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**FOOD BIOACTIVES FIGHTING RENAL CANCER PROGRESSION**

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Many studies suggest a protective role of certain dietary patterns (e.g., consumption of cruciferous vegetables) against the risk of renal cancer. However, the current body of evidence is yet insufficient to establish a link between diet and the risk of renal cancer development and progression. Our work aims at contributing to fill this gap by dissecting the effects of two structurally distinct redox-active food components on kidney cell characteristics related to renal cancer progression. A special emphasis was given to cell motility due to its critical importance for the development of metastases. Thymoquinone is a monoterpene isolated from the oil of *Nigella sativa* seeds, which is widely used as a spice as well as in traditional medicine. We have shown that thymoquinone reduced the viability and promoted apoptosis of 786-O human renal cancer cells. At non-cytotoxic/genotoxic concentrations, thymoquinone significantly decreased the collective migration and the invasiveness potential of these cells. Erucin is an isothiocyanate that can be generated by *in vivo* reduction of sulforaphane or by enzymatic hydrolysis of glucoerucin. Contrarily to sulforaphane, limited studies have addressed the anticancer properties of erucin. Erucin induced a concentration-dependent decrease of cell viability, more pronounced in 786-O cancer cells than in the "normal-like" Vero-E6 cells. The exposure of cells to this bioactive led to an increase of the G2/M population. Collective cell migration, chemotaxis and chemoinvasion abilities, as well as cell adhesion, were impaired in erucin-treated cells. Additionally, erucin induced concentration-dependent changes on cell morphology and impaired tubulin polymerization. Overall, our results suggest that thymoquinone and erucin may have a beneficial impact in reducing renal cancer cells migration, contributing to explore the mechanisms of possible dietary approaches for chemoprevention.

*Keywords: Renal cancer; thymoquinone; erucin; cell motility*

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