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Lesofavumab

| | |
|-----------|---|
| Cat. No.: | HY-P99699 |
| CAS No.: | 1807960-57-1 |
| Target: | Influenza Virus |
| Pathway: | Anti-infection |
| Storage: | Please store the product under the recommended conditions in the Certificate of Analysis. |

BIOLOGICAL ACTIVITY

| | | | | | | | | | | |
|-------------------------------------|---|--|---------------|---|---------|----------|-----------------|--|---------|--|
| Description | Lesofavumab (MHAB5553A) is a human IgG1κ anti-influenza B virus antibody ^[1] . | | | | | | | | | |
| IC₅₀ & Target | Influenza B virus ^[1] | | | | | | | | | |
| In Vitro | <p>Lesofavumab (MHAB5553A) binds to a conserved epitope in the vestigial esterase domain of hemagglutinin (HA), a major immunodominant surface glycoprotein on influenza B viruses, and blocks HA-mediated membrane fusion in the endosome^[1].</p> <p>Lesofavumab (46B8) is able to neutralize all eleven IBV strains tested that were associated with human infections spanning >70 years, with IC₅₀ values ranging from 0.58 to 0.95 nM^[2].</p> <p>Lesofavumab (46B8; 0-100 ng/mL) blocks membrane fusion and induces antibody-dependent cellular cytotoxicity (ADCC) in vitro^[2].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> | | | | | | | | | |
| In Vivo | <p>Lesofavumab (46B8; 15 mg/kg; i.v.; single dose) is able to protect mice against lethal challenge of the mutant influenza B virus^[2].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <table border="1"> <tr> <td>Animal Model:</td> <td>DBA/2 J mice, B/Wisconsin/1/2010, B/Brisbane/60/2008, B/Victoria/504/2000, B/Russia/1/1969 or B/Massachusetts/3/1966 infection model^[2]</td> </tr> <tr> <td>Dosage:</td> <td>15 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>Intravenous administration, single dose, 24, 48 or 72 h post infection</td> </tr> <tr> <td>Result:</td> <td>Treatment at 24 and 48 h post infection resulted in 100% protection, with the only exception of mice infected with B/Russia/1/1969 that showed a 60% protection when treatment commenced at 48 h. Even when given at 72 h post infection, 46B8 was highly efficacious resulting in 100% protection against the B/Brisbane/60/2008, B/Victoria/504/2000 and B/Massachusetts/3/1966 strains, 75% protection against B/Wisconsin/1/2010 and 50% protection against B/Russia/1/1969.</td> </tr> </table> | | Animal Model: | DBA/2 J mice, B/Wisconsin/1/2010, B/Brisbane/60/2008, B/Victoria/504/2000, B/Russia/1/1969 or B/Massachusetts/3/1966 infection model ^[2] | Dosage: | 15 mg/kg | Administration: | Intravenous administration, single dose, 24, 48 or 72 h post infection | Result: | Treatment at 24 and 48 h post infection resulted in 100% protection, with the only exception of mice infected with B/Russia/1/1969 that showed a 60% protection when treatment commenced at 48 h. Even when given at 72 h post infection, 46B8 was highly efficacious resulting in 100% protection against the B/Brisbane/60/2008, B/Victoria/504/2000 and B/Massachusetts/3/1966 strains, 75% protection against B/Wisconsin/1/2010 and 50% protection against B/Russia/1/1969. |
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REFERENCES

[1]. Rao GK, et al. In Vivo Assessment of Antibody-Dependent Enhancement of Influenza B Infection. Toxicol Sci. 2019 Jun 1;169(2):409-421.

[2]. Chai N, et al. A broadly protective therapeutic antibody against influenza B virus with two mechanisms of action. Nat Commun. 2017 Jan 19;8:14234.

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

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