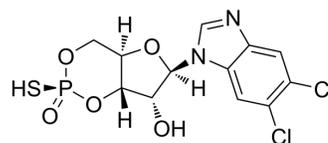


Sp-5,6-DCl-cBIMPS

| | |
|---------------------------|---|
| Cat. No.: | HY-137638 |
| CAS No.: | 120912-54-1 |
| Molecular Formula: | C ₁₂ H ₁₁ Cl ₂ N ₂ O ₅ PS |
| Molecular Weight: | 397.17 |
| Target: | PKA |
| Pathway: | Stem Cell/Wnt |
| Storage: | Please store the product under the recommended conditions in the Certificate of Analysis. |



BIOLOGICAL ACTIVITY

| | | | | | | | | | |
|--------------------|--|------------|---------------------------------|----------------|--------|------------------|--------|---------|---|
| Description | Sp-5,6-DCl-cBIMPS is a potent and specific cAMP-dependent protein kinases (cAMP-PK) activator. Sp-5,6-DCl-cBIMPS stimulates insulin release. Sp-5,6-DCl-cBIMPS inhibits U46619-induced activation of Rho, Gq and G12/G13 in platelets ^{[1][2][3]} . | | | | | | | | |
| In Vitro | <p>Sp-5,6-DCl-cBIMPS (0.005, 0.05, 0.5 mM; 60 min) stimulates insulin release from isolated pancreatic islets in a dose and time-dependent manner^[2].</p> <p>Sp-5,6-DCl-cBIMPS (0.005, 0.05, 0.5 mM; 60 min) increases glucose (8.5 mM)-stimulated insulin release^[2].</p> <p>Sp-5,6-DCl-cBIMPS (100 μM; 20 min) inhibits U46619-induced Rho activation in wild-type mouse platelets, and inhibits U46619-induced activation of both Gq and G12/G13 in human platelets^[3].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Western Blot Analysis^[3]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>Wild-type mice, Human platelets</td> </tr> <tr> <td>Concentration:</td> <td>100 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>20 min</td> </tr> <tr> <td>Result:</td> <td>Strongly suppressed U46619-induced Rho activation in wild-type, and inhibited U46619-induced activation of both Gq and G12/G13.</td> </tr> </table> | Cell Line: | Wild-type mice, Human platelets | Concentration: | 100 μM | Incubation Time: | 20 min | Result: | Strongly suppressed U46619-induced Rho activation in wild-type, and inhibited U46619-induced activation of both Gq and G12/G13. |
| Cell Line: | Wild-type mice, Human platelets | | | | | | | | |
| Concentration: | 100 μM | | | | | | | | |
| Incubation Time: | 20 min | | | | | | | | |
| Result: | Strongly suppressed U46619-induced Rho activation in wild-type, and inhibited U46619-induced activation of both Gq and G12/G13. | | | | | | | | |

REFERENCES

- [1]. Sandberg M, et al. Characterization of Sp-5,6-dichloro-1-beta-D-ribofuranosylbenzimidazole-3',5'-monophosphorothioate (Sp-5,6-DCl-cBiMPS) as a potent and specific activator of cyclic-AMP-dependent protein kinase in cell extracts and intact cells. *Biochem J.* 1991 Oct 15;279 (Pt 2)(Pt 2):521-7.
- [2]. Laychock SG. Sp-5,6-dichloro-1-beta-D-ribofuranosylbenzimidazole-3',5'-cyclic monophosphorothioate is a potent stimulus for insulin release. *Endocr Res.* 1993;19(2-3):113-22.
- [3]. Gratacap MP, et al. Differential regulation of Rho and Rac through heterotrimeric G-proteins and cyclic nucleotides. *J Biol Chem.* 2001 Dec 21;276(51):47906-13.

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

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