.2 (-) cells (mutations Chartres 1 and 2). Haematologica Aug; **95**(8):1278-86.