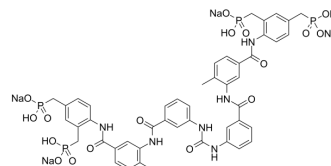


## NF546

<b>Cat. No.:</b>	HY-108661
<b>CAS No.:</b>	1006028-37-0
<b>Molecular Formula:</b>	C <sub>47</sub> H <sub>44</sub> N <sub>6</sub> Na <sub>4</sub> O <sub>17</sub> P <sub>4</sub>
<b>Molecular Weight:</b>	1180.74
<b>Target:</b>	P2Y Receptor
<b>Pathway:</b>	GPCR/G Protein
<b>Storage:</b>	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	NF546 is a selective non-nucleotide P2Y <sub>11</sub> agonist with a pEC <sub>50</sub> of 6.27. NF546 stimulates release of interleukin-8 from human monocyte-derived dendritic cells <sup>[1]</sup> .
<b>IC<sub>50</sub> &amp; Target</b>	pEC <sub>50</sub> : 6.27 (P2Y <sub>11</sub> ) <sup>[1]</sup>
<b>In Vitro</b>	NF546 is relatively selective for P2Y <sub>11</sub> over P2Y <sub>1</sub> , P2Y <sub>2</sub> , P2Y <sub>4</sub> , P2Y <sub>6</sub> , P2Y <sub>12</sub> , P2X <sub>1</sub> , P2X <sub>2</sub> , and P2X <sub>2</sub> -X <sub>3</sub> . NF546 (100 μM; 24 hours) is equi-efficacious in stimulating TSP-1 release and inhibiting the LPS-induced release of IL-12p70 in human monocyte-derived dendritic cells <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

### REFERENCES

[1]. Meis S, et al. NF546 [4,4'-(carbonylbis(imino-3,1-phenylene-carbonylimino-3,1-(4-methyl-phenylene)-carbonylimino))-bis(1,3-xylene-alpha, alpha'-diphosphonic acid) tetrasodium salt] is a non-nucleotide P2Y<sub>11</sub> agonist and stimulates release of interleukin-8 from human monocyte-derived dendritic cells. *J Pharmacol Exp Ther.* 2010 Jan;332(1):238-47.

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

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