43811

Pyroptosis Antibody Sampler Kit

1 Kit (9 x 20 microliters)



| Orders: | 877-616-CELL (2355) orders@cellsignal.com |
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| Support: | 877-678-TECH (8324) |
| Web: | info@cellsignal.com cellsignal.com |

3 Trask Lane | Danvers | Massachusetts | 01923 | USA

For Research Use Only. Not for Use in Diagnostic Procedures.

| Product Includes | Product # | Quantity | Mol. Wt | Isotype/Source |
|---|-----------|----------|--------------------|----------------|
| Gasdermin D (E8G3F) Rabbit mAb | 97558 | 20 µl | 53, 43, 30, 21 kDa | Rabbit IgG |
| Cleaved Gasdermin D (Asp275) (E7H9G) Rabbit mAb | 36425 | 20 µl | 30 kDa | Rabbit IgG |
| Caspase-1 (D7F10) Rabbit mAb | 3866 | 20 µl | 48, 20 kDa | Rabbit IgG |
| Cleaved Caspase-1 (Asp297) (D57A2) Rabbit mAb | 4199 | 20 µl | 20, 22 kDa | Rabbit IgG |
| IL-1β (D3U3E) Rabbit mAb | 12703 | 20 µl | 17, 31 kDa | Rabbit IgG |
| Cleaved-IL-1β (Asp116) (D3A3Z) Rabbit mAb | 83186 | 20 µl | 17 kDa | Rabbit IgG |
| Caspase-4 Antibody | 4450 | 20 µl | 45 kDa | Rabbit |
| Caspase-5 (D3G4W) Rabbit mAb | 46680 | 20 µl | 50, 44, 35 kDa | Rabbit IgG |
| HMGB1 (D3E5) Rabbit mAb | 6893 | 20 µl | 29 kDa | Rabbit IgG |
| Anti-rabbit IgG, HRP-linked Antibody | 7074 | 100 µl | | Goat |

Please visit cellsignal.com for individual component applications, species cross-reactivity, dilutions, protocols, and additional product information.

| Description | The Pyroptosis Antibody Sampler Kit provides an economical means of detecting proteins that are used as readouts for pyroptosis. The kit includes enough antibodies to perform two western blot experiments with each primary antibody. |
|-----------------------|---|
| Storage | Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. <i>Do not aliquot the antibody.</i> |
| Background | Pyroptosis is a regulated pathway of cell death with morphological features of necrosis, including cell swelling, plasma membrane pore formation, and engagement of an inflammatory response with the release of a number of damage-associated molecular patterns (DAMPs), such as HMGB1 and inflammatory cytokines like IL-1 β and IL-18 (1,2). Pyroptosis is generally induced in cells of the innate immune system, such as monocytes, macrophages, and dendritic cells in the presence of pathogen-associated molecular patterns (PAMPs) expressed on microbial pathogens or by cell-derived DAMPs. It is induced through assembly of inflammasomes triggering proteolytic activation of caspase-1 which then cleaves inflammatory cytokines like IL-1 β and IL-18 to their mature forms (3). A critical feature of pyroptosis is the cleavage of Gasdermin D by caspase-1 and mouse caspase-11 (or human caspase-4/5) (4-6). Upon cleavage, the N-terminal fragment of Gasdermin D oligomerizes to form a pore, allowing secretion of inflammatory DAMPs and cytokines. Canonical inflammasome assembly typically consists of a cytosolic-pattern recognition receptor (PPR; a nucleotide binding domain and leucine-rich repeat [NLR] or AIM2-like family members), an adaptor protein (ASC/TMS1), and pro-caspase-1. Distinct inflammasome complexes can recognize distinct PAMPs and DAMPs to trigger pyroptosis. The best characterized pathway triggered by the NLR, NLRP3, occurs through a two-step process. The first step is a priming signal, NF-kB is activated to induce the expression of a number of inflammasome components including NLRP3, pro-IL-1 β , and pro-IL-18. In the second activation step, caspase-1 is activated and Gasdermin D and cytokines are proteolytically activated. In a non-canonical pathway, caspase-4 and caspase-5 can directly trigger Gasdermin D cleavage in monocytes following LPS stimulation (5,7). |
| Background References | 1. Frank, D. and Vince, J.E. (2019) <i>Cell Death Differ</i> 26, 99-114. 2. Shi, J. et al. (2017) <i>Trends Biochem Sci</i> 42, 245-54. 3. Malik, A. and Kanneganti, T.D. (2017) <i>J Cell Sci</i> 130, 3955-63. 4. He, W.T. et al. (2015) <i>Cell Res</i> 25, 1285-98. 5. Shi, J. et al. (2015) <i>Nature</i> 526, 660-5. 6. Kayagaki, N. et al. (2015) <i>Nature</i> 526, 666-71. 7. Viganò, E. et al. (2015) <i>Nat Commun</i> 6, 8761. |

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