Supplementary Figure Legends

Supplementary Figure S1. Elimination of Molm-14 from murine c-kit⁺ progenitor cells.

The elimination of Molm-14 cells in murine $c-kit^+$ progenitor cells isolated from the BM of xenografted animals was analyzed by flow cytometry before (**A**) and after (**B**) AC220 treatment. Data is representative of three independent experiments.

Supplementary Figure S2. AML exosomes isolated from primary AML patient samples suppressed clonogenicity of c-kit⁺ progenitor cells in IF-injection experiments

(**A**) AML exosomes were harvested from three different primary AML samples and IF-injected into the femoral cavity alongside contralateral diluent controls of 5 NSG mice. CFU-C assays were performed on murine c-kit⁺ progenitor cells isolated from the BM cells of experimental animals after 48 hrs post IF-injection (one was 96 hrs post IF-injection). * Represents the significance (P<0.05) between the exosome and vehicle treatments determined by pair-t test.

Supplementary Figure S3. Viability of AML cells under hypoxic conditions.

(**A**) The viability of HL-60 and Molm-14 cells cultured in gas-permeable flasks under normoxia or hypoxia did not differ after more than 22 passages of cell culture.

Supplementary Figure S4. NTA profile of leukemia exosomes.

Leukemia exosomes were isolated from the culture media of Molm-14 or HL-60 cells after 48 hrs culture under hypoxia or normoxia condition with 10% VF-FBS/RMPI media. (**A**) NTA measurement of the size and concentration of HL-60 exosomes. (**B**) NTA measurement of the size and concentration of HL-60 exosomes.

Supplementary Figure S5. AML exosomes down-regulated HSC maintenance factors in primary murine BM stroma.

(A) HSC maintenance factors (*Scf, Cxcl12* and *Angpt1*) in primary mouse BM stroma were down-regulated after culture with Molm-14 exosomes for 48 hrs. Data represents two different primary cell isolates with three replicates. S1-S6 represent six individual samples.









