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产品名称: **Mutated EGFR-IN-1**

产品别名: **Osimertinib analog**

生物活性:

| Description | Mutated EGFR-IN-1 (Osimertinib analog) is a useful intermediate for the inhibitors design for mutated EGFR, such as L858R EGFR, Exon19 deletion activating mutant and T790M resistance mutant. IC50 value: Target: Mutated EGFR inhibitor More information can be found in Patent WO 2013014448 A1.2 - (2, 4, 5 - substituted -anilino) pyrimidine derivatives as egfr modulators useful for treating cancer. | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
|-------------------------------------|---|------|--|-----------------------|---------------------------|---------|------|---------------|--|--|------|--|------|-------|------|-----------|--|------------|------------|------|-----------|--|-----------|-----------|-------|-----------|--|-----------|-----------|
| IC₅₀ & Target | EGFR ^{L858R} | | EGFR ^{Exon 19 deletion/T790M} | EGFR ^{T790M} | | | | | | | | | | | | | | | | | | | | | | | | | |
| Solvent&Solubility | <p>In Vitro:</p> <p>DMSO : ≥ 75 mg/mL (168.33 mM)</p> <p>* "≥" means soluble, but saturation unknown.</p> <table border="1"><thead><tr><th rowspan="2">Preparing Stock Solutions</th><th>Solvent</th><th>Mass</th><th>Concentration</th><th></th></tr><tr><th></th><th>1 mg</th><th></th><th>5 mg</th><th>10 mg</th></tr></thead><tbody><tr><td>1 mM</td><td>2.2444 mL</td><td></td><td>11.2218 mL</td><td>22.4437 mL</td></tr><tr><td>5 mM</td><td>0.4489 mL</td><td></td><td>2.2444 mL</td><td>4.4887 mL</td></tr><tr><td>10 mM</td><td>0.2244 mL</td><td></td><td>1.1222 mL</td><td>2.2444 mL</td></tr></tbody></table> <p>*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。</p> <p>储备液的保存方式和期限 -80°C, 6 months; -20°C, 1 month。 -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。</p> | | | | Preparing Stock Solutions | Solvent | Mass | Concentration | | | 1 mg | | 5 mg | 10 mg | 1 mM | 2.2444 mL | | 11.2218 mL | 22.4437 mL | 5 mM | 0.4489 mL | | 2.2444 mL | 4.4887 mL | 10 mM | 0.2244 mL | | 1.1222 mL | 2.2444 mL |
| Preparing Stock Solutions | Solvent | Mass | Concentration | | | | | | | | | | | | | | | | | | | | | | | | | | |
| | | 1 mg | | 5 mg | 10 mg | | | | | | | | | | | | | | | | | | | | | | | | |
| 1 mM | 2.2444 mL | | 11.2218 mL | 22.4437 mL | | | | | | | | | | | | | | | | | | | | | | | | | |
| 5 mM | 0.4489 mL | | 2.2444 mL | 4.4887 mL | | | | | | | | | | | | | | | | | | | | | | | | | |
| 10 mM | 0.2244 mL | | 1.1222 mL | 2.2444 mL | | | | | | | | | | | | | | | | | | | | | | | | | |
| References | [1]. Patent WO 2013014448 A1. 2 - (2, 4, 5 - substituted -anilino) pyrimidine derivatives as egfr modulators useful for treating cancer . | | | | | | | | | | | | | | | | | | | | | | | | | | | | |

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