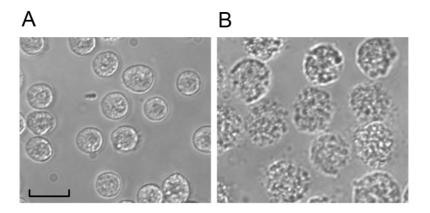
## **Supplementary Material**

## Mast Cell Changes the Phenotype of Microglia via Histamine and ATP

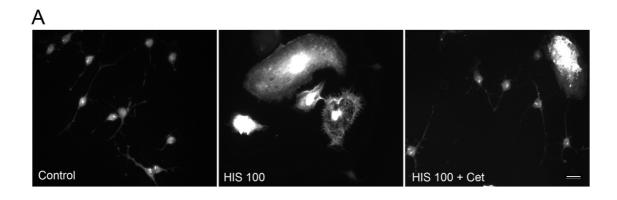
María Pilar Ramírez-Ponce<sup>a</sup> Alejandro Sola-García<sup>a</sup> Santiago Balseiro-Gómez<sup>a,b</sup> María Dolores Maldonado<sup>c</sup> Jorge Acosta<sup>a</sup> Eva Alés<sup>a</sup> Juan Antonio Flores<sup>a,c</sup>

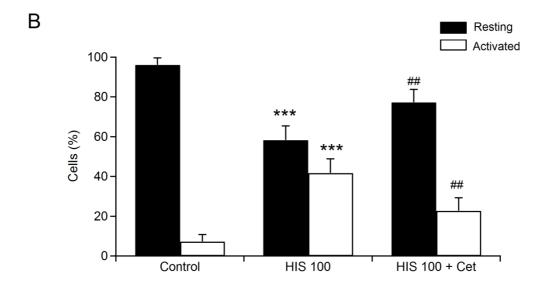
<sup>a</sup>Departamento de Fisiología Médica y Biofísica, Facultad de Medicina, Universidad de Sevilla, Sevilla, Spain, <sup>b</sup>Department of Neuroscience, Yale School of Medicine, New Haven, CT, USA, <sup>c</sup>Departamento de Bioquímica Médica y Biología Molecular e Inmunología, Facultad de Medicina, Universidad de Sevilla, Sevilla, Spain



## Supplementary Figure 1. Representative images of cultured mast cells.

A) after incubation for 1h at 37°C and B) 53°C. Scale bar 10 μm.





**Supplementary Figure 2. Histamine-induced phenotypic change: from resting to activated morphology.** A) Typical morphology of cells observed in a non-tripsinized culture of microglia in control conditions (Control) (where most cells showed a ramified phenotype), after incubation with histamine 100 μM (HIS 100) and incubation with histamine 100 μM in the presence of the H1R antagonist, cetirizine (HIS 100 + Cet) for 48 h. B) Percentage of resting and activated cells in control, HIS 100 and HIS 100 + Cet. Statistically significant from control cells (\*\*\*p<0.001). Statistically significant from histamine-treated cells (##p<0.01), using the Mann-Whitney Rank Sum test.

## **Supplementary Table 1.** Cytosolic calcium signals values in microglia under treatment with HRs antagonists

Treatments	[Ca <sup>2+</sup> ] <sub>I</sub> Peak (F360/F380)	AUC
Histamine (n = 66)	0.35 ± 0.01	3.40 ± 0.23
Histamine + Ceterizine (n = 74)	No response	No response
Histamine + Ranitidine(n = 32)	0.38 ± 0.02	3.68 ± 0.28
Histamine + Carcinine (n = 28)	0.38 ± 0.02	$3.68 \pm 0.32$
Histamine + A943931 (n = 44)	0.34 ± 0.01	$3.65 \pm 0.24$