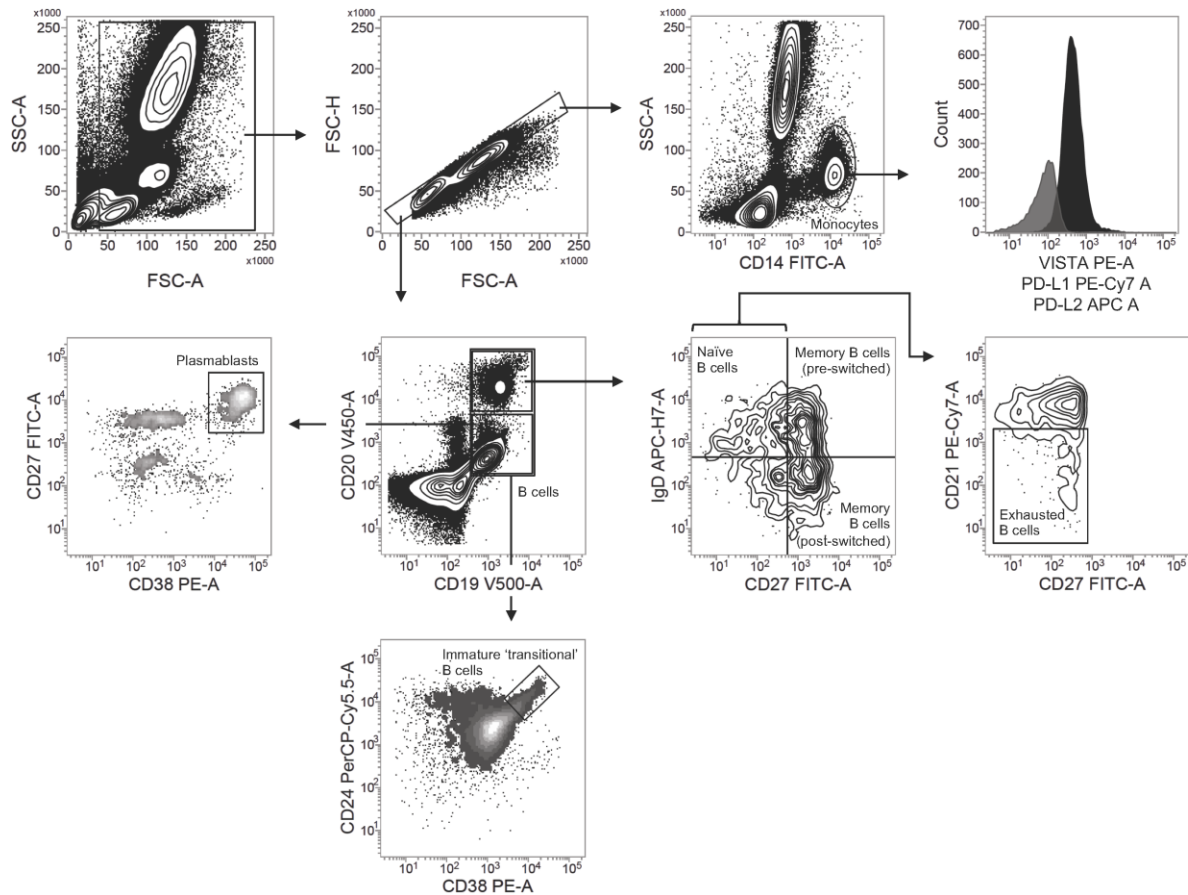
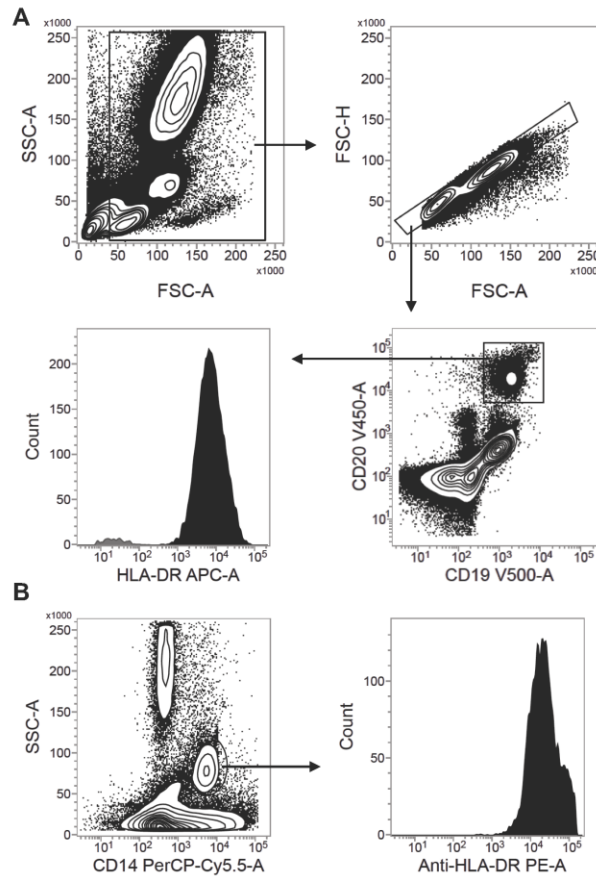


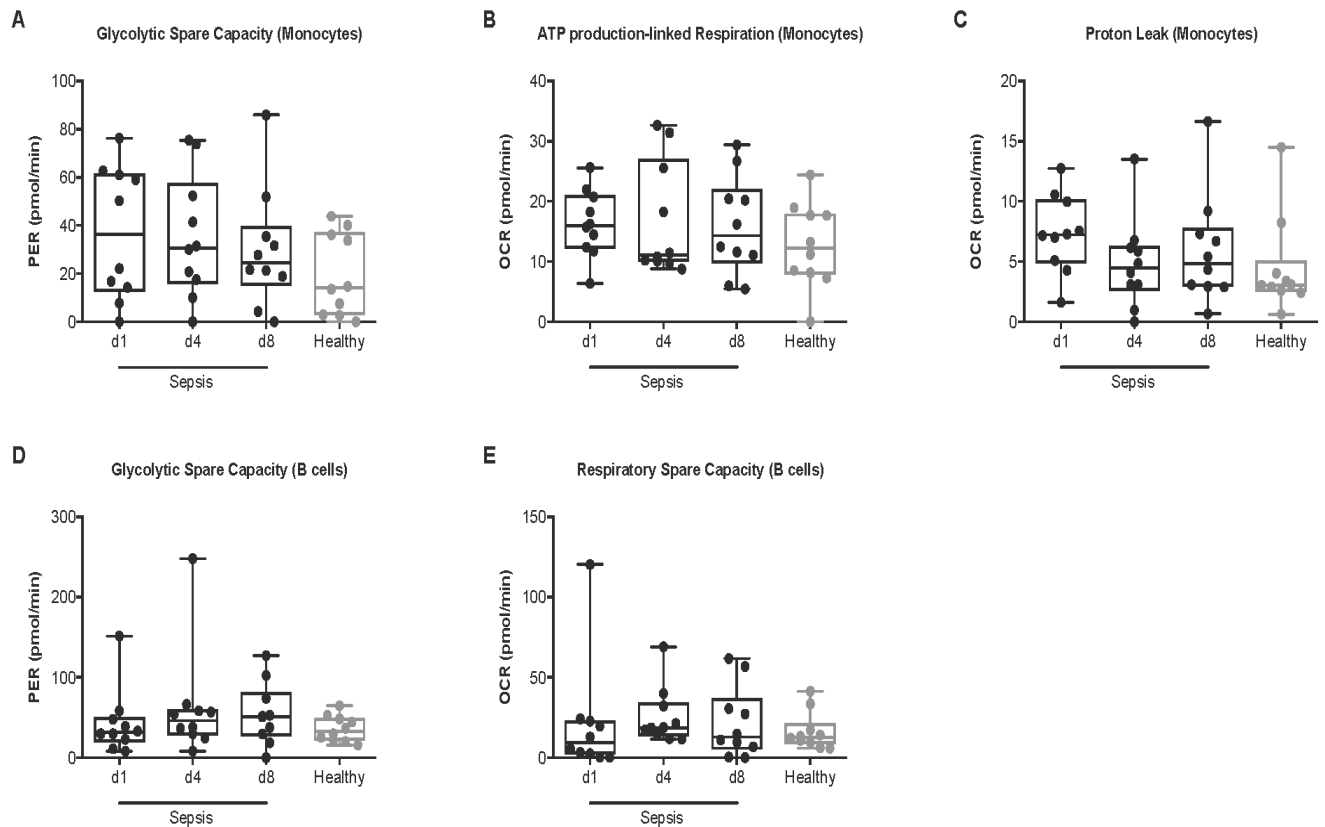
Supplementary Figure 1. Purity control of isolated **(A)** CD19<sup>+</sup> B cells and **(B)** CD14<sup>+</sup> monocytes at d1, d4 and d8. Each data point represents an individual patient/healthy control (n = 10 per group). Horizontal line within the box marks the median, boxes depict the IQR and whiskers indicate the total range.



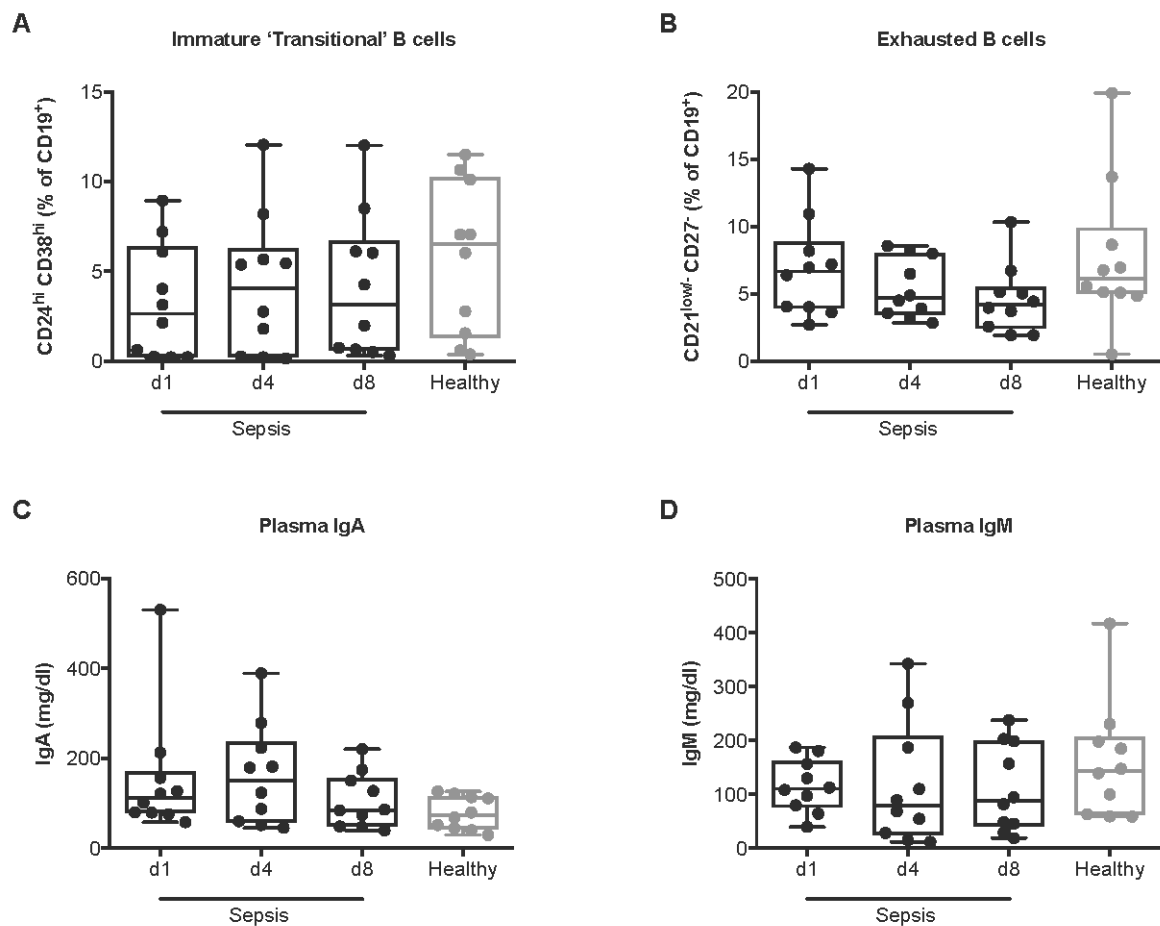
Supplementary Figure 2. **Representative gating strategy for the analysis of monocytes and B cells in whole blood.** Single cells were gated from all cellular events. Monocytes were identified as CD14<sup>+</sup>. PD-L1, PD-L2 and VISTA expression levels were determined via Mean Fluorescence Intensity (MFI) minus background MFI of FMO controls (depicted in grey). B cells were identified as CD19<sup>+</sup> and further characterized as pre-switched memory B cells (CD20<sup>+</sup>, CD27<sup>+</sup>, IgD<sup>+</sup>), post-switched memory B cells (CD20<sup>+</sup>, CD27<sup>+</sup>, IgD<sup>-</sup>), naïve B cells (CD20<sup>+</sup>, CD27<sup>+</sup>, IgD<sup>+</sup>), plasmablasts (CD20<sup>-</sup>, CD38<sup>+</sup>), exhausted B cells (CD20<sup>+</sup>, CD27<sup>-</sup>, CD21<sup>low/-</sup>) and immature 'transitional' B cells (CD24<sup>bright</sup>, CD38<sup>bright</sup>). FMO controls were used for proper gating of CD27 and IgD positive cells.



Supplementary Figure 3. **Representative gating strategy for quantification of HLA-DR.** **(A)** Single cells were gated from all cellular events. B cells were identified as CD19<sup>+</sup> and CD20<sup>+</sup> and HLA-DR was determined by MFI minus background MFI of FMO control (depicted in grey). **(B)** Monocytes were identified as CD14<sup>+</sup>. The average number of HLA-DR molecules per monocyte was quantified using BD Quantibrite PE tubes.



Supplementary Figure 4. **(A)** Glycolytic spare capacity, **(B)** ATP-production linked respiration and **(C)** proton leak of monocytes (CD14<sup>+</sup>) and **(D)** glycolytic spare capacity and **(E)** respiratory spare capacity of B cells (CD19<sup>+</sup>) was calculated at d1, d4 and d8 using PER and OCR. Each data point represents an individual patient/healthy control (n = 10 per group). Horizontal line within the box marks the median, boxes depict the IQR and whiskers indicate the total range. Group comparisons were performed by Mann-Whitney U test.



Supplementary Figure 5. Frequencies of **(A)** immature 'transitional' B cells (CD19<sup>+</sup>, CD24<sup>bright</sup>, CD38<sup>bright</sup>) and **(B)** exhausted B cells (CD19<sup>+</sup>, CD20<sup>+</sup>, CD27<sup>+</sup>, CD21<sup>low/-</sup>). Plasma concentration of **(C)** IgA and **(D)** IgM. Each data point represents an individual patient/healthy control (n = 10 per group). Horizontal line within the box marks the median, boxes depict the IQR and whiskers indicate the total range. Group comparisons were performed by Mann-Whitney U test.