

## Product datasheet

### anti-Yersinia enterocolitica O:9 mouse monoclonal, 8E9, purified

#### Short overview

<b>Cat. No.</b>	8E9-P
<b>Quantity</b>	1 ml
<b>Concentration</b>	50 µg/ml (50 µg)

#### Product description

<b>Host</b>	Mouse
<b>Antibody Type</b>	Monoclonal
<b>Isotype</b>	IgG3
<b>Clone</b>	8E9
<b>Immunogen</b>	Yersinia enterocolitica serogroup O:9 strain
<b>Formulation</b>	PBS, pH 7.4 with 0.09% sodium azide and 0.5% BSA
<b>Conjugate</b>	Unconjugated
<b>Purification</b>	Affinity chromatography
<b>Storage</b>	2-8°C
<b>Intended use</b>	Research use only
<b>Application</b>	Agglutination, ICC/IF
<b>Reactivity</b>	Y. enterocolitica O:9

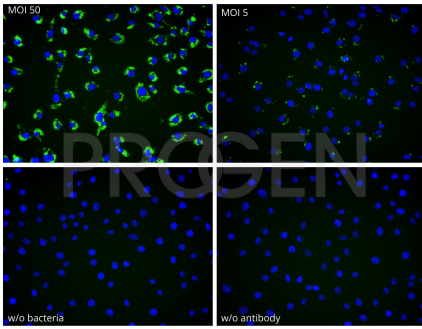
#### Applications

<b>Agglutination</b>	Ready-to-use
<b>Immunocytochemistry (ICC)</b>	1:20

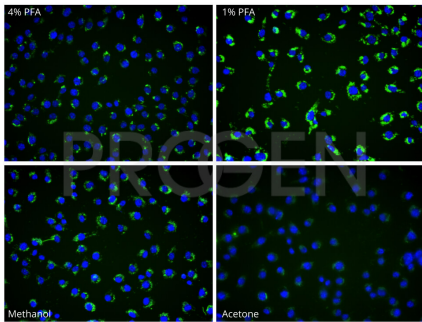
#### Background

8E9 specifically reacts with Yersinia enterocolitica serogroup O:9 strains in both agglutination and immunofluorescence assays. The antibody does not react with Y. enterocolitica O:3 and O:8 serogroups.

#### Product images



Immunofluorescence analysis of *Yersinia enterocolitica* infected EMT6 mouse breast cancer cells with anti-*Yersinia enterocolitica* O:9. Cells were infected with defined amounts of *Yersinia enterocolitica* (multiplicity of infection MOI 50, MOI 5 or without bacteria) for 45 minutes. Fixation was performed using 1% paraformaldehyde for 10 min at RT. Cells were blocked and permeabilized with 1% BSA and 0.3% Triton-X100 in PBS for 1 h at RT. The primary antibody anti-*Yersinia enterocolitica* O:9 mouse monoclonal, 8E9, FITC Conjugate (Cat. No. 8E9-FITC) was diluted in blocking buffer (1:20) and incubated over-night at 4°C. DNA was stained with Hoechst in blue. MOI = multiplicity of infection; w/o = without



Immunofluorescence analysis of *Yersinia enterocolitica* infected EMT6 mouse breast cancer cells with anti-*Yersinia enterocolitica* O:9. Cells were infected with defined amounts of *Yersinia enterocolitica* (multiplicity of infection MOI 50) for 45 minutes. Fixation was performed using either 4% paraformaldehyde (PFA), 1% PFA, 100% methanol or 100% acetone for 10 min at RT. Cells were blocked and permeabilized with 1% BSA and 0.3% Triton-X100 in PBS for 1 h at RT. The primary antibody anti-*Yersinia enterocolitica* O:9 mouse monoclonal, 8E9, FITC Conjugate (Cat. No. 8E9-FITC) was diluted in blocking buffer (1:20) and incubated over-night at 4°C. DNA was stained with Hoechst in blue.