**Review Article** 

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# **CURCUMIN: A PLEIOTROPIC DRUG**

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#### ABSTRACT

Curcumin in the form of Turmeric powder along with hot milk is widely consumed in India for its antiinflammatory effects. Curcumin is widely used in traditional Indian ayurvedic medicine to treat hepatic disorders, anorexia, cough, diabetic wounds, rheumatoid arthritis, and sinusitis. Turmeric paste in slaked lime is a popular home remedy for the treatment of inflammation and wounds. Ancient texts of Indian medicine describe the use of curcumin in inflammatory diseases, wound healing, and abdominal problems. Curcumin, exhibits pleiotropic effects such as anti-inflammatory, antioxidant, anticancer, antiviral and neurotropic activity and therefore holds a promise as a therapeutic agent to prevent and treat several diseases. The purpose of this review is to provide a brief overview of the plethora of research regarding the five major pleiotropic effects of curcumin.

### **KEYWORDS**

Curcumin; Turmeric; Antioxidant; Anti-inflammatory; Anticancer; Antiviral; Neurotropic.

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#### **INTRODUCTION**

Curcumin, a constituent from the rhizome of the herb Curcuma Longa (turmeric), has received widespread attention over the past few decades<sup>1</sup>. Excitement originated after encouraging in-vitro discoveries showed its influence on many biological mechanisms associated with several diseases. In particular, in vitro and animal models confirmed that curcumin has Pleiotropic effects (shown in Figure No.1) anti-inflammatory, antioxidant, anticancer, antiviral and neurotropic effects just to name a few<sup>1</sup>. Curcumin was first extracted from turmeric in 1815 by Vogel and Pelletier. The two scientists Milobedzka German and Lampe determined its chemical structure in  $1910^2$ . Curcumin (1, 7-bis (4-hydroxy-3-methoxyphenyl)-

1, 6-heptadiene-3, 5-dione) is also known as diferuloylmethane<sup>3</sup>. It has been shown to benefit inflammatory conditions<sup>4</sup>, metabolic syndrome<sup>5</sup>, pain<sup>6</sup>, and to helps in the management of inflammatory and degenerative eye conditions<sup>7,8</sup>. Curcuma longa has been traditionally used in Asian countries as a medical herb due to its antioxidant, anti-inflammatory properties<sup>9</sup>. Curcumin, а polyphenol, has been shown to target multiple signaling molecules while also demonstrating activity at the cellular level, and thus support its multiple health benefits<sup>9</sup>. The yellow color of turmeric is mainly due to the presence of polyphenolic curcuminoids, constitute approximately 3% to 5% of most turmeric preparations. The alcoholic extract of turmeric mainly contains three curcuminoids, namely curcumin (also referred as curcumin I or diferuloylmethane), desmethoxycurcumin (curcumin II), and bisdesmethoxy curcumin (curcumin III)<sup>10,11</sup>.

In a study from in India, daily administration of 1,200 - 2,100 mg of oral curcumin to patients with rheumatoid arthritis for 2–6 weeks did not cause any toxicity<sup>12</sup>. In another study of high-dose oral curcumin (500, 1,000, 2,000, 4,000, and 8,000 mg) daily for three months to patients with pre-invasive malignant or high risk premalignant conditions, no noticeable adverse effects were detected<sup>13</sup>.

The clinical trials conducted so far have indicated the therapeutic potential of curcumin against a wide range of human diseases. It has also shown protection against hepatic conditions, chronic arsenic exposure and alcohol intoxication. It is marketed in several forms including capsule, tablets ointment, energy drinks soaps and cosmetics<sup>14,15</sup>.

# 2. Pleiotropic Effects

# 2.1. Anti-inflammatory

The anti-inflammatory effect of curcumin is most likely mediated through its ability to inhibit cyclooxygenase-2 (COX-2), lipoxygenase (LOX), and inducible nitric oxide synthase (INOS). COX-2, LOX, and INOS are important enzymes that mediate inflammatory processes. Improper up regulation of COX-2 and/or INOS has been

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associated with the pathophysiology of certain types of human cancer as well as inflammatory disorders. Because inflammation is closely linked to tumor promotion, curcumin with its potent antiinflammatory property is anticipated to exert chemo-preventive effects on carcinogenesis. Hence, the past few decades have witnessed intense research devoted to the antioxidant and antiinflammatory properties of curcumin<sup>16</sup>.

Tumor necrosis factor  $\alpha$  (TNF- $\alpha$ ) is a major mediator of inflammation in most diseases, and this effect is regulated by the activation of a transcription factor, nuclear factor (NF)-KB. Whereas, TNF- $\alpha$  is said to be the most potent NF- $\kappa B$  activator, the expression of TNF- $\alpha$  is also regulated by NF- $\kappa$ B. In addition, TNF- $\alpha$ , NF- $\kappa$ B is also activated by most inflammatory cytokines; gram (-ve) bacteria; various disease-causing viruses: environmental pollutants; chemical. physical, mechanical, and psychological stress; high glucose; fatty acids; ultraviolet radiation; cigarette smoke; and other disease-causing factors. Therefore, agents that down regulate NF-KB and NF-kB-regulated gene products have potential efficacy against several of these diseases. Curcumin has been shown to block NF-kB activation increased by several different inflammatory stimuli<sup>17</sup>. This study demonstrated that curcumin had anti-inflammatory activity in LTA-stimulated microglial cells may through inhibiting NF-κB and p38 MAPK activation, and may induce the expression of Nrf2 and HO-1. Furthermore, curcumin does not have cytotoxic effects in BV-2 microglial cells at its anti-inflammatory dose. Curcumin may have therapeutic potential for some neuro inflammation-associated disorders caused by Gram-positive bacteria<sup>18</sup>.

# 2.2. Antioxidant

Oxidative damage is believed to be one of the mechanisms behind aging and many diseases. It involves free radicals, highly reactive molecules with unpaired electrons<sup>19</sup>. Many people consume antioxidants as a defense against oxidative stress. Antioxidants in the form of commercial food additives have been manufactured synthetically and may contain high amounts of preservatives<sup>20</sup>. January – March 33

Identifying potential natural antioxidant sources can be a useful alternative to ensure sound health $^{21}$ . Anti-oxidant constituents in plant material have piqued the interest of scientists, food manufacturers, cultivators, and consumers for their roles in the maintenance of human health<sup>20</sup>. Curcumin is a lipophilic compound, making it an efficient scavenger of peroxyl radicals; However, the actual reaction site and the mechanism of free radical scavenging have not been clarified yet<sup>22</sup>. It is generally assumed that the phenolic moieties are responsible for radical scavenging properties of the pertinent antioxidant reactants. The clinical use of curcumin is limited because of its low bioavailability, due to the hydrophobic nature of the molecule $^{23}$ .

### 2.3. Anticancer

Over half of the publications were published since 2014, which mainly focused on the effects of against cancer, inflammation, curcumin and oxidative stress. Frequently investigated cancer types were breast, colon, colorectal, pancreatic, and prostate cancer<sup>24</sup>. Curcumin has shown considerable anticancer effects against several different types of cancer, including prostate cancer, breast cancer, colorectal cancer, pancreatic cancer, and head and neck cancer both *in vitro* and *in vivo*<sup>25</sup>. Cancer is the second most life-threatening disease and one of the main public health problems worldwide. In 2018, there were around 1.73 million new cases of cancer and more than 609,000 deaths in the United States alone<sup>25</sup>. Curcumin and its derivatives have received immense attention in the past two decades due to their bio functional properties such as anti-tumor, antioxidant, and anti-inflammatory activities<sup>26</sup>.

A recent report suggests that curcumin inserts deep into the cellular membrane in a trans bilayer orientation, anchored by hydrogen bonding to the phosphate group of lipids, thus inducing negative curvature in the bilayer<sup>27</sup>. The promotion of negative curvature by curcumin may have a direct effect on apoptosis by increasing the permeabilizing activity of the apoptotic protein tBid<sup>28</sup>. Curcumin has been shown to suppress multiple signaling pathways and inhibit cell proliferation, invasion, metastasis, and angiogenesis. The chemo preventive

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action of curcumin might be due to its ability to induce apoptosis by several pathways. Curcumin directly or indirectly controls different gene or gene products involved in cell death pathways.

Curcumin inhibits the STAT3 (Signal transducer and activator of transcription 3) and NF-kB signaling pathways, which play key-roles in cancer development and progression. Constitutive activation of the STAT3 and NF-kB signaling pathways has been demonstrated in prostate cancer cell lines and clinical samples of prostate cancer<sup>29-</sup> <sup>32</sup>. The nuclear factor (NF)- $\kappa$ B, is a ubiquitous transcription factor that regulates many genes implicated in growth regulation, inflammation, carcinogenesis, and apoptosis. *In vitro* and *in* vivo studies have documented that constitutive activation of NF-kB results in inhibition of chemotherapy-induced apoptosis in a number of cancer cells. Signal transducer and activator of transcription 3 (STAT3) is a ubiquitously expressed member of the STAT family of transcription factors activated that is by tyrosine phosphorylation *via* upstream receptors, such as epidermal growth factor (EGF), platelet-derived growth factor (PDGF) and cytokines, such as interleukin-6 (IL-6). Recent studies have demonstrated that STAT3 may confer cancer resistance to chemotherapeutic agents<sup>33</sup>.

# 2.4. Antiviral

Curcumin has been studied as an antiviral agent against numerous viruses like Herpes simplex virus (HSV), dengue virus (DENV), Human Immunodeficiency Virus (HIV), Hepatitis viruses, influenza type A virus, and Ebola virus<sup>34</sup>. Dengue is the most common infective disease caused by dengue virus (DENV). Until now, there is no specific antiviral drug for dengue infection. Curcumin as major active compound has been identified for its antiviral effect<sup>35</sup>. A number of research studies and scientific evidence suggest that turmeric has a potent antiviral activity<sup>36</sup> Curcumin, a relatively non-toxic natural product isolated from Curcuma longa, is a modest inhibitor of the HIV-1 and HIV-2 proteases. Boron complexes of curcumin cause time-dependent inactivation of the HIV-1 and HIV-2 proteases<sup>37</sup>. Oncogenesis by January – March 34

human T-cell leukemia virus type-1 as an etiology factor of adult t-cell leukemia (ATL) Is critically dependent of the activator protein-1 (AP-1)<sup>40</sup>. DNA Binding and transcriptional effect of AP-1 HTLV-1-Infected T-Cell lines were suppressed by curcumin treatment curcumin also inhibited the expression of junD protein as an important factor in AP-1-DNA complex in HTLV-1 infected T-cells as well as HTLV-1(Human T-Cell Leukemia Virus Type 1) Tax-induced AP-1 transcriptional effect. Cell cycle arrest and including of apoptosis were found to be possible mechanism against HTLV-1 application in infected T cell line by curcumin. Suppression of AP-1 activity possibly trough decreasing the expression of Jun D protein is introduces as possible pathway of anti-ATL activity of curcumin<sup>38</sup>.

#### **2.5. Neurotropic Effect**

Diets rich in curcumin have been reported to be associated with a lower risk of Alzheimer's disease (AD) or cognitive impairment in epidemiological studies. *In vitro* and *in vivo* experimental studies with AD models have indicated that curcumin has anti-amyloidogenic effects in addition to their antioxidant and anti-inflammatory effects. In addition, it has been reported that curcumin has neuroprotective and anti-inflammatory effects in *Invitro* and *In vivo* experimental studies on Parkinson's disease<sup>39</sup>.

Findings indicate that curcumin can enhance postsynaptic electrical reactivity and cell viability in intact neural circuits with antidepressant-like effects, possibly through the up regulation of BDNF (Brain-derived neurotrophic factor)) and reduction brain<sup>40</sup>. inflammatory factors in the of In Alzheimer's disease (AD), a peptide called  $\beta$ amyloid (A $\beta$  peptide) aggregates into oligomers and fibrils and forms deposits known as amyloid (or senile) plaques outside neurons in the hippocampus and cerebral cortex of patients. Another feature of AD is the accumulation intracellular of

neurofibrillary tangles formed by phosphorylated protein<sup>41</sup>. Abnormal Tau microglial activation, oxidative stress, and neuronal death are also associated with the progression of the disease. Curcumin has been found to inhibit AB fibril formation and extension and to destabilize preformed fibrils *in vitro*<sup>42-44</sup>. Metal chelation by curcumin might interfere with metal ion  $(Cu^{2+}/Zn^{2+})$ -induced AB aggregation. Curcumin might also affect the trafficking of Aβ peptide precursor (APP) and the generation of  $A\beta$ peptides from APP<sup>45-46</sup>. Abnormally activated microglia and hypertrophic astrocytes around amyloid plaques in AD brains release cytotoxic molecules. such as proinflammatory cytokines and ROS (reactive oxygen species), which enhance  $A\beta$  formation and deposition and further damage neurons. Curcumin was found to reduce the inflammatory response triggered by Αβ peptide-induced microglial activation and increase neuronal cell survival<sup>47</sup>. When injected into the carotid artery of a transgenic mouse model of AD, curcumin was found to cross the blood-brain barrier, bind to amyloid plaques, and block the formation of AB oligomers and fibrils<sup>45</sup>. In other animal models of AD, dietary curcumin decreased biomarkers of inflammation and oxidative damage, increased  $A\beta$  peptide clearance by macrophages, dismantled amyloid plaques in the brain, stimulated neuronal cell growth in the hippocampus, and improved Aβinduced memory deficits<sup>48</sup>.



Figure No.1: Mechanism of action of curcumin in various conditions

# CONCLUSION

Recently, many natural substances have been increasingly found to have significant biological properties which make them useful in the treatment of various diseases. Nature has plenty of substances having healing properties making them invaluable for its role towards promotion of a healthy world. Curcumin has received worldwide attention for its multiple health benefits, reported to act primarily through its pleiotropic effect. From preclinical studies to clinical trials, tremendous progress has been made with respect to this compound in the medical field. Research suggests that curcumin could help in the management of oxidative and inflammatory conditions, metabolic syndrome, arthritis, anxiety, and hyperlipidemia. It may also help in the management of exercise-induced inflammation and muscle soreness, thus enhancing recovery and subsequent performance in active people. Thus, we can conclude that a relatively low dose of curcumin or in the form of turmeric powder could provide health benefits for people that do not have diagnosed health conditions.

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### **CONFLICT OF INTEREST**

We declare that we have no conflict of interest.

# BIBLIOGRAPHY

- 1. Adrian L Lopresti. The Problem of Curcumin and Its Bioavailability: Could Its Gastrointestinal Influence Contribute to Its overall Health-Enhancing Effects?, *Adv Nutr.*, 9(1), 2018, 41-50.
- Muhammad Imran, Muhammad Nadeem, Muhammad Asif Khan, Sheraz Ahmed, Ali Imran, Rai Muhammad Amir, Muhammad Umair Arshad, Syed Amir Gilani, Farhan Saeed, Abdur Rauf, Zaffar Mehmood, Shaista Khan and Hafiz Ansar Rasul Suleria. Curcumin and its Allied Analogues: Epigenetic and Health Perspectives – a

Review, Czech J. Food Sci., 35(4), 2017, 285-310.

- 3. Panahi Y, Hosseini M S, Khalili N, Naimi E, Majeed M and Sahebkar A: Antioxidant and anti-inflammatory effects of Curcuminoid-piperine combination in subjects with metabolic syndrome: а randomized controlled trial and an updated meta-analysis, Clinical Nutrition, 34(6), 2015, 1101-1108.
- 4. Aggarwal B B, Harikumar K B. Potential therapeutic effects of curcumin, the antiinflammatory agent, against neurodegenerative, cardiovascular, pulmonary, metabolic, autoimmune and neoplastic diseases, *Int. J. Biochem. Cell Biol.*, 41(1), 2009, 40-59.
- Panahi Y, Hosseini M S, Khalili N, Naimi E, Simental-Mendia L E, Majeed M, Sahebkar A. Effects of curcumin on serum cytokine concentrations in subjects with metabolic syndrome: A post-hoc analysis of a randomized controlled trial, *Biomed. Pharmacother.*, 82, 2016, 578-582.
- Kuptniratsaikul V, Dajpratham P, Taechaarpornkul W, Buntragulpoontawee M, Lukkanapichonchut P, Chootip C, Saengsuwan J, Tantayakom K, Laongpech S. Efficacy and safety of *Curcuma domestica* extracts compared with ibuprofen in patients with knee osteoarthritis: A multicenter study, *Clin. Interv. Aging.*, 9, 2014, 451-458.
- 7. Mazzolani F, Togni S. Oral administration of a curcumin-phospholipid delivery system for the treatment of central serous chorioretinopathy: A 12-month follow-up study, *Clin. Ophthalmol.*, 7, 2013, 939-945.
- Allegri P, Mastromarino A, Neri P. Management of chronic anterior uveitis relapses: Efficacy of oral phospholipidic curcumin treatment. Long-term followup, *Clin. Ophthalmol.*, 4, 2010, 1201-1206.
- 9. Lestari M L, Indrayanto G. Curcumin. Profiles Drug Subst. Excip. Relat. Methodol., 39, 2014, 113-204.

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- 10. Vogel H A, Pelletier J. Curcumin-biological and medicinal properties, *J. Pharma.*, 2, 1815, 50.
- 11. Milobedzka J, Kostanecki V, Lampe V. Structure of curcumin, *Ber. Dtsch. Chem. Ges.*, 43, 1910, 2163-2170.
- 12. Deodhar S D, Sethi R, Srimal R C. Preliminary study on anti-rheumatic activity of curcumin (diferuloyl methane), *Indian J. Med. Res.*, 71, 1980, 632-634.
- Cheng A L, Hsu C H, Lin J K, Hsu M M, Ho Y F, Shen T S, Ko J Y, Lin J T, Lin B R, Ming-Shiang W. Phase I clinical trial of curcumin, a chemopreventive agent, in patients with high-risk or pre-malignant lesions, *Anticancer Res.*, 21, 2001, 2895-2900.
- 14. He Y, Yee Y, Zheng X, Zhangk, Chens, DVZ. Curcumin, inflammation and Chronic disease: How are they linked, *Molecule*, 20, 2015, 9183-9213.
- 15. Akila Srinivasan, Sandhiya Selvarajan and Steven Aibor Dkhar. The Pleiotropic Effects of Curcumin, *Int. J. Adv. Res.*, 5(7), 2749-2756, 2320-5407.
- 16. Venugopal Padmanabhan Menon, Adluri Ram Sudheer. Antioxidant and antiinflammatory properties of curcumin, 2007.
- Panahi Y, Hosseini M S, Khalili N, Naimi E, Simental-Mendia L E, Majeed M, Sahebkar A. Effects of curcumin on serum cytokine concentrations in subjects with metabolic syndrome: A post-hoc analysis of a randomized controlled trial, *Biomed. Pharmacother.*, 82, 2016, 578-582.
- 18. Yangyang Yu, Qian Shen, Yihong Lai, Sun Y. Park, ingmei Ou, Dongxu Lin, Meiling Jin and Weizhen Zhang1. Antiinflammatory Effects of Curcumin in Microglial Cells Front, *Pharmacol.*, 20 April 2018.
- 19. Kris Gunnars, BSc 10 Proven Health Benefits of Turmeric and Curcumin on July 13, 2018.
- 20. E. M. Tanvir, R. Afroz, N. Karim et al., "Antioxidant and antibacterial activities of

methanolic extract of BAU kul (*Ziziphus mauritiana*), an improved variety of fruit from Bangladesh," *Journal of Food Biochemistry*, 39, 2015, 139-147.

- 21. Afroz R, Tanvir E, Zheng W and Little P. "Molecular Pharmacology of Honey," *Journal of Clininal Experimental Pharmacology*, 6(3), 2016.
- 22. Dora Tatraaljai, Balazs Kirschweng, Janos Kovacs, Eniko Foldes, Bela Pukanszky. Processing Stabilization of PE with a Natural Antioxidant, Curcumin.
- 23. Aggarwal B B, Sundaram C, Malani N, Ichikawa H. *Adv. Exp. Med. Biol.*, 595, 2007, 1-75.
- 24. Andy Wai Kan Yeung, Michal Horba nczuk, Nikolay T. Tzvetkov, Andrei Mocan, Simone Carradori, Filippo Maggi, Joanna Marchewka, Stefania Sut, Stefano Dall Acqua, Ren-You Gan, Lyubka P. Tancheva, Timea Polgar, Ioana Berindan-Neagoe, Vasil Pirgozliev, Karel Smejkal and Atanas G. Atanasov. Curcumin: *Total-Scale Analysis of the Scientific Literature*, 2019.
- 25. Mhd Anas Tomeh, Roja Hadianamrei and Xiubo Zhao A. Review of Curcumin and Its Derivatives as Anticancer Agents, *Int J Mol Sci.*, 20(5), 2019, 1033.
- 26. Nagahama K, Utsumi T, Kumano T, Maekawa S, Oyama N, Kawakami J. Discovery of a new function of curcumin which enhances its anticancer therapeutic potency, *Sci. Rep.*, 6, 2016, 30962.
- 27. Barry J, Fritz M, Brender J R, Smith P E, Lee D K, Ramamoorthy A. Determining the effects of lipophilic drugs on membrane structure by solid-state NMR spectroscopy: the case of the antioxidant curcumin, *J Am Chem Soc.*, 131, 2009, 4490-8.
- 28. Epand R F, Martinou J C, Fornallaz-Mulhauser M, Hughes D W, Epand R M. The apoptotic protein tBid promotes leakage by altering membrane curvature, *J Biol Chem.*, 277, 2002, 32632.
- 29. Kasdagly M, Radhakrishnan S, Reddivari L, Veeramacheni DN, Vanamala J. Colon

Available online: www.uptodateresearchpublication.com

carcinogenesis: Influence of Western dietinduced obesity and targeting stem cells using dietary bioactive compounds, *Nutrition*, 11-12: 1242-1256, 2014.

- 30. Anthwal A, Thakur B K, Rawat M S, Rawat D S, Tyagi A K, Aggarwal B B. Synthesis, characterization and *in vitro* anticancer activity of C-5 curcumin analogues with potential to inhibit TNF-a-induced NF-kB activation, *Biomed Res Int*, 2014, 524161.
- 31. Chiablaem K, Lirdprapamongkol K, Keerati chamroen S, Surarit R, Svasti J. Curcumin suppresses vasculogenic mimicry capacity of hepatocellular carcinoma cells through STAT3 and PI3K/AKT inhibition, Anticancer Res, 4, 1857-1864, 2014.
- Abdulghani J, Gu L, Dagvadorj A: Stat3 promotes metastatic progression of prostate cancer, *Am J Pathol*, 172, 2008, 1717-1728, 2008.
- 33. Natalia G. Vallianou, Angelos Evangelopoulos, Nikos Schizas and Christos Kazazis Potential Anticancer Properties and Mechanisms of Action of Curcumin.
- 34. Dony Chacko Mathew, Wei-Li Hsu. "Antiviral potential of curcumin", *Journal of functional food*, 40(692:699).
- 35. Ichsyani M, Ridhanya A, Risanti M, Desti H, Ceria R, Putri D H, Sudiro T M and Dewi B E. "Antiviral effects of *Curcuma longa L.*, against dengue virus *in vitro* and *in vivo*".
- 36. Shruti (BE Biotech. and amp; PGD Clinical Research), "How Turmeric fights these 7 serious viral infections".
- 37. Sui Z, Salto R, Li J, Craik C and Ortiz de Montellano P R. "Inhibition of the HIV-1 and HIV-2proteases by curcumin and curcumin boron complexes," *Bioorganic* and amp; Medicinal Chemistry, 1(6), 415-422, 1993.
- 38. Tomita M, Kawakami H, Uchihara J N. "Curcumin suppresses constitutive activation of AP-1 by down regulation of

Jun D protein in Htlv-1 Hyprerinfected Tcell lines," *Leukemia research*, 30(3), 313-321, 2006.

- 39. Moeko Noguchi-Shinohara, Tsuyoshi Hamaguchi, and Masahito Yamada. "The potential roll of curcumin in treatment and prevention of neurological disorder".
- 40. Ga-Young Choi, Hyun-Bum Kim, Eun-Sang Hwang, Seok Lee, Min-Ji Kim, Ji-Young Choi, Sung-Ok Lee, Sang-Seong Kim and Park. Curcumin Alters Ji-Ho Neural and Plasticity Viability of Intact Hippocampal Circuits and Attenuates Behavioral Despair and COX-2 Expression in Chronically Stressed Rats. 2017, 9 pages.
- 41. Prvulovic D, Hampel H. Amyloid beta (Aβ) and phospho-tau (p-τ) as diagnostic biomarkers in Alzheimer's disease, *Clin Chem Lab Med.*, 49(3), 2011, 367-374.
- 42. Ono K, Hasegawa K, Naiki H, Yamada M. Curcumin has potent anti-amyloidogenic effects for Alzheimer's β-amyloid fibrils *in vitro*, *J Neurosci Res.*, 75(6), 2004, 742-750.
- 43. Reinke A A, Gestwicki J E. Structureactivity relationships of amyloid  $\beta$ aggregation inhibitors based on curcumin: influence of linker length and flexibility, *Chem Biol Drug Des.*, 70(3), 2007, 206-215.

- 44. Yang F, Lim G P, Begum A N, et al. Curcumin inhibits formation of amyloid  $\beta$  oligomers and fibrils, binds plaques, and reduces amyloid *in vivo*, *J Biol Chem.*, 280(7), 2005, 5892-5901.
- 45. Lin R, Chen X, Li W, Han Y, Liu P, Pi R. Exposure to metal ions regulates mRNA levels of APP and BACE1 in PC12 cells: blockage by curcumin, *Neurosci Lett.*, 440(3), 2008, 344-347.
- 46. Zhang C, Browne A, Child D, Tanzi R E. Curcumin decreases amyloid-β peptide levels by attenuating the maturation of amyloid-β precursor protein, *J Biol Chem.*, 285(37), 2010, 28472-28480.
- 47. Shi X, Zheng Z, Li J, *et al.* Curcumin inhibits A β-induced microglial inflammatory responses in vitro: Involvement of ERK1/2 and p38 signaling pathways, *Neurosci Lett.*, 594, 2015, 105-110.
- 48. Goozee K G, Shah T M, Sohrabi H R, *et al.* Examining the potential clinical value of curcumin in the prevention and diagnosis of Alzheimer's disease, *Br J Nutr.*, 2015, 1-17.

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