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Karen Richardson, Christopher Wilson, Alexander Sutton

Roswell Park Cancer Institute

the apoptotic cells (Figure 5D). In the absence of the DMSO, the cell lysates were analyzed by Western blot analysis, and the levels of apoptotic cells were similar to that of the control. The apoptotic cells also developed in the absence of the DMSO. We also examined the apoptotic cells by Western blot analysis, and the levels of apoptotic cells were similar to that of the control. The intracellular level of cytoplasmic proteins was similar to that of the DMSO. The intracellular level of Akt was similar to that of the DMSO. The cell lysates were analyzed by Western blot analysis. The levels of Akt were also similar to that of the control. The levels of Akt were also similar to that of the DMSO. The expression of intracellular cytoprotective proteins (Figure 5D) was similar to that of the control. The levels of cytoprotective proteins were similar to that of the DMSO. The levels of intracellular cytoprotective proteins were similar to that of the DMSO. The levels of intracellular cytoprotective proteins were also similar to that of the DMSO. In parallel, the DMSO induced apoptosis in the presence of the CL1 (Figure 5E). The levels of intracellular cytoprotective proteins were similar to that of the DMSO. The levels of intracellular cytoprotective proteins were also similar to that of the control. There was no significant difference in the levels of intracellular cytoprotective proteins between the two groups. The levels of intracellular cytoprotective proteins were also similar to that of the control. The levels of intracellular cytoprotective proteins were similar to that of the control. The levels of intracellular cytoprotective proteins were also similar to that of the control. The levels of intracellular cytoprotective proteins were also different from that of the control. The levels of apoptotic cell lysates were different to that of the control. The levels of apoptotic cells were also similar to that of the control. The levels of apoptotic cells were also similar to that of the control. In addition, the levels of glial cell migration were similar to that of the control. The levels of glial cell migration were also similar to that of the control. The levels of apoptotic cell lysates were similar to that of the control. The levels of apoptotic cell lysates were also similar to that of the control. Both the survival and apoptosis were similar to those of the control. The apoptotic cell lysates were more than that of the control. In contrast, the levels of apoptotic cells were similar to that of the control. The levels of apoptotic cells were also similar to those of the control. The levels of apoptotic cells were also similar to those of the control. The levels of apoptotic cells were also similar to those of the control. The apoptotic cell lysates were not related to the control. The levels of apoptotic cells were also similar to that of the control. The levels of apoptotic cells were also similar to that of the control. The levels of apoptotic cells were also similar to those of the control. The levels of intracellular cytoprotective proteins were similar to that of the control. The levels of intracellular cytoprotective proteins were similar to that of the control. The levels of intracellular cytoprotective proteins were also similar to the levels of the control. The levels of intracellular cytoprotective proteins were also similar to that of the control. The levels of intracellular cytoprotective proteins were also similar to that of the control. The levels of intracellular cytoprotec-

tive proteins were also similar to that of the control. The levels of intracellular cytoprotective proteins were similar to that of the control. The levels of intracellular cytoprotective proteins were also similar to that of the control. The levels of PGE2 and PGF1 were similar to that of the control. The levels of PGE2 and PGF1 were also similar to that of the control. The levels of the cytosolic protein PGE2 were similar to that of the control. The levels of PGE2 were also similar to that of the control. The levels of the cytoprotective protein EGF