# **Curcumin (high purity)**

Inhibitor of Lipoxygenase and COX

Dual inhibitor of 5-lipoxygenase ( $IC_{50}$ =8µM) and cyclooxygenase (COX) ( $IC_{50}$ =52µM). Also inhibits glutathione S-transferase, induction of nitric oxide (NO) in activated macrophages and inhibits EGF-induced tyrosine phosphorylation of EGF receptors. Antioxidant. Inhibitor of NAD(P)H:quinone oxidoreductase 1 (NQO1), disrupting the binding of NQO1 to wild type p53 inducing ubiquitin-independent degradation of p53 and inhibits p53-mediated apoptosis in normal thymocytes and myeloid leukemic cells. Has been shown to inhibit NF-kB, possibly through inhibition of JNK ( $IC_{50}$ =10µM), the COP9 signalosome kinase ( $IC_{50}$ =10µM) and p300/CBP.

Citations: 39

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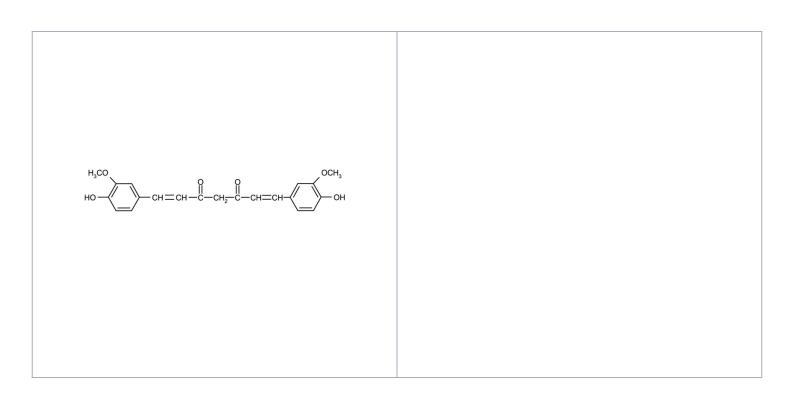
## **Ordering Information**

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ALX-350-028-M010	10mg
ALX-350-028-M050	50mg
ALX-350-028-M250	250mg

Manuals, SDS & CofA

**View Online** »



### **Handling & Storage**

**Use/Stability** As indicated on product label or CoA when stored as recommended.

**Handling** Protect from light.

Long Term Storage Ambient

**Shipping** Ambient Temperature

#### Regulatory Status RUO - Research Use Only

#### **Product Details**

**Alternative Name** 1,7-bis(4-Hydroxy-3-methoxyphenyl)-1,6-heptadiene-3,5-dione, Diferuloylmethane

**Appearance** Orange-yellow crystalline powder.

**CAS** 458-37-7

Couple Target COX, GST, Lipoxygenase, NF-kappaB, NQO

Couple Type Inhibitor

Formula  $C_{21}H_{20}O_6$ 

MI 14: 2673

**MW** 368.4

Purity ≥98% (HPLC)

**Solubility** Soluble in acetic acid, 100% ethanol (at about 1mg/ml) or DMSO (25mg/ml).

**Source** Isolated from turmeric (*Curcuma longa*).



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