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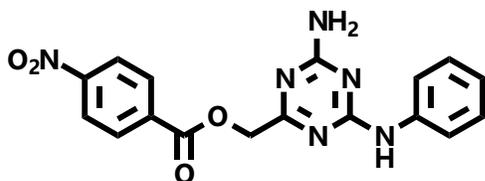
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Rad6 Ubiquitin Inhibitor – TZ9

Chemical Name: (4-amino-6-(phenylamino)-1,3,5-triazin-2-yl)methyl 4-nitrobenzoate



| | |
|-------------------|---|
| Molecular Weight: | 366.34 |
| Formula: | C ₁₇ H ₁₄ N ₆ O ₄ |
| Purity: | ≥98% |
| CAS#: | 1002789-86-7 |
| Solubility: | DMSO up to 50 mM |
| Storage | Powder: 4°C 1 year DMSO: 4°C 3 month -20°C 1 year |

Biological Activity:

TZ9 is a selective and cell permeable inhibitor of human E2 ubiquitin-conjugating enzyme Rad6B. It binds to the Rad6B catalytic site, inhibits Rad6B-induced histone H2A ubiquitination, downregulates intracellular β -catenin. It can induce G2/M arrest and apoptosis, and inhibit proliferation and migration of metastatic human breast cancer cells (IC_{50} ~6 μ M).

How to Use:

In vitro: TZ9 was used at 10-50 μ M final concentration in vitro and in cellular assays.

In vivo: n/a

Reference:

1. Sanders MA, et al. Novel inhibitors of Rad6 ubiquitin conjugating enzyme: design, synthesis, identification, and functional characterization. (2013) Mol Cancer Ther. 12(4):373-83.

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