## 4'-Bromoflavone

BIOLOGICAL ACTIVITY	
Description	4'-Bromoflavone, a cancer chemopreventive agent, is a potent phase II detoxification enzymes inducer <sup>[1]</sup> .
In Vitro	<ul> <li>4'-Bromoflavone significantly induces quinone reductase (QR) activity in cultured murine hepatoma 1c1c7 cells (concentration to double activity: 10 nM) and effectively induces the α- and mu-isoforms of glutathione S-transferase in cultured H4IIE rat hepatoma cells with no observed toxicity<sup>[1]</sup>.</li> <li>4'-Bromoflavone is a potent inhibitor of cytochrome P4501A1-mediated ethoxyresorufin-O-deethylase activity, with an IC<sub>50</sub> of 0.86 μM. In HepG2 or MCF-7 cells, 4'-Bromoflavone significantly reduces the covalent binding of metabolically activated benzo[a]pyrene to cellular DNA<sup>[1]</sup>.</li> <li>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</li> </ul>
In Vivo	In short-term dietary studies, 4'-Bromoflavone is also shown to increase QR activity and glutathione levels in rat liver, mammary gland, colon, stomach, and lung in a dose-dependent manner. In studies conducted with female Sprague Dawley rats ,4'-Bromoflavone significantly increases QR activity (phase II enzyme) <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

## REFERENCES

[1]. L L Song, et al. Cancer chemopreventive activity mediated by 4'-bromoflavone, a potent inducer of phase II detoxification enzymes. Cancer Res. 1999 Feb 1;59(3):578-85.

Caution: Product has not been fully validated for medical applications. For research use only.

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## Product Data Sheet

