IWP-4

| Cat. No.: | HY-12879 | | |
|--------------------|---|-----------------------|---------|
| CAS No.: | 686772-17-8 | 8 | |
| Molecular Formula: | C ₂₃ H ₂₀ N ₄ O ₃ S | b ₃ | |
| Molecular Weight: | 496.62 | | |
| Target: | Wnt | | |
| Pathway: | Stem Cell/Wnt | | |
| Storage: | Powder | -20°C | 3 years |
| | | 4°C | 2 years |
| | In solvent | -80°C | 2 years |
| | | -20°C | 1 year |

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SOLVENT & SOLUBILITY

| | Solvent Mass Concentration | 1 mg | 5 mg | 10 mg |
|------------------------------|-------------------------------|-----------|------------|------------|
| Preparing Stock Solution: | 1 mM | 2.0136 mL | 10.0681 mL | 20.1361 mL |
| | 5 mM | | | |
| | 10 mM | | | |

| BIOLOGICAL ACTIVITY | | | | |
|---------------------------|---|--|--|--|
| BIOLOGICAL ACTIVITY | | | | |
| Description | IWP-4 is a small molecule Wnt inhibitor with an IC $_{50}$ of 25 nM. | | | |
| IC ₅₀ & Target | IC50: 25 nM (Wnt) ^[1] | | | |
| In Vitro | IWP-4 is a small molecule Wnt inhibitor with an IC ₅₀ of 25 nM. IWP-4 induces the expression of cardiac markers, including cardiac troponin I (CTNI) and cardiac myosin heavy chain bright cells (MYH ^{hi+}). IWP-4 also results in the appearance of beating foci (0.44±0.10 SEM beats per second), which is absent in all cultures not receiving IWP-4. Further, flow cytometric analysis shows that there are significantly more MYH ^{lo+} cells in IWP-4 treated cultures (P<0.0002) compare with untreated cultures at day 16, being 17.0±1.3 SD% and 5.4±1.4 SD%, respectively. Quantification of NKX2-5 protein expression shows that 63% (481/817) of IWP-4 treated cells display nuclear NKX2-5 expression ^[1] . Mesenchymal precursor cells (MPCs) treated with IWP-4 show no significant changes in the expression of AXIN2, CTNNB1 and GSK3β as compare to osteogenic medium alone on day 7, but MPCs treated with IWP-4 express elevates levels of DKK1 and GSK3β on day 21. IWP-4 also causes a significant down regulation of SPARC and COL1A1 ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. | | | |

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PROTOCOL

Cell Assay^[1]

hESC cultures are obtained in mTeSR-1 medium and expanded with daily medium exchange until colonies reach the desired level of confluence (~70% to 80%). At this time (marked day 0), mTeSR-1 is replaced with a basal medium comprised of RPMI 1640 medium supplemented with 2% B27 supplement and 1% penicillin/streptomycin. 20 ng/mL BMP-4 and/or 6 ng/mL activin A are added to the basal medium for primitive streak induction, and exchanged daily until day 3. Then, basal media with or without 5 mM IWP-4 is added to the cells and exchanged every 2 days [dimethyl sulfoxide (DMSO) at the same concentration is used as a vehicle control] until day 15, after which basal medium is supplied every 2 days^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

CUSTOMER VALIDATION

• Chin Med. 2022 Jan 6;17(1):11.

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REFERENCES

[1]. Hudson J, et al. Primitive cardiac cells from human embryonic stem cells. Stem Cells Dev. 2012 Jun 10;21(9):1513-23.

[2]. Frith JE, et al. Microbioreactor array screening of Wnt modulators and microenvironmental factors in osteogenic differentiation of mesenchymal progenitor cells. PLoS One. 2013 Dec 23;8(12):e82931.

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

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