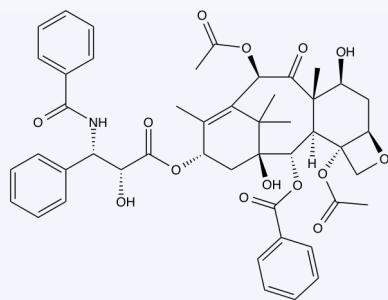


# Cytoskeleton

## Tools for modifying actin or microtubule polymerization



Taxol

Many other cytoskeleton modulators are available. Get lower prices and next day delivery on:

Cytochalasins

Colchicine

Vinblastine sulfate

And more. . .

### References

1. Rowinsky and Donehower (1995), N. Engl. J. Med. **332** 1004
2. Fabbri *et al.* (2008), J. Cell Physiol, **217** 494
3. Spector *et al.* (1989), Cell Motil. Cytoskeleton, **13** 127
4. Suwanborirux *et al.* (1990) Experientia **46** 117
5. Goodin *et al.* (2004), J. Clin. Oncol., **22** 2015
6. Park *et al.* (2012) ChemMedChem **7** 53

### Taxol

Chemotherapeutic agent for the treatment of breast, non-small cell lung and ovarian cancer<sup>1</sup>. Acts as a promoter of tubulin polymerization and stabilizes microtubules in vitro and in vivo resulting in arrest of cells in the G2 and M phase of the cell cycle.

10-2095

5 mg , 25mg

### Docetaxel

Antimitotic chemotherapeutic which inhibits via reversible high-affinity binding to microtubules<sup>2</sup>. Induces apoptosis in a variety of cancer cell lines. Can act in synergy with a other anticancer agents including kinase inhibitors.

10-2286

5 mg , 25mg

### Latrunculins

Inhibit actin polymerization and disrupt microfilament organization. Significantly more potent than cytochalasins in the disruption of microfilament mediated processes<sup>3</sup>.

Latrunculin A

10-2254

100 µg , 1mg

Latrunculin B

10-4303

100 µg , 1mg

### Ansamitocin P-3

Maytansinoid with potent cytotoxic activity<sup>4</sup>. Binds to the rhizoxin/phomopsin binding site on tubulin causing microtubule disassembly and preventing tubulin spiralization<sup>4</sup>.

10-2199

1 mg , 5mg

### Epothilone B

Stabilizes microtubules and promotes tubulin polymerization inducing G<sub>2</sub>-M cell cycle arrest<sup>5</sup>. Displays potent cytotoxic activity in a variety of cell lines and mouse models.

10-2133

1 mg , 5 mg

### Nocodazole

A microtubule polymerization inhibitor that is widely used to induce mitotic arrest and cell synchronization. Recently has been shown to inhibit a number of cancer-related kinases including ABL, c-Kit, BRAF, MEK1, MEK2, and MET<sup>6</sup>.

10-2387

10 mg , 50 mg