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产品名称: **AMG-510; Sotorasib**

CAS 号: **2296729-00-3**

产品货号: **S89986**

生物活性:

Description	Sotorasib (AMG-510) is a first-in-class, orally bioavailable, and selective KRAS G12C covalent inhibitor. Sotorasib irreversibly inhibits KRAS G12C by locking it in an inactive GDP-bound state. Sotorasib leads to the regression of KRAS G12C-mutated locally advanced or metastatic non-small cell lung cancer (NSCLC)[1][2][3][4][5][6].																								
IC50 & Target	KRAS(G12C)																								
In Vitro	在细胞测定中, Sotorasib (AMG-510) 共价修饰 KRAS G12C 并抑制 KRAS G12C 信号传导, 通过所有 KRAS p.G12C 突变细胞系中的 ERK1/2 (p-ERK) 磷酸化来测量[2]。 Sotorasib (AMG-510; 1-10 μM; 72 小时) 也有效损害 NCI-H358 和 MIA PaCa-2 中的细胞活力, IC50 分别为 0.006 μM 和 0.009 μM。非 KRASG12C 细胞系对 Sotorasib (IC50>7.5 μM) 不敏感[3]。 Sotorasib (AMG-510) (0-50 μM, 72 小时) 可抑制细胞生长, 并与 Cisplatin 在 H23 和 H358 细胞中产生协同作用[5]。 Sotorasib (AMG-510) (100 nM, 4-72 小时) 对活性 KRAS 具有持续抑制作用, 但仍能观察到 RAS-MAPK 的自适应反馈再激活[6]。 Cell Viability Assay[3] <table><tr><td>Cell Line:</td><td>NCI-H358 and MIA PaCa-2 cells</td></tr><tr><td>Concentration:</td><td>1-10 μM</td></tr><tr><td>Incubation Time:</td><td>72 hours</td></tr><tr><td>Result:</td><td>Potently impaired cellular viability in both NCI-H358 and MIA PaCa-2 (IC50=$\sim$0.006 μM and $\sim$0.009 μM respectively).</td></tr></table> Cell Viability Assay[5] <table><tr><td>Cell Line:</td><td>NCI-H23 (H23) and NCI-H358 (H358) cells (Lung adenocarcinoma cell lines harbor KRAS G12C mutations)</td></tr><tr><td>Concentration:</td><td>0-50 μM</td></tr><tr><td>Incubation Time:</td><td>72 h</td></tr><tr><td>Result:</td><td>The IC50 values in H358 and H23 cell lines were 0.0818 μM, 0.6904 μM, respectively. Had a synergistic effect with Cisplatin (HY-17394).</td></tr></table> Western Blot Analysis[6] <table><tr><td>Cell Line:</td><td>H358, MGH1088-1, MGH1062, MIA PaCa-2, B8182, LIM2099, SW837, SW1463 (KRAS_{G12C} mutant cell lines)</td></tr><tr><td>Concentration:</td><td>100 nM</td></tr><tr><td>Incubation Time:</td><td>4, 24, 48, 72 h</td></tr><tr><td>Result:</td><td>Showed effective inhibition of the MAPK pathway at 4 h, comparable rapid and robust reactivation of RAS-MAPK signaling was observed, such that phospho-ERK levels returned to an average of $\sim$75% of baseline levels by just 72 h. Adaptive feedback is likely to be a key driver of resistance.</td></tr></table>	Cell Line:	NCI-H358 and MIA PaCa-2 cells	Concentration:	1-10 μ M	Incubation Time:	72 hours	Result:	Potently impaired cellular viability in both NCI-H358 and MIA PaCa-2 (IC50= $\sim$ 0.006 μ M and $\sim$ 0.009 μ M respectively).	Cell Line:	NCI-H23 (H23) and NCI-H358 (H358) cells (Lung adenocarcinoma cell lines harbor KRAS G12C mutations)	Concentration:	0-50 μ M	Incubation Time:	72 h	Result:	The IC50 values in H358 and H23 cell lines were 0.0818 μ M, 0.6904 μ M, respectively. Had a synergistic effect with Cisplatin (HY-17394).	Cell Line:	H358, MGH1088-1, MGH1062, MIA PaCa-2, B8182, LIM2099, SW837, SW1463 (KRAS _{G12C} mutant cell lines)	Concentration:	100 nM	Incubation Time:	4, 24, 48, 72 h	Result:	Showed effective inhibition of the MAPK pathway at 4 h, comparable rapid and robust reactivation of RAS-MAPK signaling was observed, such that phospho-ERK levels returned to an average of $\sim$ 75% of baseline levels by just 72 h. Adaptive feedback is likely to be a key driver of resistance.
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In Vivo	在临床前肿瘤模型中, Sotorasib (AMG-510) 快速且不可逆地结合 KRAS G12C, 持久抑制丝裂原活化																								



	蛋白激酶 (MAPK) 信号通路。Sotorasib (口服; 每天一次) 能够在 KRAS G12C 癌症小鼠模型中诱导肿瘤消退[3]。 Sotorasib (AMG-510) (30 mg/kg; 口服; 每日一次, 共持续 28 天) 可减小 NCI-H358 细胞衍生异种移植小鼠模型中的肿瘤大小, 并与 Cisplatin 具有协同作用[5]。 <table><tr><td>Animal Model:</td><td>Balb/C nude mice (injected with H358 cells under the skin)[5]</td></tr><tr><td>Dosage:</td><td>30 mg/kg</td></tr><tr><td>Administration:</td><td>Oral gavage (p.o.); daily for 28 days</td></tr><tr><td>Result:</td><td>The mean tumor volume were 426.66mm³. Reduced tumor size, and had a synergistic effect with Cisplatin. Had little effects on body weight.</td></tr></table>	Animal Model:	Balb/C nude mice (injected with H358 cells under the skin)[5]	Dosage:	30 mg/kg	Administration:	Oral gavage (p.o.); daily for 28 days	Result:	The mean tumor volume were 426.66mm ³ . Reduced tumor size, and had a synergistic effect with Cisplatin. Had little effects on body weight.													
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Solvent&Solubility	细胞实验: DMSO 中的溶解度 : 50 mg/mL (89.19 mM; 超声助溶; 吸湿的 DMSO 对产品的溶解度有显著影响, 请使用新开封的 DMSO) H2O 中的溶解度 : 33.33 mg/mL (59.46 mM; ultrasonic and adjust pH to 11 with NaOH) <table><tr><th rowspan="4">Preparing Stock Solutions</th><th>Solvent Concentration</th><th>Mass</th><th>1 mg</th><th>5 mg</th><th>10 mg</th></tr><tr><td>1 mM</td><td></td><td>1.7838 mL</td><td>8.9192 mL</td><td>17.8383 mL</td></tr><tr><td>5 mM</td><td></td><td>0.3568 mL</td><td>1.7838 mL</td><td>3.5677 mL</td></tr><tr><td>10 mM</td><td></td><td>0.1784 mL</td><td>0.8919 mL</td><td>1.7838 mL</td></tr></table> * 请根据产品在不同溶剂中的溶解度选择合适的溶剂配制储备液; 一旦配成溶液, 请分装保存, 避免反复冻融造成的产品失效。 储备液的保存方式和期限: -80°C, 6 months; -20°C, 1 month (stored under nitrogen)。-80°C 储存时, 请在 6 个月内使用, -20°C 储存时, 请在 1 个月内使用。 动物实验: 请根据您的 实验动物和给药方式 选择适当的溶解方案。 以下溶解方案都请先按照 In Vitro 方式配制澄清的储备液, 再依次添加助溶剂: ——为保证实验结果的可靠性, 澄清的储备液可以根据储存条件, 适当保存; 体内实验的工作液, 建议您现用现配, 当天使用; 以下溶剂前显示的百分比是指该溶剂在您配制终溶液中的体积占比; 如在配制过程中出现沉淀、析出现象, 可以通过加热和/或超声的方式助溶 方案 一 请依序添加每种溶剂: 10% DMSO → 40% PEG300 → 5% Tween-80 → 45% Saline Solubility: ≥ 2.08 mg/mL (3.71 mM); 澄清溶液 此方案可获得 ≥ 2.08 mg/mL (饱和度未知) 的澄清溶液。 以 1 mL 工作液为例, 取 100 μL 20.8 mg/mL 的澄清 DMSO 储备液加到 400 μL PEG300 中, 混合均匀; 再向上述体系中加入 50 μL Tween-80, 混合均匀; 然后再继续加入 450 μL 生理盐水 定容至 1 mL。 生理盐水的配制: 将 0.9 g 氯化钠, 溶解于 ddH₂ O 并定容至 100 mL, 可以得到澄清透明的生理盐水溶液。 方案 二 请依序添加每种溶剂: 10% DMSO → 90% (20% SBE-β-CD in Saline) Solubility: ≥ 2.08 mg/mL (3.71 mM); 澄清溶液 此方案可获得 ≥ 2.08 mg/mL (饱和度未知) 的澄清溶液。	Preparing Stock Solutions	Solvent Concentration	Mass	1 mg	5 mg	10 mg	1 mM		1.7838 mL	8.9192 mL	17.8383 mL	5 mM		0.3568 mL	1.7838 mL	3.5677 mL	10 mM		0.1784 mL	0.8919 mL	1.7838 mL
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	以 1 mL 工作液为例, 取 100 μL 20.8 mg/mL 的澄清 DMSO 储备液加到 900 μL 20% 的 SBE-β-CD 生理盐水水溶液 中, 混合均匀。 2 g SBE-β-CD (磺丁基醚 β-环糊精) 粉末定容于 10 mL 的生理盐水中, 完全溶解至澄清透明。 方案 三 请依序添加每种溶剂: 10% DMSO → 90% Corn Oil Solubility: ≥ 2.08 mg/mL (3.71 mM); 澄清溶液 此方案可获得 ≥ 2.08 mg/mL (饱和度未知) 的澄清溶液, 此方案实验周期在半个月以上的动物实验酌情使用。 以 1 mL 工作液为例, 取 100 μL 20.8 mg/mL 的澄清 DMSO 储备液加到 900 μL 玉米油中, 混合均匀。 以下溶解方案, 请直接配制工作液。建议现用现配, 在短期内尽快用完。 以下溶剂前显示的百分比是指该溶剂在您配制终溶液中的体积占比; 如在配制过程中出现沉淀、析出现象, 可以通过加热和/或超声的方式助溶。 方案 一 请依序添加每种溶剂: 20% HP-β-CD in Saline Solubility: 10 mg/mL (17.84 mM); 悬浊液; 超声助溶				
参考文献	[1]. Marwan Fakih, et al, Phase 1 study evaluating the safety, tolerability, pharmacokinetics (PK), and efficacy of AMG 510, a novel small molecule KRASG12Cinhibitor, in advanced solid tumors. Journal of Clinical Oncology. [2]. Karen Rex, et al. Abstract 3090: In vivo characterization of AMG 510 - a potent and selective KRASG12Ccovalent small molecule inhibitor in preclinical KRASG12C cancer models. Experimental and Molecular Therapeutics. [3]. Canon J, et al. The clinical KRAS(G12C) inhibitor AMG 510 drives anti-tumour immunity. Nature. 2019 Nov;575(7781):217-223. [4]. Brian A. Lanman, et al.Abstract 4455: Discovery of AMG 510, a first-in-human covalent inhibitor of KRASG12C for the treatment of solid tumors. Cancer Chemistry. [5]. Wu LL, et al. AMG-510 and cisplatin combination increases antitumor effect in lung adenocarcinoma with mutation of KRAS G12C: a preclinical and translational research. Discov Oncol. 2023 Jun 7;14(1):91. [6]. Ryan MB, et al. KRASG12C-independent feedback activation of wild-type RAS constrains KRASG12C inhibitor efficacy. Cell Rep. 2022 Jun 21;39(12):110993.				
完整储备液配制表: 请根据产品在不同溶剂中的溶解度选择合适的溶剂配制储备液; 一旦配成溶液, 请分装保存, 避免反复冻融造成的产品失效。 储备液的保存方式和期限: -80℃, 6 months; -20℃, 1 month (stored under nitrogen)。-80℃ 储存时, 请在 6 个月内使用, -20℃ 储存时, 请在 1 个月内使用。					
可选溶剂	质量 溶剂体积 浓度	1 mg	5 mg	10 mg	25 mg
H2O / DMSO	1 mM	1.7838 mL	8.9192 mL	17.8383 mL	44.5959 mL
	5 mM	0.3568 mL	1.7838 mL	3.5677 mL	8.9192 mL
	10 mM	0.1784 mL	0.8919 mL	1.7838 mL	4.4596 mL



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	15 mM	0.1189 mL	0.5946 mL	1.1892 mL	2.9731 mL
	20 mM	0.0892 mL	0.4460 mL	0.8919 mL	2.2298 mL
	25 mM	0.0714 mL	0.3568 mL	0.7135 mL	1.7838 mL
	30 mM	0.0595 mL	0.2973 mL	0.5946 mL	1.4865 mL
	40 mM	0.0446 mL	0.2230 mL	0.4460 mL	1.1149 mL
	50 mM	0.0357 mL	0.1784 mL	0.3568 mL	0.8919 mL
DMSO	60 mM	0.0297 mL	0.1487 mL	0.2973 mL	0.7433 mL
	80 mM	0.0223 mL	0.1115 mL	0.2230 mL	0.5574 mL

备注：如您选择水作为储备液，请稀释至工作液后，再用 0.22 μ m 的滤膜过滤除菌后使用。



源叶