



CURCUMIN ACTION AGAINST PRE-CANCEROUS LESION

DR.M.ELUMALAI* , DR. S.BHUMINATHAN AND DR. D. RAMYAA

*Department of Pharmacology and Prosthodontics, Sree Balaji Dental College and Hospital,
Bharath University, Pallikaranai, Chennai-600100, Tamil Nadu, India.*

ABSTRACT

Oral precancerous lesions are one of the most common lesions present worldwide today. They are usually neglected by the common population when compared to cancerous lesion. However, they also may be extremely fatal which changes as cancer if left untreated at a very initial stage of the lesion. Early detection and treatment gives the best chance for its cure. The detection and diagnosis are currently based on clinical examination, histopathological evaluation. Several medications have been developed over the years for early cure of oral precancerous lesion. The purpose of this article is to review the available curcumin herbal to treat these precancerous lesions.

KEY WORDS: curcumin herbal, oral precancerous lesions, anti-oxidant, apoptosis



*Corresponding author



DR.M.ELUMALAI

Department of Pharmacology and Prosthodontics, Sree Balaji Dental College and Hospital,
Bharath University, Pallikaranai, Chennai-600100, Tamil Nadu, India.

INTRODUCTION

Extensive research within the past half-century has indicated that curcumin (diferuloylmethane), a yellow pigment in curry powder, exhibits anti-oxidant, anti-inflammatory, and pro-apoptotic activities. The anti-pre-cancer activities assigned to curcumin are mediated through an anti-oxidant and DNA protecting mechanism¹. Values for serum and salivary vitamins C and E showed a significant decrease in oral leukoplakia, submucous fibrosis and lichen planus, in contrast to healthy individuals, but increased significantly in all groups subsequent to curcumin administration after clinical cure of lesions. Oral leukoplakia and submucous fibrosis are two major precancerous lesions, but only 8-10% of these lesions which ultimately become malignant². The ability to clinically predict malignant transformation is limited and routine histopathological diagnosis has limited prognostic value. The presence of epithelial dysplasia is an important parameter used in the prognostication of leukoplakia. However, there are limitations in its usage where all lesions exhibiting dysplasia do not eventually become malignant and some may even regress, and carcinoma can develop from lesions in which epithelial dysplasia was not diagnosed in previous biopsies³. Therefore, it is necessary to develop other methods for predicting the malignant potential of pre-malignant lesions and preventive Curcumin, the principal curcuminoid found in turmeric, which is generally considered to be an active component. Other curcuminoids found in turmeric include demethoxycurcumin and bisdemethoxycurcumin. measures⁴.

Antioxidant Activity

Curcumin is an effective scavenger of reactive oxygen species and reactive nitrogen species in the test tube (*in vitro*)^{5, 6}. However, the finding that oral curcumin supplementation (3.6 g/day) for seven days decreased the number of oxidative DNA adducts in malignant colorectal tissue suggests that curcumin taken orally may reach sufficient concentrations in the gastrointestinal tract to inhibit oxidative DNA damage^{5,6}.

Anti-inflammatory Activity

The metabolism of arachidonic acid in cell membranes plays an important role in the inflammatory response by generating potent chemical messengers known as eicosanoids⁷. Curcumin has been found to inhibit PLA2, COX-2, and 5-LOX activities in cultured cells⁸. Although curcumin inhibited the catalytic activity of 5-LOX directly, it inhibited PLA2 by preventing its phosphorylation and COX-2 mainly by inhibiting its transcription. In inflammatory cells, such as macrophages, iNOS catalyzes the synthesis of nitric oxide, which can react with superoxide to form peroxynitrite, a reactive nitrogen species that can damage proteins and DNA. Curcumin has been found to inhibit NF-κB-dependent gene transcription⁹ and the induction of COX-2 and iNOS in cell culture and animal studies^{10,11}

Induction of Cell Cycle Arrest and Apoptosis

After a cell divides, it passes through a sequence of stages—collectively known as the cell cycle—before it can divide again. Following DNA damage, the cell cycle can be transiently arrested to allow for DNA repair or, if the damage cannot be repaired, for activation of pathways leading to cell death (apoptosis)¹². Curcumin has been found to induce cell-cycle arrest and apoptosis in a variety of cancer cell lines grown in culture¹³⁻¹⁷. The mechanisms by which curcumin induces apoptosis are varied but may include inhibitory effects on several cell-signaling pathways. However, studies have found that curcumin induces apoptosis in cancer cells. Curcumin inhibited apoptosis induced by the tumor suppressor protein p53 in cultured human colon cancer cells^{18,19}, and one study found that curcumin inhibited apoptosis induced by several chemotherapeutic agents in cultured breast cancer cells at concentrations of 1-10 micromoles/liter²⁰. The antiprecancerous effects of curcumin are mediated through an antioxidant mechanism. Serum and salivary vitamin C and E levels were found to increase, while MDA and 8-OHdG levels decreased in oral leukoplakia, submucous fibrosis and lichen-planus patients after intake of curcumin, as compared to pre-

treatment levels¹. The mechanisms by which curcumin mediates its prooxidant effects remain unclear. It has been suggested that mitochondria play a role in curcumin-induced apoptosis. It is possible that curcumin activates the mitochondrial enzymes that lead to the production of reactive oxygen species (ROS)^{21, 22}. The induction of ROS by curcumin may occur through its interaction with thioredoxin reductase, thus altering its activity to NADPH oxidase, which could lead to the production of ROS²³. The mechanism of action of TO in oncoprevention can be speculated to be through its antioxidant action and protection against DNA damage. An adverse effect on lipids was noticed in only one case in this study. Prakasunand et al have reported

on the biochemical safety of turmeric, 3gms/day, in 54 cases of pepticulcer²⁴

CONCLUSION

Curcumin acts against transcription factors, which are like a master switch," said lead researcher, Bharat Aggarwal. "Transcription factors regulate all the genes needed for tumors to form. When we make the genes inactive that are involved in the growth and invasion of cancer cells. Curcumin helps the body to destroy mutated cancer cells, so they cannot spread in the body. Thus curcumin is best herbal for treating pre-cancerous lesion.

REFERENCES

- Balwant Rai, Jasdeep Kaur, Reinhilde Jacobs and Jaipaul Singh. (2010). Possible action mechanism for curcumin in pre-cancerous lesions based on serum and salivary markers of oxidative stress. *J Oral Sci.* 52(2):251-6.
- Gupta PC, Bhonsle RB, Murti PR, Daftary DK, Mehta FS, Pindborg JJ (1989). An epidemiological assessment of cancer risk in oral precancerous lesions in India with special reference to nodular leucoplakia. *Cancer* 63, 2247-2252.
- Reibel J (2003). Prognosis of oral pre-malignant lesions: significance of clinical, histopathological, and molecular biological characteristics. *Crit Rev Oral Biol Med* 14, 47-62.
- Allison P, Locher S, Feine JS (1998). The role of diagnostic delays in the prognosis of oral cancer: review of literature. *Oral Oncol* 34, 161-170.
- Sreejayan, Rao MN. (1997). Nitric oxide scavenging by curcuminoids. *J Pharm Pharmacol.* 49(1):105-107. (PubMed)
- Sreejayan N, Rao MN. (1996). Free radical scavenging activity of curcuminoids. *Arzneimittelforschung.* 46(2):169-171. (PubMed)
- Steele VE, Hawk ET, Viner JL, Lubet RA. (2003). Mechanisms and applications of non-steroidal anti-inflammatory drugs in the chemoprevention of cancer. *Mutat Res.* 523-524:137-144. (PubMed)
- Hong J, Bose M, Ju J, et al. (2004). Modulation of arachidonic acid metabolism by curcumin and related beta-diketone derivatives: effects on cytosolic phospholipase A(2), cyclooxygenases and 5-lipoxygenase. *Carcinogenesis.* 25(9):1671-1679. (PubMed)
- Plummer SM, Holloway KA, Manson MM, et al. (1999). Inhibition of cyclo-oxygenase 2 expression in colon cells by the chemo preventive agent curcumin involves inhibition of NF-kappa activation via the NIK/IKK signalling complex. *Oncogene.* 18(44):6013-6020. (PubMed)
- Brouet I, Ohshima H. (1995). Curcumin, an anti-tumour promoter and anti-inflammatory agent, inhibits induction of nitric oxide synthase in activated macrophages. *Biochem Biophys Res Commun.* 206(2):533-540. (PubMed)
- Nanji AA, Jokelainen K, Tipoe GL, Rahemtulla A, Thomas P, Dannenberg AJ. (2003). Curcumin prevents alcohol-induced liver disease in rats by inhibiting the expression of NF-kappa B-dependent genes. *Am J Physiol Gastrointest Liver Physiol.* 284(2):G321-327. (PubMed)
- Stewart ZA, Westfall MD, Pietenpol JA. (2003). Cell-cycle dysregulation and anticancer therapy. *Trends Pharmacol Sci.* 24(3):139-145. (PubMed)
- Duvoix A, Blasius R, Delhalle S, et al. (2005) Chemopreventive and therapeutic

- effects of curcumin. *Cancer Lett.* 223(2):181-190. (PubMed)
14. Surh YJ, Chun KS. (2007). Cancer chemo preventive effects of curcumin. *AdvExp Med Biol.* 595:149-172. (PubMed)
 15. Singh S, Khar A. (2006). Biological effects of curcumin and its role in cancer chemoprevention and therapy. *Anticancer Agents Med Chem.* 6(3):259-270. (PubMed)
 16. Kuttan G, Kumar KB, Guruvayoorappan C, Kuttan R. (2007). Antitumor, anti-invasion, and antimetastatic effects of curcumin. *AdvExp Med Biol.* 595:173-184. (PubMed)
 17. Kunnumakkara AB, Anand P, Aggarwal BB. (2008). Curcumin inhibits proliferation, invasion, angiogenesis and metastasis of different cancers through interaction with multiple cell signalling proteins. *Cancer Lett.* 269(2):199-225. (PubMed)
 18. Moos PJ, Edes K, Mullally JE, Fitzpatrick FA. (2004). Curcumin impairs tumor suppressor p53 function in colon cancer cells. *Carcinogenesis.* 25(9):1611-1617. (PubMed)
 19. Tsvetkov P, Asher G, Reiss V, Shaul Y, Sachs L, Lotem J. (2005). Inhibition of NAD(P)H:quinoneoxidoreductase 1 activity and induction of p53 degradation by the natural phenolic compound curcumin. *ProcNatlAcadSci U S A.* 102(15):5535-5540. (PubMed)
 20. Somasundaram S, Edmund NA, Moore DT, Small GW, Shi YY, Orlowski RZ. (2002). Dietary curcumin inhibits chemotherapy-induced apoptosis in models of human breast cancer. *Cancer Res.* 62(13):3868-3875. (PubMed)
 21. Uddin S, Hussain AR, Manogaran PS, Al-Hussein K, Platanias LC, Gutierrez MI, Bhatia KG (2005). Curcumin suppresses growth and induces apoptosis in primary effusion lymphoma. *Oncogene* 24,7022-7030.
 22. Atsumi T, Fujisawa S, Tonosaki K (2005). Relationship between intracellular ROS production and membrane mobility in curcumin- and tetrahydrocurcumin-treated human gingival fibroblasts and human submandibular gland carcinoma cells. *Oral Dis* 11, 236-242.
 23. Fang J, Lu J, Holmgren A (2005). Thioredoxinreductase is irreversibly modified by curcumin: a novel molecular mechanism for its anticancer activity. *J BiolChem* 280, 25284-25290.
 24. Prackasunand C, Indrasukhsri B, Leethochawalit M, Hungspreugs K. (2001). Phase II clinical trial on effect of the long turmeric (*Curcuma longa* Linn) on healing of peptic ulcer. *Southeast Asian J Trop Med Public Health*; 32:208-15.