Lipopolysaccharides (LPS)		
Store	Orders: 877-616-CELL (2355) orders@cellsignal.com	
10 mg	Support: 877-678-TECH (8324)	
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Background	Lipopolysaccharide (LPS), also known as endotoxin, is a major glycolipid constituent of the outer cell wall of gram-negative bacteria. LPS molecules typically consist of a strain-specific distal polysaccharide side chain known as the O-antigen, a hydrophilic core oligosaccharide, and a hydrophobic domain referred to as lipid A. Lipid A is covalently bound to the outer bacterial membrane and is responsible for the toxicity of LPS (1-3). LPS is a potent activator of the proinflammatory response in many mammalian cell types, including macrophages, monocytes, and endothelial cells. Investigators have demonstrated that LPS binds to the CD14/TLR4/MD2 receptor complex, which in turn induces inflammatory cytokines including TNF-α, Interleukin-1, and IFN-α, as well as numerous inflammatory proteins such as iNOS, NF-κB, RIG-1, and IRF-3 (4-6).
Description	LPS is supplied as a lyophilized powder and is from <i>E. coli</i> serotype O111:B4. It is purified via phenol extraction.
Solubility	Soluble in PBS and H2O at 5mg/ml and cell culture medium at 1mg/ml.
Storage	Store lyophilized at 4°C. In lyophilized form, the product is stable for 24 months. Once in solution store at -80°C and use within 3 months to prevent loss of potency. <i>Aliquot to avoid multiple freeze/thaw cycles.</i>
Directions for Use	For a 1 mg/ml stock, reconstitute the 10 mg in 10 ml sterile PBS. Working concentrations and length of treatment can vary depending on the desired effect, but it is typically used at 10-1000 ng/ml for 15 min-24 hours. Soluble in PBS and cell culture medium at 5 mg/ml and 1 mg/ml, respectively.
Background References	1. Raetz, C.R. and Whitfield, C. (2002) <i>Annu Rev Biochem</i> 71, 635-700. 2. Rietschel, E.T. et al. (1994) <i>FASEB J</i> 8, 217-25. 3. Maeshima, N. and Fernandez, R.C. (2013) <i>Front Cell Infect Microbiol</i> 3, 3. 4. Pålsson-McDermott, E.M. and O'Neill, L.A. (2004) <i>Immunology</i> 113, 153-62. 5. Wang, J. et al. (2008) <i>J Immunol</i> 180, 8011-9. 6. Zhang, F.X. et al. (1999) <i>J Biol Chem</i> 274, 7611-4.
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