

Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

Weitere Information auf den folgenden Seiten! See the following pages for more information!



Lieferung & Zahlungsart

siehe unsere Liefer- und Versandbedingungen

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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Screening Libraries

Y06036

Cat. No.: HY-111502 CAS No.: 1832671-96-1 Molecular Formula: $C_{16}H_{15}BrN_{2}O_{5}S$

Molecular Weight: 427.27

Target: **Epigenetic Reader Domain**

Pathway: **Epigenetics**

Storage: Powder -20°C 3 years

2 years

In solvent -80°C 2 years

> -20°C 1 year

Product Data Sheet

SOLVENT & SOLUBILITY

In Vitro

DMSO: 130 mg/mL (304.26 mM; Need ultrasonic)

	Solvent Mass Concentration	1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM	2.3404 mL	11.7022 mL	23.4044 mL
	5 mM	0.4681 mL	2.3404 mL	4.6809 mL
	10 mM	0.2340 mL	1.1702 mL	2.3404 mL

Please refer to the solubility information to select the appropriate solvent.

In Vivo

- 1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.17 mg/mL (5.08 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.17 mg/mL (5.08 mM); Clear solution

of both full-length (AR-fl) and AR variants levels^[1].

BIOLOGICAL ACTIVITY

Description	Y06036 is a potent and selective BET inhibitor, which binds to the BRD4(1) bromodomain with K_d value of 82 nM ^[1] . Antitumor activity ^[1] .
IC ₅₀ & Target	BRD4(1) 82 nM (Kd)
In Vitro	Y06036 (0.001-100 nM, 96 hours for LNCaP, C4-2B, and 22Rv1 cells; 144 hours for VCaP cells) exhibits low micromolar or nanomolar potencies (IC ₅₀ : 0.29-2.6 µM) in the four androgen receptor (AR)-positive prostate cancer cell lines LNCaP, C4-2B,

22Rv1, and VCaP. Treatment of 22Rv1 cells with Y06036 (1, 2, 4, 8, and 16 μ M, 48 hours) results in significant down-regulation

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MCE has not independently confirmed the accuracy of these methods. They are for reference only. Cell Viability Assay^[1]

Cell Line:	LNCaP, C4-2B, 22Rv1, and VCaP prostate cancer cells
Concentration:	0.001-100 μΜ
Incubation Time:	96 hours for LNCaP, C4-2B, and 22Rv1; 144 hours for VCaP
Result:	Inhibited LNCaP, C4-2B, 22Rv1, and VCaP cells with IC $_{50}$ s of 1.06, 2.62, 1.50, 0.63 μ M, respectively.

Cell Line:	22Rv1 cells
Concentration:	1,2,4,8, and 16 μM
Incubation Time:	48 hours
Result:	Resulted in significant down-regulation of both AR-fl and AR variants levels

In Vivo

Y06036 (50 mg/kg, i.p. injection, 5 times per week, 25 days) demonstrates therapeutic effects in a C4-2B CRPC xenograft tumor model in mice. Y06036 is well tolerated in the treated mice, based on the weight of the animal body and their general behavior^[1].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	Four-week-old male mice (strain: C.B-17/IcrHsd-Prkdc for C4-2B) with C4-2B mouse xenograft model $^{[1]}$
Dosage:	50 mg/kg, 100 μL
Administration:	Intraperitoneal (i.p.) injection, 5 times per week, 25 days
Result:	Exhibited strong antitumor activities during the 25-day treatment period, with a tumor growth inhibition (TGI) of 70%.

REFERENCES

[1]. Zhang M, et al. Structure-Based Discovery and Optimization of Benzo[d]isoxazole Derivatives as Potent and Selective BET Inhibitors for Potential Treatment of Castration-Resistant Prostate Cancer (CRPC). J Med Chem. 2018 Apr 12;61(7):3037-3058.

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

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