

Supplementary Fig. 3. Oxidative stress, DNA oxidation and DSB levels in the BMs and spleens at the endpoint. (A-F) Expression of mutated *Kras* in hematopoietic cells was induced in double (M-*Kras*<sup>G12D</sup>) and triple (*Nox2*<sup>-/-</sup>M-*Kras*<sup>G12D</sup>) transgenic mice by pIpC injections. Mice were treated with *N*MH (250  $\mu$ g/mouse; red) or NaCl (CON; blue) i.p. thrice weekly for 5 weeks. When moribund, spleen (A-C) and bone marrow (D-F) cells were collected from them M-*Kras*<sup>G12D</sup> and *Nox2*<sup>-/-</sup>M-*Kras*<sup>G12D</sup> mice and analyzed for (A and D) DCFDA expression in CD11b<sup>+</sup> cells (B and E) 8-OHdG expression and (C and F) gamma-H2AX expression in myeloid cells (for spleens: n=8 for *Kras*<sup>WT</sup>, n=12 for M-*Kras*<sup>G12D</sup>, n=8 for *Nox2*<sup>-/-</sup> M-*Kras*<sup>G12</sup>