

Supplementary Materials

Effects of Mitochondrial Transplantation on Transcriptomics in a Polymicrobial Sepsis Model

Seongmin Kim ^{1,†}, Ji Heon Noh ^{2,†}, Min Ji Lee ³, Ye Jin Park ³, Bo Mi Kim ³, Yun-Seok Kim ³, Sangik Hwang ², Chungoo Park ^{1,*} and Kyuseok Kim ^{3,*}

¹ School of Biological Science and Technology, Chonnam National University, Gwangju 61186, Republic of Korea; sgmn0223@gmail.com

² Department of Biochemistry, Chungnam National University, Daejeon 34134, Republic of Korea; journi@cnu.ac.kr (J.H.N.), hsi9740@naver.com (S.H.)

³ Department of Emergency Medicine, CHA University School of Medicine, Seongnam 13497, Republic of Korea; minji.lee29@gmail.com (M.J.L.); yejin6577@naver.com (Y.J.P.); alfks4050@naver.com (B.M.K.); loupys@naver.com (Y.-S.K.)

* Correspondence: chungoo@jnu.ac.kr (C.P.); dreinstein70@gmail.com or dremkks@cha.ac.kr (K.K.)

† These authors equally contributed to this work.

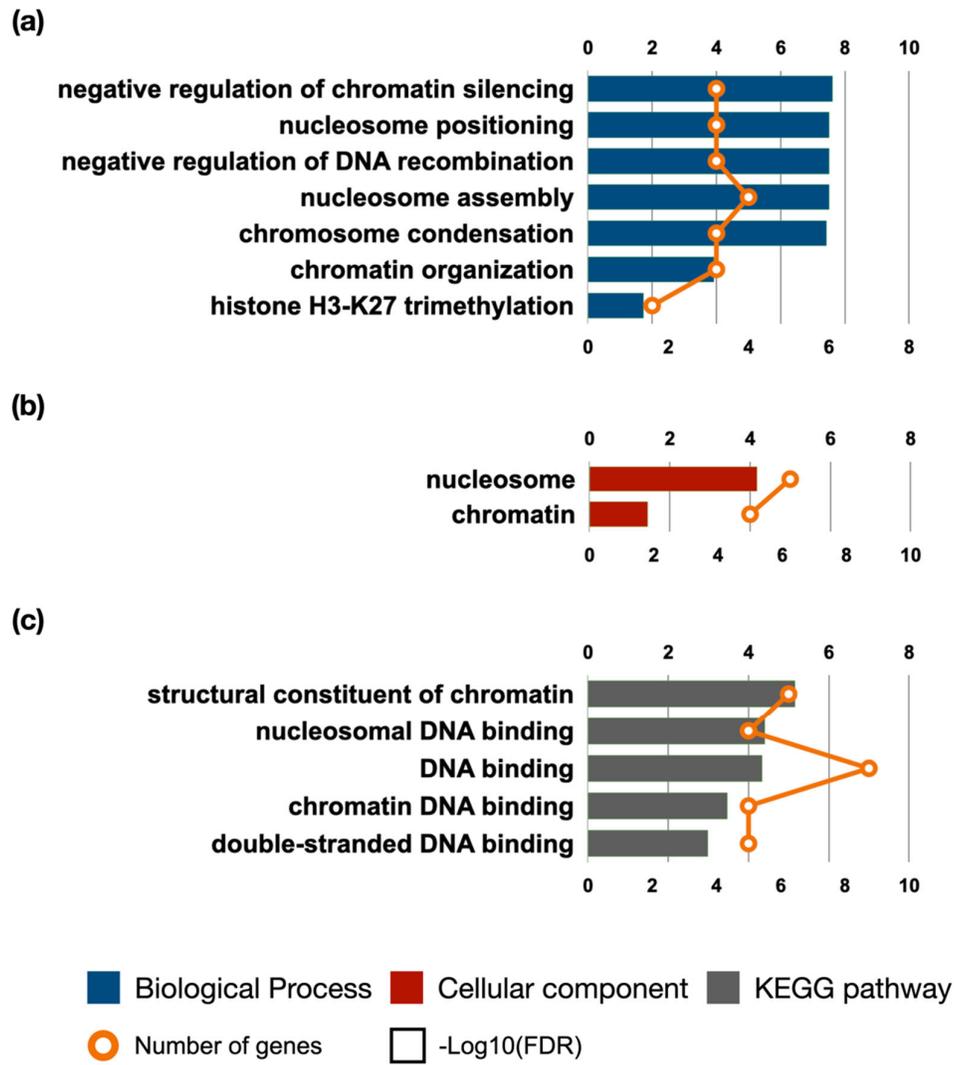


Figure S1. The significantly enriched Gene ontology (GO) biological process (a) and cellular component (b) categories and KEGG (c) pathways of 160 DEGs that were significantly upregulated in SEPSIS versus SEPSIS + MT.

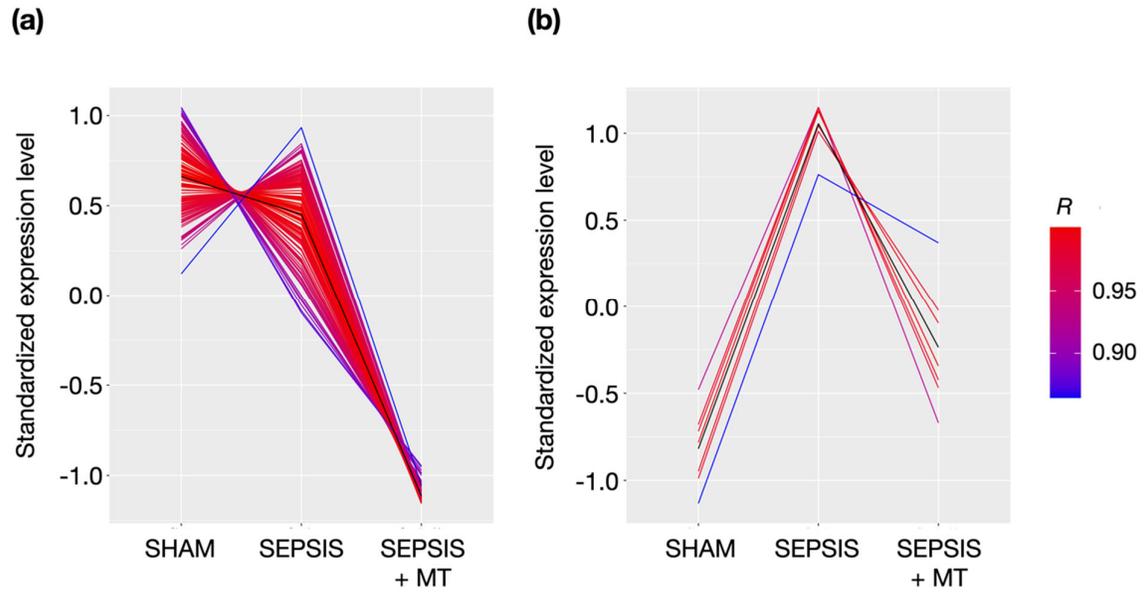


Figure S2. Trajectories of 160 DEGs expression change in three different groups. 153 genes were represented in cluster 1 (a) and the others (seven genes) were identified in cluster 2 (b). Black and other colored lines represent the centroid of clustered gene expression levels and each their own one, respectively. The gradient scale shows the value of Pearson's correlation coefficients (R).

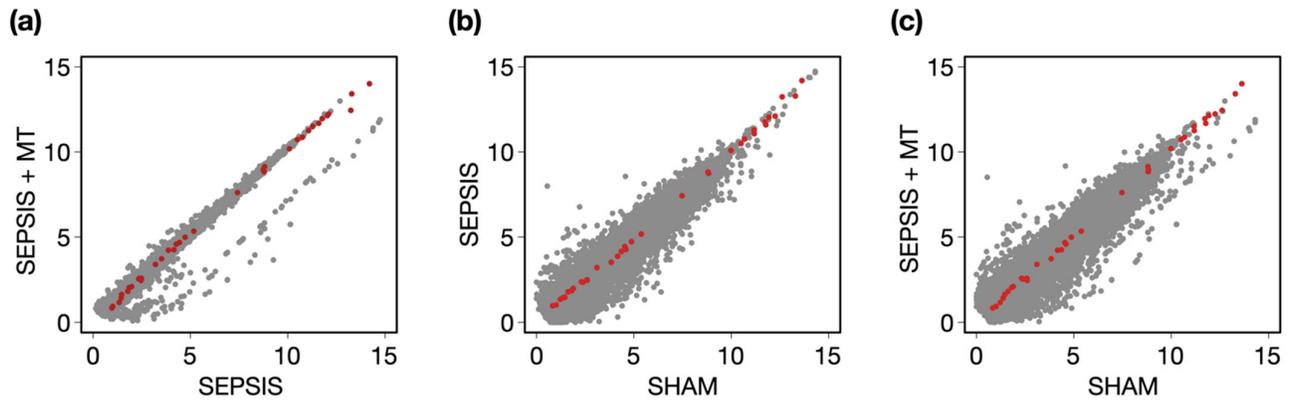


Figure S3. Scatter plots of RNA-seq data between SEPSIS and SEPSIS + MT (a), between SEPSIS and SHAM (b), and between SEPSIS + MT and SHAM (c). Red dot indicates the gene of mitochondrial genome.