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**Purified Mouse IgG1,  $\kappa$  Isotype Ctrl**

<b>Catalog # / Size:</b>	2607005 / 50 $\mu$ g 2607010 / 500 $\mu$ g
<b>Clone:</b>	MG1-45
<b>Isotype:</b>	Mouse IgG1, $\kappa$
<b>Immunogen:</b>	Trinitrophenol + KLH
<b>Preparation:</b>	The immunoglobulin was purified by affinity chromatography.
<b>Formulation:</b>	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide.
<b>Concentration:</b>	0.5

**Applications:**

<b>Applications:</b>	Other
<b>Recommended Usage:</b>	Each lot of this antibody is quality control tested by immunofluorescent staining with flow cytometric analysis as negative control, and the purity is greater than 95% by SDS-PAGE. Use at concentrations comparable to those of the specific antibody of interest.
<b>Application Notes:</b>	The MG1-45 immunoglobulin is useful as an isotype-matched control (for the relevant formats) for Western blotting, immunoprecipitation, immunohistochemistry, functional assay, immunofluorescence microscopy, immunocytochemistry and immunofluorescent staining (surface or intracellular) for flow cytometric analysis. The LEAF™ purified antibody (Endotoxin <0.1 EU/ $\mu$ g, Azide-Free, 0.2 $\mu$ m filtered) is recommended for functional assays (Cat. No. 401404) as negative control. For <i>in vivo</i> studies or highly sensitive assays, we recommend Ultra-LEAF™ purified antibody (Cat. No. 401408) with a lower endotoxin limit than standard LEAF™ purified antibodies (Endotoxin <0.01 EU/microg).
<b>Application References:</b>	1. Ozeki M, <i>et al.</i> 2008. <i>J. Leukocyte Biol.</i> 84:769. <a href="#">PubMed</a> 2. Kraetzel K, <i>et al.</i> 2008. <i>Eur. Respir. J.</i> 32:563. <a href="#">PubMed</a> 3. de Jong MA, <i>et al.</i> 2008. <i>J Gen Virol.</i> 89:2398. <a href="#">PubMed</a> 4. Romee R, <i>et al.</i> 2012 <i>Blood.</i> 120:4751. <a href="#">PubMed.</a> 5. Wong-Baeza I, <i>et al.</i> 2013. <i>J. Immunol.</i> 190:3216. <a href="#">PubMed.</a>

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<b>Description:</b>	The isotype of MG1-45 immunoglobulin is mouse IgG1, $\kappa$ . It was chosen as an isotype control after screening on a variety of resting, activated, live, and fixed mouse, rat and human tissues.
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