

Detailed analysis of apoptosis and delayed luminescence of human leukemia Jurkat T-cells after proton-irradiation and treatments with oxidant agents and flavonoids

Supplementary Material

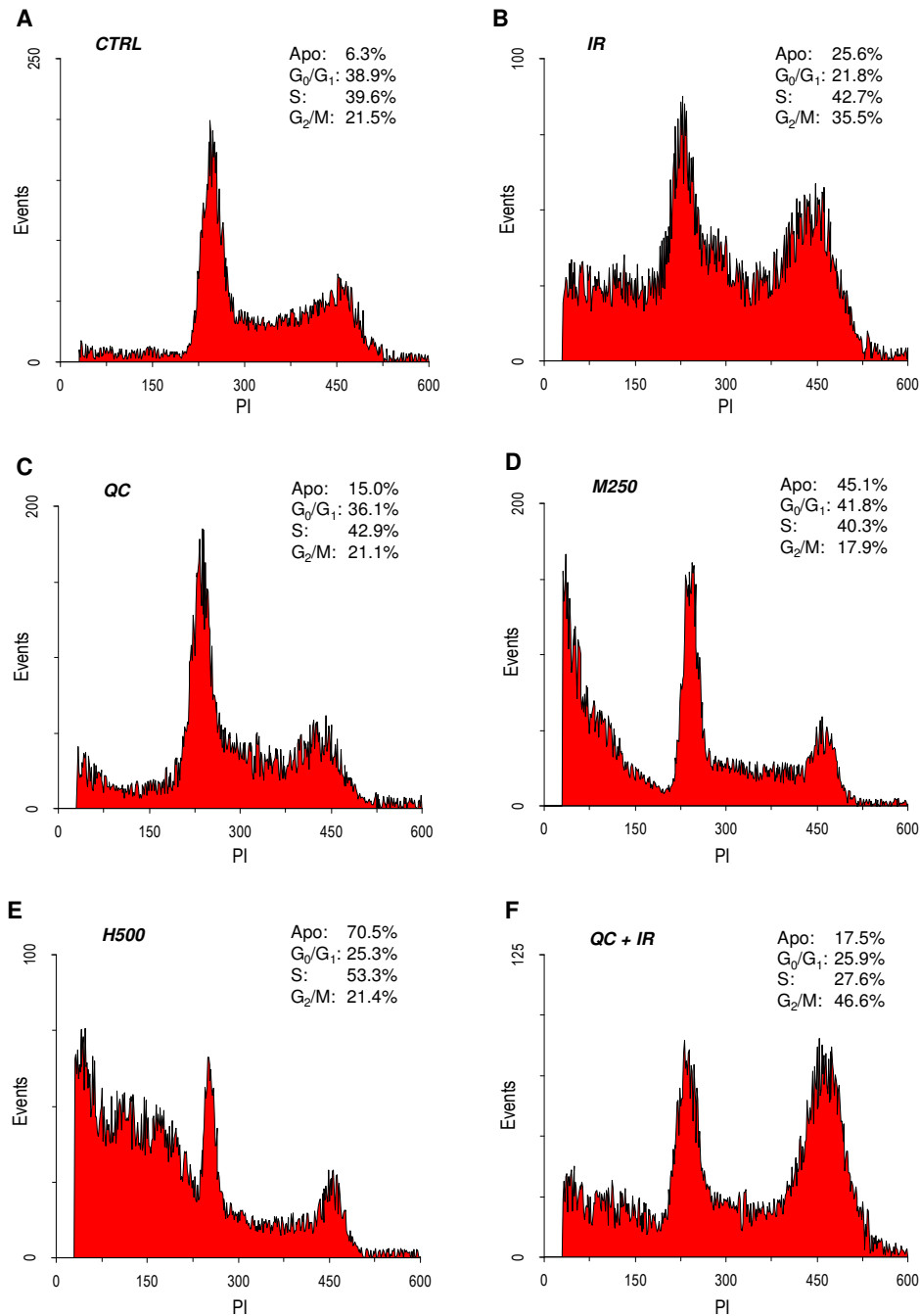


Fig. S1. Representative DNA-content histograms obtained by flow cytometry. After the treatment, cells were collected, fixed and stained with propidium iodide as described in Material and Methods. Percentages of sub-G₀/G₁ (Apo), G₀/G₁, S and G₂/M cells are shown. **(A)** CTRL: Control cells; **(B)** IR: Cells irradiated with 2 Gy of protons; **(C)** QC: Cells treated with 50 μ M QC for 1 h; **(D)** M250: Cells treated with 250 μ M MD for 20 min.; **(E)** H500: Cells treated with 500 μ M H₂O₂ for 20 min.; **(F)** QC + IR: Cells pre-treated with 50 μ M QC for 1 h and then irradiated with 2 Gy of protons. Histograms in B-F were recorded 48 h after the treatment.