

Hepatitis B Virus ImmuneSelect Peptide Pool Premium

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| Catalogue | 1215-07 |
| Components | 7 nmol/peptide (approx. 11.6 µg), for the stimulation of 1×10^8 cells. |
| Purity grade | ≥90% (HPLC) |
| Format | Lyophilized, no additive |
| Storage | Once reconstituted in sterile water, store aliquots at $-20\text{ }^{\circ}\text{C}$ or below. Aliquots are stable for 6 months. Do not add salts or buffers to the storage solution, as this may affect peptide stability. |

Description

The Hepatitis B virus (HBV) ImmuneSelect peptide pool contains 46 peptides derived from the following proteins:

- ◆ Capsid protein
- ◆ External core antigen
- ◆ Large envelope protein
- ◆ Protein P

The peptide sequences are a mixture of MHC-I and MHC-II epitopes, and the minimum purity of each peptide is ≥90% as determined by HPLC-MS.

Application

The *in vitro* stimulation of peripheral blood mononuclear cells (PBMCs) with Hepatitis B virus (HBV) ImmuneSelect Peptide Pool activates memory T cells specific to the infection, leading to cytokine secretion and the increased expression of specific surface markers. 7nmol per peptide for stimulation of up to 10^8 human PBMCs.

Reconstitution of peptide pool

1. Allow the lyophilized peptide pool vial to equilibrate to room temperature before opening.
2. Add 233 µL of sterile, nuclease-free water directly to the vial to fully dissolve the contents.
3. Vortex thoroughly to ensure complete solubilization of peptides.
4. The resulting stock solution contains each peptide at a concentration of 30 nmol/mL.
5. Prepare single-use working aliquots to avoid repeated freeze-thaw cycles.
6. Store working aliquots at $-20\text{ }^{\circ}\text{C}$.

Recommendations for *in vitro* stimulation

1. Wash PBMCs with sterile cell culture medium, centrifuge at $300 \times g$ for 10 min, and aspirate the supernatant.
2. Resuspend PBMCs in appropriate cell culture medium at $4 \times 10^6 - 1 \times 10^7$ cells/mL depending on the downstream experiment. For ELISpot assays, we recommend plating 200,000 cells/well in a 96-well plate.
3. Mix the reconstituted peptide stock thoroughly before use. Dilute the peptide stock solution in sterile cell culture medium to $2 \times$ the final working concentration (1.2 nmol/mL) and sterile-filter the solution prior to use.
4. Add the peptide solution to the wells at a 1:1 (v/v) ratio with the cell suspension, resulting in a final peptide concentration of 0.6 nmol/mL per peptide.
5. Incubate cells for desired time depending on the application.

Note: Incubation times can differ depending on the application. For IFN- γ ELISpot assays, an incubation period of 18–48 hours is recommended.

Need assistance?

Contact our technical support team at support@viraxbiolabs.com
We are happy to help via email or can schedule a call on request