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Vixtimotamab

| | |
|------------------|---|
| Cat. No.: | HY-P99958 |
| Target: | Others |
| Pathway: | Others |
| Storage: | Please store the product under the recommended conditions in the Certificate of Analysis. |

BIOLOGICAL ACTIVITY

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|-------------------------------------|---|---------------|--|----------------|------------------------|------------------|--|---------|---|
| Description | Vixtimotamab (AMV-564; TandAb T564) is a bispecific tetravalent tandem diabody (TandAb) that targets human CD33 and human CD3 antigens. Vixtimotamab can be used for the research of acute myeloid leukemia (AML) ^[1] . | | | | | | | | |
| IC₅₀ & Target | KD: 0.3 nM (CD33, HL-60 cells), 5.1 nM (CD3, Human T cells) ^[1] | | | | | | | | |
| In Vitro | <p>Vixtimotamab (TandAb T564; 24 h) induces CD25 and CD69 with EC₅₀s of 1 pM and 2 pM, respectively^[1]. Vixtimotamab (TandAb T564; 4 days) induces T-cell proliferation in PBMCs with an EC₅₀ of 3 pM^[1]. Vixtimotamab (TandAb T564; 25 pM, 48 h) shows cytotoxicity against HL-60 and KG-1a cells^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Cytotoxicity Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>HL-60 and KG-1a cells</td> </tr> <tr> <td>Concentration:</td> <td>25 pM</td> </tr> <tr> <td>Incubation Time:</td> <td>48 h</td> </tr> <tr> <td>Result:</td> <td>Showed cytotoxicity of 86.0±1.4% and 81.3±1.5% against HL-60 and KG-1a cells, respectively.</td> </tr> </table> | Cell Line: | HL-60 and KG-1a cells | Concentration: | 25 pM | Incubation Time: | 48 h | Result: | Showed cytotoxicity of 86.0±1.4% and 81.3±1.5% against HL-60 and KG-1a cells, respectively. |
| Cell Line: | HL-60 and KG-1a cells | | | | | | | | |
| Concentration: | 25 pM | | | | | | | | |
| Incubation Time: | 48 h | | | | | | | | |
| Result: | Showed cytotoxicity of 86.0±1.4% and 81.3±1.5% against HL-60 and KG-1a cells, respectively. | | | | | | | | |
| In Vivo | <p>Vixtimotamab (TandAb T564; 0.1-10 µg/mouse, i.v.; 5 days) demonstrates dose-dependent tumor growth delay in a prophylactic HL-60 graft NOD/scid mouse model^[1]. Vixtimotamab (TandAb T564; 50 µg/mouse/d, i.v.; 7 days) substantially inhibits tumor growth in an established HL-60 xenograft NOD/scid mouse model^[1]. 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>8-week-old NOD/scid female mice, weighing 20.7 ± 1.48 g, prophylactic HL-60 graft model^[1]</td> </tr> <tr> <td>Dosage:</td> <td>0.1, 1, or 10 µg/mouse</td> </tr> <tr> <td>Administration:</td> <td>Intravenous injection, on days 0, 1, 2, 3, and 4</td> </tr> </table> | Animal Model: | 8-week-old NOD/scid female mice, weighing 20.7 ± 1.48 g, prophylactic HL-60 graft model ^[1] | Dosage: | 0.1, 1, or 10 µg/mouse | Administration: | Intravenous injection, on days 0, 1, 2, 3, and 4 | | |
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| Administration: | Intravenous injection, on days 0, 1, 2, 3, and 4 | | | | | | | | |

| | |
|---------|---|
| Result: | Demonstrated dose-dependent tumor growth delay. |
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REFERENCES

[1]. Reusch U, et al. Characterization of CD33/CD3 Tetravalent Bispecific Tandem Diabodies (TandAbs) for the Treatment of Acute Myeloid Leukemia. Clin Cancer Res. 2016 Dec 1;22(23):5829-5838.

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

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