

# **POSTER PRESENTATION**

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# Effects of specific nutrients on tax-dependent activation of NF- $\kappa$ B and MMP-9 in human T-cell lymphotropic virus -1 positive malignant T-lymphocytes

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### **Background**

Adult T-cell Leukemia (ATL) is a disease with no known cure so farand it is a resistant to chemotherapy. The virus can be transmitted by exchange of bodily fluids through the placenta and from mother to child. Only 5% of those who are infected develop the disease after a long latency period ranging from 30-50 years [1,2]. The disease manifests itself as an aggressive proliferation of CD4+ cells with the human T-cell Lymphotropic virus type 1 (HTLV-1) [3]. The leukemogenesis of the virus is mainly attributed to the viral oncoprotein, Tax, that activates the Nuclear Factor kappa B (NF- $\kappa$ B) which in turn stimulates the activity and expression of the matrix metalloproteinase-9 (MMP-9)which is important in angiogenesis[2]. Our previous work has shown that using non-cytotoxic concentrations of a Specific Nutrient Synergy (SNS)mixture resulted in the induction of apoptosis in both HTLV-1 positive and negative malignant T-lymphocytes [1]. The objective of this study is to investigate the efficacy of SNS on Tax expression, NF- $\kappa$ B levels as well as on MMP-9 activity and expression both at the transcriptional and translational levels intwo HTLV-1 positive cell lines, HuT-102 and C91-PL.

## **Materials and methods**

Cell growth, experimental design, source of SNS, preparation and storage of stock solution, were previously described by our group [1]. The effects of non-cytotoxic concentrations of SNS ranging from 0-350 µg/ml were evaluated for their efficacy on proliferation, Tax expression, NF- $\kappa$ Bmobility and the activity and expression of MMP-9 at 48h and 96hof incubation. Cytotoxicity of EGCG was assayed using CytoTox 96 Non-radioactive and proliferation was measured using Cell Titer96<sup>TM</sup> Nonradioactive Cell Proliferation kit( MTT- based assay). Elisa and EMSA were used to assess the effect of SNS on NF- $\kappa$ Bmobility. Zymography was used to determine the effects of SNS on the activity and secretion of MMP-9. The expression of MMP-9 was done using RT-PCR at the translational level and Immunoblottingat the transcriptional level.

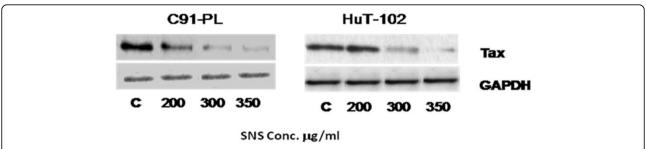
#### Results

A significantinhibition of proliferation was seen in both cell lines starting at a concentration of  $200\mu g/ml$  and in a dose dependent manner. SNS induced a dose dependent decrease in Tax expression (Fig.1),which was paralleled by a down-regulation of the nuclearization of NF- $\kappa$ B (Fig.2). This culminated in the inhibition of the activity of MMP-9 and their expression both at the transcriptional and translational levels (Fig.3).

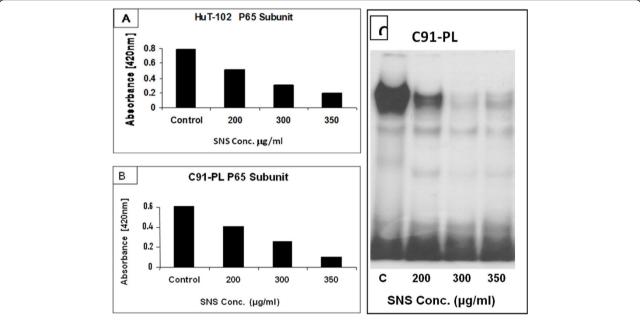
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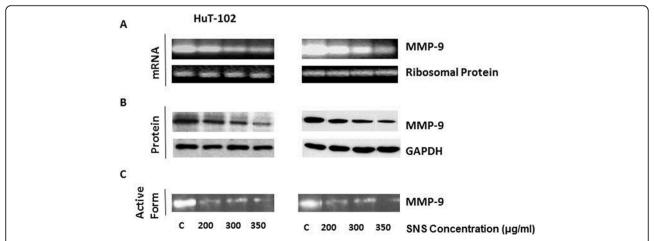
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**Figure 1** Effect of SNS on Tax expression in C91-PL and HuT-102 cell lines. Equal loading was ensured using GAPDH. The immunoblots represent results obtained in one of three independent experiments.



**Figure 2** Effect of SNS on NF- $\kappa$ B nuclear translocation in HuT-102 and C91-PL HTLV-1 positive cell lines. (A,B) Nuclear levels of P65 subunit of NF- $\kappa$ B was evaluated by ELISA at 48 and 96 h of incubation. Each value is the mean  $\pm$  SD deduced from three separate experiments done in triplicate. (C) EMSA Gel representing one of three independent experiments with nuclear extracts of C91-PL.



**Figure 3** Effect of SNS on MMP-9 mRNA (a), protein (b) and activity (c) in two ATL-HTLV-1 positive cell lines. Equal loading was ensured using ribosomal protein for mRNA expression (a) and GAPDH for protein expression (b). The results represent one out of three independent experiments.

#### **Conclusions**

The role of nutrients in the treatment of disease has been overlooked for a long time. Recently, it has been recognized that nutrients play a crucial role in the outcome of the treatment. The results of this study indicate that a specific nutrient synergy targeted multiple levels pertinent to the progression of ATL. Its activity was mediated through the NF- $\kappa$ B pathway, and hence has the potential to be integrated in the treatment of this disease as a natural, yet potent anticancer agent.

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