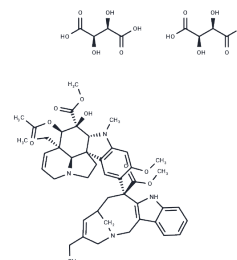


Vinorelbine ditartrate

Chemical Properties

| | |
|-------------------|--|
| CAS No. : | 125317-39-7 |
| Formula: | C ₄₅ H ₅₄ N ₄ O ₈ ·2C ₄ H ₆ O ₆ |
| Molecular Weight: | 1079.11 |
| Appearance: | no data available |
| Storage: | store at low temperature Powder: -20°C for 3 years In solvent: -80°C for 1 year |



Biological Description

| | |
|---------------|---|
| Description | Vinorelbine ditartrate (KW-2307) is a natural alkaloid and an anti-mitotic agent. Vinorelbine ditartrate has anti-tumor activity, inhibiting cell proliferation and inducing apoptosis. |
| Targets(IC50) | Microtubule Associated, Autophagy |
| In vitro | <p>METHODS: Human lung cancer cells NCI-H460 and A549 were treated with Vinorelbine ditartrate (0.01-10 μM) and DT-13 (10 μM) for 48 h. Cell viability was measured by MTT assay.</p> <p>RESULTS: DT-13 significantly increased the cytotoxicity of Vinorelbine on NCI-H460 and A549 cells. The drug-drug interaction between DT-13 and NVB was calculated by combinatorial index (CI) value, which indicated that the DT-13/Vinorelbine combination therapy showed strong synergistic effects in NSCLC cells. [1]</p> <p>METHODS: APC+ or APC- U2OS cells were treated with Vinorelbine ditartrate (1-50 μg/mL) for 4 h. Apoptosis was detected by Flow cytometry.</p> <p>RESULTS: Vinorelbine-induced cell death was more pronounced in APC-deficient cells, suggesting that the number of aCasp3-containing cells increased after 4 h of treatment with a certain concentration of Vinorelbine. This rapid response to Vinorelbine suggests that death is not associated with mitotic arrest. [2]</p> |
| In vivo | <p>METHODS: To assay antitumor activity in vivo, vinorelbine ditartrate (1-10 mg/kg, i.v.) and DT-13 (1.25 mg/kg, by gavage) were administered to BALB/c athymic nude mice bearing NCI-H460 xenografts once daily for three weeks.</p> <p>RESULTS: Treatment with Myricetin or cisplatin alone moderately inhibited tumor growth, but the combination treatment inhibited tumor growth more significantly than Myricetin or cisplatin alone. [1]</p> |

Solubility Information

| | |
|------------|--|
| Solubility | H ₂ O: 107.9 mg/mL (99.99 mM), Sonication is recommended. DMSO: 55 mg/mL (50.97 mM), Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble) |
|------------|--|

Preparing Stock Solutions

| | 1mg | 5mg | 10mg |
|-------|-----------|-----------|-----------|
| 1 mM | 0.9267 mL | 4.6334 mL | 9.2669 mL |
| 5 mM | 0.1853 mL | 0.9267 mL | 1.8534 mL |
| 10 mM | 0.0927 mL | 0.4633 mL | 0.9267 mL |
| 50 mM | 0.0185 mL | 0.0927 mL | 0.1853 mL |

Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Reference

- Li H, et al. DT-13 synergistically enhanced vinorelbine-mediated mitotic arrest through inhibition of FOXM1-BICD2 axis in non-small-cell lung cancer cells. *Cell Death Dis.* 2017 May 25;8(5):e2810.
- Qian H Y, Zhou F, Wu R, et al. Metformin Attenuates Bone Cancer Pain by Reducing TRPV1 and ASIC3 Expression. *Frontiers in Pharmacology.* 2021: 1924.
- Klotz DM, et al. The microtubule poison vinorelbine kills cells independently of mitotic arrest and targets cells lacking the APC tumour suppressor more effectively. *J Cell Sci.* 2012 Feb 15;125(Pt 4):887-95.
- Zhou YT, et al. *Asian Pac J Cancer Prev.* 2013, 14(8), 4635-4639.
- Xu YC, et al. *Breast J.* 2013, 19(2), 180-188.
- Hill BT, et al. *Eur J Cancer.* 1999, 35(3), 512-520.

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