

Borreliella burgdorferi Antigens

Lyme Disease is caused by different *Borreliella* species of the *B. burgdorferi sensu lato* complex that comprises at least 18 genospecies. Generally, the *Borreliella spp.* was long known as *Borrelia spp.* According to the findings of Adeolu and Gupta (2014) and official nomenclature changes by Oren and Garrity (2015) the species is now named *Borreliella.* This enables differentiation between Lyme Disease and Relapsing Fever spirochaetes, which are still classified *Borrelia spp.* While in Europe the pathogenic genospecies *B. afzelii, B. bavariensis, B. garinii, B. spielmanii, and B. burgdorferi* sensu stricto are present, the latter predominates in the United States (Stanek *et al.* 2012).

Several immunogenic proteins, often associated with the *Borreliella* outer membrane have been identified (Skare *et al.* 1995). These include outer surface proteins A and C

Ordering Information		
40500 40501	Borreliella burgdorferi BmpA	0.1 mg 1.0 mg
40400 40401	Borreliella burgdorferi DbpA	0.1 mg 1.0 mg
40600 40601	Borreliella burgdorferi DbpB	0.1 mg 1.0 mg
41300 41301	Borreliella burgdorferi NapA	0.1 mg 1.0 mg
41200 41201	Borreliella burgdorferi OspA	0.1 mg 1.0 mg
40300 40301	Borreliella burgdorferi OspC	0.1 mg 1.0 mg
42500 42501	Borreliella burgdorferi p28	0.1 mg 1.0 mg
42600 42601	Borreliella burgdorferi p30	0.1 mg 1.0 mg
40200 40201	Borreliella burgdorferi p41	0.1 mg 1.0 mg
41500 41501	Borreliella burgdorferi p45	0.1 mg 1.0 mg
41600 41601	<i>Borreliella burgdorferi</i> p58	0.1 mg 1.0 mg
41700 41701	Borreliella burgdorferi p66	0.1 mg 1.0 mg
40100 40101	<i>Borreliella burgdorferi</i> p100	0.1 mg 1.0 mg
45700 45701	Borreliella burgdorferi VIsE1	0.1 mg 1.0 mg

(OspA, OspC) and neutrophil activating protein A (NapA) that appear to be important for infection and immune evasion (Borchers *et al.* 2015). The protein p28, also known as Oms28, is considered to play an important role in host-pathogen interaction (Cluss *et al.* 2004).



Figure: Immunodot analyses of negative (BD) and positive sera (PS1, PS2) for Borreliella burgdorferi (Bb). The presence of IgG (left) and IgM (right) antibodies was determined spotting triplicates of recombinant DIARECT antigens derived from B. burgdorferi on nitrocellulose membrane.

The antigens p28 and p30 were found to be not detectable in all strains of *B. burgdorferi* (Das *et al.* 1996). Further immunogens are basic membrane protein A (BmpA), flagellal protein p41 (FlaB), p45, p66, p58, and p100. The most sensitive protein for IgG antibody detection in all stages of Lyme disease was found to be VIsE1 (Goettner *et al.* 2005).

DIARECT's recombinant *Borreliella burgdorferi* antigens are produced in either *E. coli* or the baculovirus/insect cell expression system.

References: Adeolu and Gupta (2014) Anton Leeuw Int J G. 105: 1049-1072 Borchers *et al.* (2015) J Autoimmun. 57: 82-115 Cluss *et al.* (2004) Infect Immun. 72: 6279-6286 Das *et al.* (1996) Res Microbiol. 147: 739-751 Goettner *et al.* (2005) J Clinical Microbio. 43: 3602-3609 Oren and Garrity (2015) Int J Syst Evol Microbiol. 65: 1105-1111 Skare *et al.* (1995) J Clin Invest. 96: 2380-2392 Stanek *et al.* (2012) Lancet. 379: 461-473

In some countries the use of certain antigens in diagnostic tests may be protected by patents. DIARECT is not responsible for the determination of these issues and suggests clarification prior to use.

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DIARECT GmbH - Bötzinger Str. 29 B - 79111 Freiburg - Germany

Tel. +49 (0) 761 47979-0 · Fax +49 (0) 761 47979-29 · orders-dia@bbisolutions.com · www.bbisolutions.com