PRODUCT INFORMATION



7-Methoxyisoflavone

Item No. 33419

CAS Registry No.: 1621-56-3

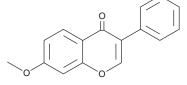
Formal Name: 7-methoxy-3-phenyl-4H-1-benzopyran-4-one

Synonym: 5-methyl-7-Methoxyisoflavone

MF: $C_{16}H_{12}O_3$ FW: 252.3 **Purity:** ≥98% λ_{max} : 250 nm A crystalline solid UV/Vis.: Supplied as:

Storage: -20°C Stability: ≥4 years

Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.



Laboratory Procedures

7-Methoxyisoflavone is supplied as a crystalline solid. A stock solution may be made by dissolving the 7-methoxyisoflavone in the solvent of choice, which should be purged with an inert gas. 7-Methoxyisoflavone is soluble in organic solvents such as DMSO and dimethyl formamide. The solubility of 7-methoxyisoflavone in these solvents is approximately 10 and 30 mg/ml, respectively.

Description

7-Methoxyisoflavone is a synthetic isoflavone and an isomer of 7-methoxyflavone (Item No. 33497). 12 It inhibits mitosis induced by concanavalin A (Item No. 14951) in human peripheral blood mononuclear cells (PBMCs) with an IC $_{50}$ value of 1.36 $\mu g/ml.^{1}$ 7-Methoxyisoflavone (10, 20, and 40 μM) inhibits the growth of HCT116 colon cancer cells in a concentration-dependent manner.²

References

- 1. Hirano, T., Oka, K., Kawashima, E., et al. Effects of synthetic and naturally occurring flavonoids on mitogen-induced proliferation of human peripheral-blood lymphocytes. Life Sci. 45(15), 1407-1411 (1989).
- 2. Hyun, J., Shin, S.Y., So, K.M., et al. Isoflavones inhibit the clonogenicity of human colon cancer cells. Bioorg. Med. Chem. Lett. 22(8), 2664-2669 (2012).

WARNING
THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

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