

## Small Molecules

### Pifithrin-mu

p53 inhibitor

Catalog # 72802  
72804

10 mg  
50 mg



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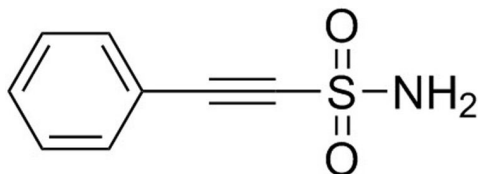
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## Product Description

Pifithrin-mu (PFT- $\mu$ ) is an inhibitor of p53-mediated apoptosis, preventing p53 binding to Bcl-xL and Bcl-2 at the mitochondrial surface, without affecting p53 transactivational activities (Strom et al.). In vitro, PFT- $\mu$  binds both p53 ( $K_d = 0.82$  mM) and Bcl-xL ( $K_d = 0.80$  mM; Hagn et al.). PFT- $\mu$  also interacts selectively with heat shock protein 70 (HSP70), leading to disruption of the association between HSP70 and many of its co-chaperones and substrate proteins (Leu et al.).

Molecular Name:	Pifithrin-mu
Alternative Names:	2-Phenylethynylsulfonamide; PFT- $\mu$ ; Pifithrin- $\mu$
CAS Number:	64984-31-2
Chemical Formula:	C <sub>8</sub> H <sub>7</sub> NO <sub>2</sub> S
Molecular Weight:	181.2 g/mol
Purity:	≥ 98%
Chemical Name:	Ethynylsulfonamide, 2-phenyl-
Structure:	



## Properties

Physical Appearance:	A crystalline solid
Storage:	Product stable at -20°C as supplied. Protect from prolonged exposure to light. Stable as supplied for 12 months from date of receipt.
Solubility:	· Absolute ethanol ≤ 75 mM · DMSO ≤ 75 mM For example, to prepare a 10 mM stock solution in DMSO, resuspend 1 mg in 552 $\mu$ L of fresh DMSO.

Prepare stock solution fresh before use. Information regarding stability of small molecules in solution has rarely been reported, however, as a general guide we recommend storage in DMSO at -20°C. Aliquot into working volumes to avoid repeated freeze-thaw cycles. The effect of storage of stock solution on compound performance should be tested for each application.

Compound has low solubility in aqueous media. For use as a cell culture supplement, stock solution should be diluted into culture medium immediately before use. Avoid final DMSO concentration above 0.1% due to potential cell toxicity.

## Published Applications

### MAINTENANCE AND SELF-RENEWAL

- In combination with Rho-associated coiled-coil containing protein kinase (ROCK) inhibitor Y-27632 (Catalog #72302), improves cell recovery after cryopreservation (Xu et al.).
- Inhibits DNA damage-induced apoptosis in human embryonic stem cells (Qin et al.).

## References

- Hahn F et al. (2010) BclxL changes conformation upon binding to wild-type but not mutant p53 DNA binding domain. *J Biol Chem* 285(5): 3439–50.
- Leu JI-J et al. (2009) A small molecule inhibitor of inducible heat shock protein 70. *Mol Cell* 36(1): 15–27.
- Qin H et al. (2007) Regulation of apoptosis and differentiation by p53 in human embryonic stem cells. *J Biol Chem* 282(8): 5842–52.
- Strom E et al. (2006) Small-molecule inhibitor of p53 binding to mitochondria protects mice from gamma radiation. *Nat Chem Biol* 2(9): 474–9.
- Xu X et al. (2010) Enhancement of cell recovery for dissociated human embryonic stem cells after cryopreservation. *Biotechnol Prog* 26(3): 781–8.

## Related Small Molecules

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**This product is hazardous. Please refer to the Safety Data Sheet (SDS).**

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