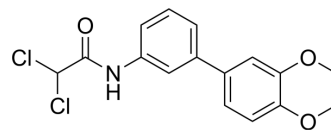


CCTA-1523

| | |
|--------------------|---|
| Cat. No.: | HY-129301 |
| CAS No.: | 1616271-41-0 |
| Molecular Formula: | C ₁₆ H ₁₅ Cl ₂ NO ₃ |
| Molecular Weight: | 340.2 |
| Target: | BCRP |
| Pathway: | Membrane Transporter/Ion Channel |
| Storage: | Please store the product under the recommended conditions in the Certificate of Analysis. |



BIOLOGICAL ACTIVITY

| | | | | | | | | | |
|--------------------|---|---------------|---|---------|--|-----------------|--------------|---------|--|
| Description | CCTA-1523 is a potent, selective, reversible and orally active ABCG2 inhibitor. CCTA-1523 shows cytotoxicity. CCTA-1523 shows anticancer activity ^[1] . | | | | | | | | |
| In Vitro | CCTA-1523 (0-1000 μM; 72 h) shows cytotoxicity for H460, H460/MX20, HEK293/pcDNA3.1, HEK/R482, HEK/R482G and HEK/R482T cells ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. | | | | | | | | |
| In Vivo | CCTA-1523 (75 mg/kg; p.o.; q3d x 6) combined with doxorubicin (HY-15142A) (1.8 mg/kg, i.p., q3d X 6) inhibits tumor growth and increases the survival time in mice ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. | | | | | | | | |
| | <table border="1"> <tr> <td>Animal Model:</td> <td>NCR nude mice (H460/MX20 xenografts)^[1]</td> </tr> <tr> <td>Dosage:</td> <td>75 mg/kg (DOX ;1.8 mg/kg, i.p., q3d X 6)</td> </tr> <tr> <td>Administration:</td> <td>P.o.;q3d x 6</td> </tr> <tr> <td>Result:</td> <td>Significantly inhibited the growth of H460/MX20 tumor xenografts when combined with doxorubicin compared to the vehicle control group.</td> </tr> </table> | Animal Model: | NCR nude mice (H460/MX20 xenografts) ^[1] | Dosage: | 75 mg/kg (DOX ;1.8 mg/kg, i.p., q3d X 6) | Administration: | P.o.;q3d x 6 | Result: | Significantly inhibited the growth of H460/MX20 tumor xenografts when combined with doxorubicin compared to the vehicle control group. |
| Animal Model: | NCR nude mice (H460/MX20 xenografts) ^[1] | | | | | | | | |
| Dosage: | 75 mg/kg (DOX ;1.8 mg/kg, i.p., q3d X 6) | | | | | | | | |
| Administration: | P.o.;q3d x 6 | | | | | | | | |
| Result: | Significantly inhibited the growth of H460/MX20 tumor xenografts when combined with doxorubicin compared to the vehicle control group. | | | | | | | | |

REFERENCES

[1]. Patel A, et al. Suppression of ABCG2 mediated MDR in vitro and in vivo by a novel inhibitor of ABCG2 drug transport. Pharmacol Res. 2017 Jul;121:184-193.

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

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