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FIG. 1. Expression of p-ERK and p38 in Rps7- and MHC-labeled Rps7-E2 cells. Rps7 cells were treated with 5 mM DMSO, 5 mM MgCl2, 10 mM Na2VO4, 2 mM DTT, 0.1 mM m-FGF, and 10 mM Na2VO4 and were treated with 0.1 M of DMSO for 1 h. The cells were washed twice with PBS and then resuspended in RPMI 1640. The cells were then incubated with 10 mM DMSO for 1 h. After washing, cells were collected, and the cells were collected and analyzed for apoptosis. FIG. 2. Expression of p-ERK and p38 in Rps7- and MHC-labeled Rps7-E2 cells. Rps7 cells were treated with 5 mM DMSO, 5 mM MgCl2, 10 mM Na2VO4, 2 mM DTT, 0.1 mM m-FGF, and 10 mM Na2VO4 and were treated with 0.1 M of DMSO for 1 h. The cells were washed twice with PBS and then resuspended in RPMI 1640. The cells were incubated with 10 mM DMSO for 1 h. The cells were detected by a fluorescence- microscopy. (A) Growth inhibition of Rps7 cells was measured by 3S-ADP-conjugated secondary antibody (Dako IV, Japan) using a Zeiss Axiofermeine (Zeiss, Co., Japan). The cells were treated with 25 mM DMSO for 1 h, and the cells were washed twice with PBS. The cells were then incubated with 10 mM DMSO for 1 h. The cells were collected, and analyzed for apoptosis. (B) Growth inhibition of Rps7 cells was measured by 3S-ADP-conjugated secondary antibody (Dako IV, Japan) using a Zeiss Axiofermeine (Zeiss, Co., Japan). The cells were treated with 25 mM DMSO for 1 h, and the cells were washed twice with PBS. The cells were then incubated with 10 mM DMSO for 1 h. The cells were collected, and analyzed for apoptosis. FIG. 3. Mapping of apoptosis on Rps7- and MHC-labeled Rps7-E2 cells. Cells were treated with 5 mM DMSO, 5 mM MgCl2, 10 mM Na2VO4, 2 mM DTT, 0.1 mM m-FGF, and 10 mM Na2VO4 and were treated with 0.1 M of DMSO for 1 h. The cells were washed twice with PBS, and the cells were incubated with 10 mM DMSO for 1 h. (A) Mapping of apoptosis on Rps7- and MHC-labeled Rps7-E2 cells. The cells were treated with 5 mM DMSO, 5 mM MgCl2, 10 mM Na2VO4, 2 mM DTT, 0.1 mM M-FGF, and 10 mM Na2VO4 and were treated with 0.1 M of DMSO for 1 h. The cells were washed twice with PBS, and the cells were incubated with 10 mM DMSO for 1 h. (B) Mapping of apoptosis on Rps7-E2 cells. The cells were treated with 5 mM DMSO, 5 mM MgCl2, 10 mM Na2VO4, 2 mM DTT, 0.1 mM m-FGF, and 10 mM Na2VO4 and were treated with 0.1 M of DMSO for 1 h. The cells were washed twice with PBS, and the cells were incubated with 10 mM DMSO for 1 h. (C) Mapping of apoptosis on Rps7- and MHC-labeled Rps7-E2 cells. The cells were treated with 5 mM DMSO, 5 mM MgCl2, 10 mM Na2VO4, 2 mM DTT, 0.1 mM m-FGF, and 10 mM Na2VO4 and were treated with 0.1 M of DMSO for 1 h. The cells were washed