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Proteins

Product Data Sheet

3-Deoxyglucosone

Cat. No.: HY-N7426 CAS No.: 4084-27-9 Molecular Formula: $C_{6}H_{10}O_{5}$ **Molecular Weight:** 162.14

Target: **GLP Receptor** Pathway: GPCR/G Protein

Please store the product under the recommended conditions in the Certificate of Storage:

Analysis.

BIOLOGICAL ACTIVITY

Description

3-Deoxyglucosone (3-Deoxy-D-glucosone) is a reactive intermediate of the Maillard reaction and the polyol pathway. 3-Deoxyglucosone rapidly reacts with protein amino groups to form advanced glycation end products (AGEs), such as imidazolone, it is the most specific AGE for 3-DG. 3-Deoxyglucosone synergizes with low glucose to potentiate GLP-1 secretion and is considered as a biomarker for diabetes^{[1][2][3]}.

In Vitro

- 3-Deoxyglucosone (80 ng/ml-1000 ng/ml; 1 hour) markedly increases GLP-1 secretion by 1.23-folds in 300 ng/ml or 1000 ng/ml 3DG-treated group. But at allower concentration (80 ng/ml) has no effects^[1].
- 3-Deoxyglucosone (300 ng/ml; 1 hour) dramatically increases intracellular Ca²⁺ levels by Fluo-3/AM determination (2.5 µM for 30 mins). But 3DG does not affect intracellular cAMP levels in a cAMP Elisa assay^[1].
- 3-Deoxyglucosone (300 ng/ml; 1 hour) significantly increases the protein expression levels of TAS1R2, TAS1R3, and TRPM5 under both glucose-free and highconditions^[1].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Western Blot Analysis^[1]

Cell Line:	STC-1 cells
Concentration:	300 ng/ml
Incubation Time:	1 hour
Result:	Upregulated TAS1R2, TAS1R3, and TRPM5 expression.

In Vivo

- 3-Deoxyglucosone (intragastric administration; 20 mg/kg; single dose) impairs glucose tolerance with increased AUC, but the plasma glucagon levels are not significantly different. It developes impaired glucose regulation (IGR) with obviously pancreatic islet cell dysfunction in kunming mice and SD-rats^[2].
- 3-deoxyglucosone (gastric gavage; 5-50 mg/kg; once daily; 2 weeks) is significantly increased in the upper small intestine (1.4-fold), lower small intestine (1.4-fold), ileum (1.4-fold) and colon (two fold) compared with the basal levels in the corresponding control group. In addition, the protein expressions of TAS1R2, TAS1R3 and TRPM5 in both duodenum and colon are significantly decreased^[3].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model: SD rats ^[3]

Dosage:	5, 20 and 50 mg/kg
Administration:	oral administration; once daily; 2 weeks
Result:	Was capable of accumulating in intestinal tissue and thereby decreased secretion of GLF 1 and insulin.

REFERENCES

- [1]. Song X, et al. 3-Deoxyglucosone Induces Glucagon-Like Peptide-1 Secretion from STC-1 Cells via Upregulating Sweet Taste Receptor Expression under Basal Conditions.Int J Endocrinol. 2019 Oct 23;2019:4959646.
- [2]. Wang F, et al. Acute Reduction of Incretin Effect and Glucose Intolerance in Rats by Single Intragastric Administration of 3-deoxyglucosone. Exp Clin Endocrinol Diabetes. 2017 Jan;125(1):4-11.
- [3]. Zhang L, et al. Accumulation of intestinal tissue 3-deoxyglucosone attenuated GLP-1 secretion and its insulinotropic effect in rats. Diabetol Metab Syndr. 2016 Nov 29;8:78.
- [4]. Niwa T, et al. 3-Deoxyglucosone: metabolism, analysis, biological activity, and clinical implication. J Chromatogr B Biomed Sci Appl. 1999 Aug 6;731(1):23-36.

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

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