Roslin 2 bromide

MedChemExpress

| Cat. No.: CAS No.: Molecular Formula: Molecular Weight: Target: | HY-A0280 29574-21-8 C ₁₃ H ₁₉ BrN ₄ 311.22 MDM-2/p53; FAK | |
|---|--|--|
| Molecular Weight: Target: Pathway: | 311.22 MDM-2/p53; FAK Apoptosis; Protein Tyrosine Kinase/RTK | |
| Storage: | -20°C, sealed storage, away from moisture * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture) | |

| BIOLOGICAL ACTIVITY | | |
|---------------------|---|--|
| Description | Roslin 2 bromide (Benzylhexamethylenetetramine bromide) is a p53 reactivator with anticancer effects. Roslin 2 bromide binds FAK, disrupts the binding of FAK and p53 ^[1] . | |
| In Vitro | Roslin 2 bromide decreases cancer cell viability and clonogenicity in a p53-dependent manner. Roslin 2 bromide increases p53 transcriptional activity that is inhibited by FAK using p21, Mdm-2, and Bax-promoter targets. Roslin 2 bromide also causes increased expression of p53 targets: p21, Mdm-2 and Bax proteins ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. | |
| In Vivo | Roslin 2 (60 mg/kg; i.p; 5 days/week) bromide significantly decreases tumor growth, disrupts the complex of FAK and p53, and up-regulates p21 in HCT116 p53 ^{+/+} but not in HCT116 p53 ^{-/-} xenografts in vivo ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. | |

REFERENCES

[1]. Vita M Golubovskaya, et al. Disruption of focal adhesion kinase and p53 interaction with small molecule compound R2 reactivated p53 and blocked tumor growth. BMC Cancer. 2013 Jul 11;13:342.

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

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Product Data Sheet