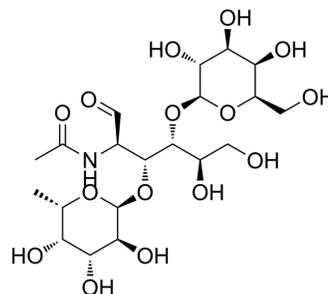


## Lewis X trisaccharide

Cat. No.:	HY-N10534	
CAS No.:	71208-06-5	
Molecular Formula:	C <sub>20</sub> H <sub>35</sub> NO <sub>15</sub>	
Molecular Weight:	529.49	
Target:	Parasite	
Pathway:	Anti-infection	
Storage:	Powder	-20°C 3 years
	In solvent	-80°C 6 months
		-20°C 1 month



### BIOLOGICAL ACTIVITY

<b>Description</b>	<p>Lewis X trisaccharide (Lewis X, Le<sup>x</sup>) is a potent T<sub>H</sub>2 regulator, antagonizes LPS-induced IL-12 immune expression. Lewis X trisaccharide is a human histo-blood group antigen, plays an key role in cell-cell adhesion, and servers as a tumor marker. Lewis X trisaccharide is highly expressed in the outer membrane of the parasite, can be used for the immunology research of schistosomiasis<sup>[1][2][3]</sup>.</p>								
<b>IC<sub>50</sub> &amp; Target</b>	Schistosome								
<b>In Vitro</b>	<p>Lewis X trisaccharide-BSA (25 µg/mL; 30 min; before LPS stimulation of 10 ng/mL for 4 h) IL-12p40 and suppresses IL-12p70 protein expression induced by <a href="#">Lipopolysaccharide</a> (LPS, HY-D1056)<sup>[1]</sup>.</p> <p>Lewis X trisaccharide (2 µM; 30 min; before LPS stimulation of 10 ng/mL for 2 h) decreases nuclear NF-κB concentration in mice bone marrow derived dendritic cells (BDDCs)<sup>[1]</sup>.</p> <p>Lewis X trisaccharide-BSA (25 µg/mL; 48 h) or Lewis X trisaccharide (5 µg/mL; 48 h) plus ovalbumin (OVA, 25 µg/mL) increases cytokines (IL-4, IL-13, and INF-γ) level in mice splenocytes<sup>[1]</sup>.</p> <p>Lewis X trisaccharide-containing glycoconjugates stimulates B cells to proliferate and to produce factors that down-regulates the TH<sub>1</sub> immune response and up-regulates the T<sub>H</sub>2 immune response<sup>[2]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p>								
<b>In Vivo</b>	<p>Lewis X trisaccharide ( 5 µg, combined with 50 µg ovalbumin; s.c.; once a week for 2 weeks) regulates IgE/T<sub>H</sub>2 responses, and selectively increases the IgE and IgG1 responses in C3H mice, independent of the LPS-TLR4 signaling<sup>[1]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <td style="background-color: #f2f2f2; width: 30%;">Animal Model:</td> <td>Mice (BALB/c, IL-12 deficient on a BALB/c background, TLR4-defective C3H/hej, or TLR4-wild type C3H/HeOuj mice) (6-8 weeks old)<sup>[1]</sup></td> </tr> <tr> <td style="background-color: #f2f2f2;">Dosage:</td> <td>5 µg, combined with 50 µg ovalbumin</td> </tr> <tr> <td style="background-color: #f2f2f2;">Administration:</td> <td>Subcutaneous injection; once a week for 2 weeks</td> </tr> <tr> <td style="background-color: #f2f2f2;">Result:</td> <td>In C3H mice, coupled with BSA (Le<sup>x</sup>-BSA) and elicited higher levels of specific IgE and IgG1, but not IgG2a, which were associated with increased levels of splenic T<sub>H</sub>2 cytokines when compared with those seen in BSA-sensitized mice. In BALB/c mice, induced by ovalbumin, significantly increased levels of specific IgE,</td> </tr> </table>	Animal Model:	Mice (BALB/c, IL-12 deficient on a BALB/c background, TLR4-defective C3H/hej, or TLR4-wild type C3H/HeOuj mice) (6-8 weeks old) <sup>[1]</sup>	Dosage:	5 µg, combined with 50 µg ovalbumin	Administration:	Subcutaneous injection; once a week for 2 weeks	Result:	In C3H mice, coupled with BSA (Le <sup>x</sup> -BSA) and elicited higher levels of specific IgE and IgG1, but not IgG2a, which were associated with increased levels of splenic T <sub>H</sub> 2 cytokines when compared with those seen in BSA-sensitized mice. In BALB/c mice, induced by ovalbumin, significantly increased levels of specific IgE,
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resulted IgG2a antibodies concomitant with reduced levels of serum IL-12p70.  
In IL-12-deficient BALB/c mice, attenuated the downward trend of the above indicators.

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## REFERENCES

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- [1]. Hsu SC, et al. Antigen coupled with Lewis-x trisaccharides elicits potent immune responses in mice. *J Allergy Clin Immunol*. 2007 Jun;119(6):1522-8.
- [2]. van Roon AM, et al. Structure of an anti-Lewis X Fab fragment in complex with its Lewis X antigen. *Structure*. 2004 Jul;12(7):1227-36.
- [3]. Topin J, et al. The Hidden Conformation of Lewis x, a Human Histo-Blood Group Antigen, Is a Determinant for Recognition by Pathogen Lectins. *ACS Chem Biol*. 2016 Jul 15;11(7):2011-20.
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**Caution: Product has not been fully validated for medical applications. For research use only.**

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