

**Introduction:** Recent studies confirmed that a moderate amount of wine may have health benefits. Experimental evidence suggested that these beneficial effects could be due to the polyphenol content of red wine.

**Objectives:** In this study, we evaluated the effect of red wine polyphenols (RWP) in J774, a cellular model widely used for studies on oxidative stress and redox capacity of plant extracts and molecules.

**Results:** In the presence of RWP, J774 cells showed an opposite effect on cell viability. Increased cell growth of about 20% was measured at low concentration (4 µg/ml GAE [gallic acid equivalents]), while 50% cytotoxicity was observed at higher concentrations (40 µg/ml GAE). We also detected a rapid (15 min) and dose-dependent increase of superoxide anions (15%–and 180% at 4 and 40 µg/ml GAE, respectively compared to untreated controls). However, while at the lower concentration applied GSH and H<sub>2</sub>O<sub>2</sub> production did not change significantly, at the higher RWP value, GSH and H<sub>2</sub>O<sub>2</sub> contents decreased significantly of 38% and 15%, respectively. To counteract the intracellular ROS production, cells triggered a rapid dose-dependent activation of all the antioxidant enzymes investigated at 4 µg/ml GAE (41% SOD; 18% GSH peroxidase; 24% GSH reductase; 23% G6PD) and 40 µg/ml GAE (150% SOD; 97% GSH peroxidase; 100% GSH reductase and G6PD).

**Conclusions:** Our preliminary results indicate a pro-oxidant activity of RWP, which may promote cell proliferation at low doses and cell death at higher ones.

#### A5

##### VITAMIN D ANALOGUE EB1089 SENSITIZES HUMAN TRANSFORMED CELL LINES TO GAMMA RADIATIONS BY INDUCING LETHAL AUTOPHAGY AND APOPTOSIS

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**Introduction:** The gold standard for unresectable solid tumours and advanced forms of leukemia is represented by chemo-radiation. However, therapeutic options are limited and patients undergo systemic cytotoxicity. Vitamin D insufficiency is a widespread problem and its low serum levels are linked to higher cancer incidence. Previous studies showed a strong correlation between serum concentration of vitamin D and time of first treatment in chronic lymphocytic leukemia (CLL). Other studies demonstrated that vitamin D and its ipocalcemic analogue EB1089 are able to bypass radio-resistance in breast and lung cancer cell lines by activating cytostatic/cytotoxic forms of autophagy. However, only few data are available on the intracellular and molecular effects of vitamin D in osteosarcoma (OS) and CLL-derived cells and on its association with gamma-radiation in sensitizing these cancer types.

**Objectives:** We studied the antiproliferative effects of an active form of vitamin D, the analogue EB1089, in two cell lines U2Os and HG3 derived from a human OS and a CLL, respectively, and its efficacy after treatment with gamma-radiations in terms of cytotoxicity, autophagic and apoptotic effects.

**Results:** EB1089, used at physiological concentration (100 ng/ml), is able to bypass gamma-radiations resistance in U2Os and HG3 cell lines by activating cytotoxic autophagy and apoptosis. The co-treatment resulted highly synergic in terms of combination index (C.I. <1) inducing 85% of cell death at higher doses of radiation after 24h of co-treatment.

**Conclusions:** The results obtained will be discussed at the light of the cytostatic/cytotoxic function of autophagy mediated by vitamin D and involving MAPK/ERK and AMPK pathways in enhancing the therapeutic response to gamma-radiations.

#### A6

##### AUTOPHAGY FLUX MODULATION BY A CAROTENOID-ENRICHED EXTRACT FROM THE PUMPKIN CUCURBITA MOSCHATA ON HUMAN CHRONIC LYMPHOCYTIC LEUKEMIA CELL LINE

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**Introduction:** Chronic lymphocytic leukemia (CLL) is the most frequent form of leukemia in adult population and chemotherapy resistance

occurs in 15–30% of patients with elevated genomic complexity. Apoptosis resistance and induction of a protective form of autophagy are possible explanations of the poor responsiveness of CLL to conventional and novel therapeutic drugs. Given the difficulties to maintain in culture B-CLL lymphocytes, the HG3 cell line represents an interesting preclinical model to study the effects of natural bioactive molecules or extracts derived from food matrix as potential, chemo-sensitizers in CLL.

**Objective:** A previous study demonstrated the induction of “not-protective” autophagy on osteosarcoma and colon adenocarcinoma cell lines after prolonged treatment with a carotenoid-enriched extract (CE) obtained from the pumpkin *Cucurbita moschata*, variety “long of Naples”. To extend and confirm these data, the present communication focuses on the anti-proliferative effect of the same extract in HG3 cell line derived from EBV immortalization of B-CLL cells.

**Results:** CE was obtained from pumpkin by supercritical CO<sub>2</sub> extraction and delivered to HG3 cells in combination with foetal bovine serum. After 96 h, we detected a 40% delay in cell proliferation compared to untreated cells, without signs of cytotoxicity. This delay was due to p27/KIP1 over-expression and modulation of autophagic flux, measured by different autophagy markers (LC3II; p62) and 30% autophagosome intracellular increase.

**Conclusions:** The results obtained will be discussed at the light of the functional cross-talk between the modulation of the autophagy flux by the CE extract and the retard in cell growth observed in HG3 cells, as an opportunity to prolong the asymptomatic phase of CLL before disease occurrence.

#### A7

##### ANTI-INFLAMMATORY EFFECTS OF BLUEBERRY EXTRACT IN MICROGLIAL CELLS

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**Background:** Microglia (MG), the immunocompetent cells of the CNS, respond to brain injury activating and modifying their morphology. Microglia can exist broadly between two different activation states, namely the classical (M1) and the alternative activated (M2) phenotype. The first one is characterized by the production of pro-inflammatory cytokines, in contrast, the latter is characterized by the production of anti-inflammatory cytokines (Kettenmann et al., *Neuron*. 2013; 77:10–18). Blueberry is involved in the control of the redox state of the cell, cooperating with antioxidant mechanisms, whereas its anti-inflammatory activity is still poorly understood (Businaro et al., *Curr. Alzheimer Res.* 2018; 15: 363–380). The aim of the present study is to determine the effect of blueberry extract in resting form or lipopolysaccharide (LPS)-stimulated BV-2 murine MG cells.

**Methods:** The hydroalcoholic extract obtained from fresh blueberries was analyzed by UHPLC/MS. The cellular viability was evaluated by MTT test and Trypan blue assay. Cellular migration was determined by Boyden chamber and Scratch assay. Cytokines mRNA levels were determined by qPCR. Actin cytoskeletal organization and M1/M2 marker expression were analyzed by immunofluorescence.

**Results:** Isomers of the chlorogenic acid, a powerful antioxidant, were detected in the blueberry extract, which, added to the cultures, had no cytotoxic effect, but induced increased cell viability and reduced LPS-driven migration. mRNA expression of pro-inflammatory cytokines IL-1β, IL-6 and TNF-α and that of iNOS (M1 marker) was decreased, whereas Arg-1 expression (M2 marker) was increased.

**Conclusion:** Our results suggest that blueberry may promote MG polarization towards the M2 phenotype, and therefore may be used as a nutraceutical in the treatment of neuroinflammatory diseases.

#### A8

##### INTESTINAL EPITHELIUM RESPONSES TO TITANIUM DIOXIDE NANOPARTICLES

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