

Supplementary Information

AHR signaling is induced by infection with coronaviruses

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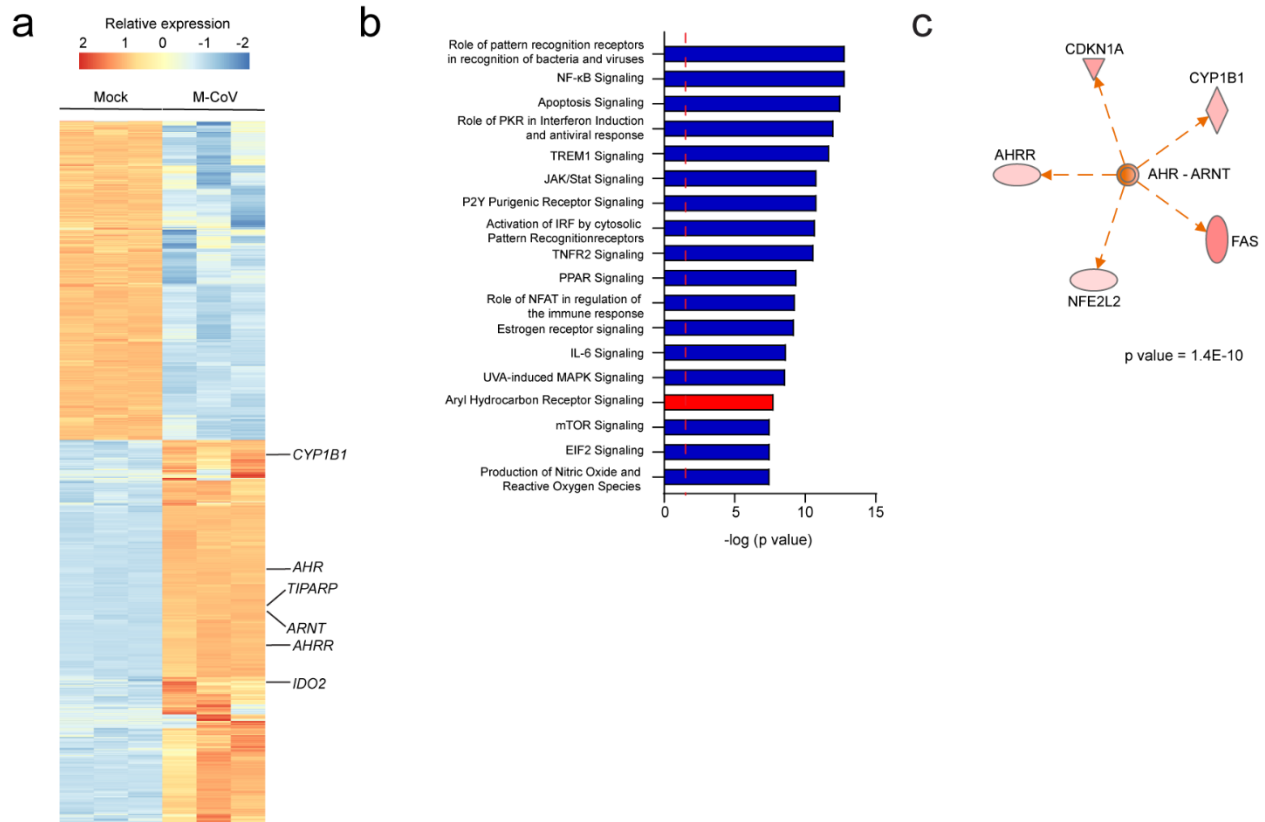
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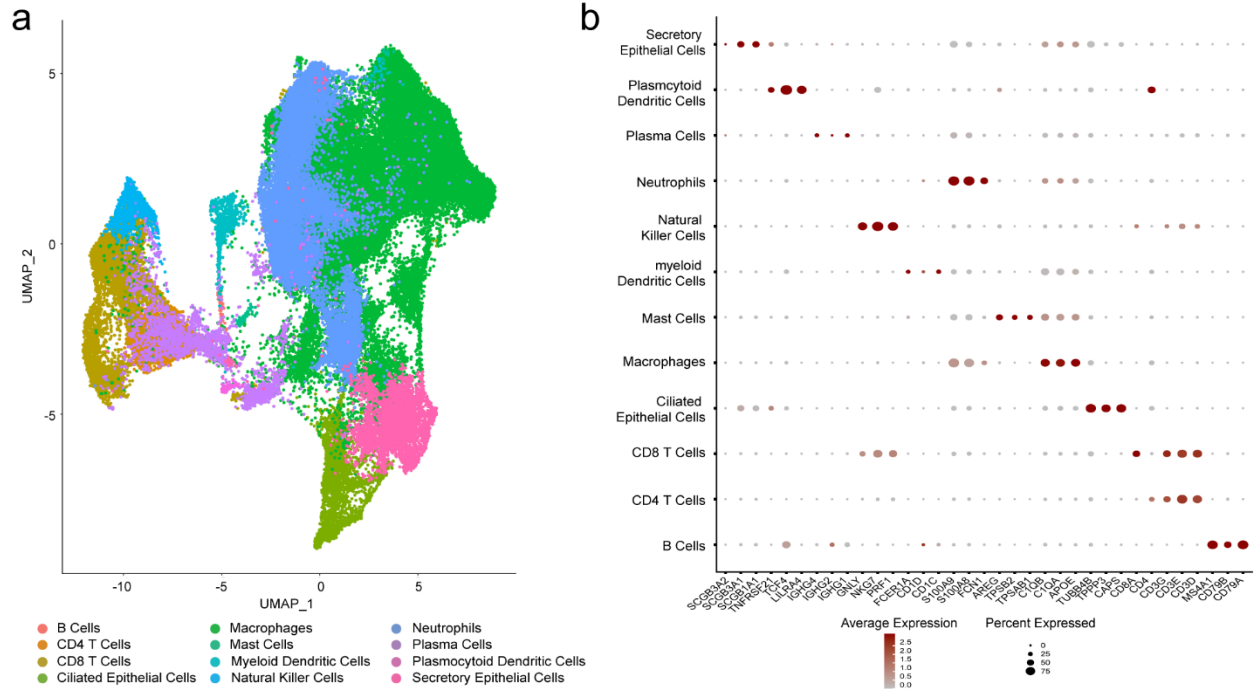
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Supplementary Figure 1. RNA-Seq analyses of M-CoV-infected macrophages. (a) Heatmap showing gene expression levels detected by RNA-seq analysis of mock-infected and M-CoV-infected bone marrow-derived macrophages (n=3 independent experiments per condition) (b) IPA of pathways enriched in M-CoV-infected cells compared to mock-infected cells (n=3 independent experiments per condition). Dashed red line indicates $p=0.05$. p values were determined using a right-tailed Fisher's exact test. (c) IPA Upstream regulator analysis identified AHR as a transcriptional regulator of gene expression in response to M-CoV infection. p value was determined using a right-tailed Fisher's exact test.



Supplementary Figure 2. scRNA-Seq on BALF cells identified 12 major cell types. (a) Uniform Manifold Approximation and Projection (UMAP) plot of BALF cells. (b) Dot plot of marker genes for each major cluster of cells identified. Dot size represents the percentage of cells in each cluster expressing a given gene. Color coding indicates average expression level for each gene.

Supplementary Table 1. Primers used for RT-qPCR analysis.

Gene	Sense	Sequence
<i>IFNB1</i>	Forward	TAGCACTGGCTGGAATGAGA
	Reverse	TCCTTGGCCTTCAGGTAATG
<i>IFNL1</i>	Forward	CTTCCAAGCCCACCACAAC
	Reverse	GGCCTCCAGGACCTTCAGC
<i>IFNL2/3</i>	Forward	GCCACATAGCCCAGTTCAAGTC
	Reverse	GGCATCTTTGGCCCTCTTAAA
<i>IDO</i>	Forward	GGGACACTTTGCTAAAGGCG
	Reverse	GTCTGATAGCTGGGGGTTGC
<i>AHR</i>	Forward	ACATCACCTACGCCAGTCGC
	Reverse	TCTATGCCGCTTGGAAGGAT
<i>PML</i>	Forward	TTCTGCTCCAACCCCAAC
	Reverse	CGCTGATGTCGCACTTGA
<i>RSAD2</i>	Forward	CGTGAAGAGGACATGACGGAAC
	Reverse	CCGCTCTACCAAATCCAGCTTC