



PerKit™ HRP-Peptide Conjugation Kit (CM32401x1 and CM32401x3) User Reference Guide

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Important Notes & Contact Information

READ BEFORE USING ANY RESOURCES PROVIDED HEREIN

The information provided in this document and the methods included in this package are for information purposes only. CellMosaic provides no warranty of performance or suitability for the purpose described herein. The performance of labeling using this kit may be affected by many different variables, including but not limited to: purity and complexity of the peptide, differences in preparation techniques, operator abilities, and environmental conditions.

Sample data are provided for illustration and example purposes only and represent a small dataset used to verify kit performance in the CellMosaic laboratory. Information about the chemicals and reagents used in the kit are provided as necessary.

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Kit Components

This kit provides materials to perform HRP labeling of one (CM32401x1) or three peptide samples (CM32401x3).

Name	Part #	Quantity (CM32401x1)	Quantity (CM32401x3)	Storage condition
Activated HRP (red color insert)	CM53211	1x2 mg	3x2 mg	-20°C
Solution A (green label)	CM01003	1.5 mL	3 mL	RT
Solution B (purple label)	CM01007	0.5 mL	0.5 mL	RT
1xPBS buffer	CM02013	3 mL	10 mL	RT
Centrifuge filter device	CM03CD030A	2	6	RT
Collection tubes	N/A	4	12	RT
1.5 mL centrifuge tube	N/A	1	3	RT
0.5 mL centrifuge tube	N/A	1	3	RT
Cys-peptide	N/A		NOT PROVIDED (User Supplied Material. 0.5 µmol for each reaction)	

Safety Information

Warning: some of the chemicals used can be potentially hazardous and can cause injury or illness. Please read and understand the Material Safety Data Sheets (MSDS) available at CellMosaic.com before you store, handle, or use any of the materials.

Labeling Chemistry

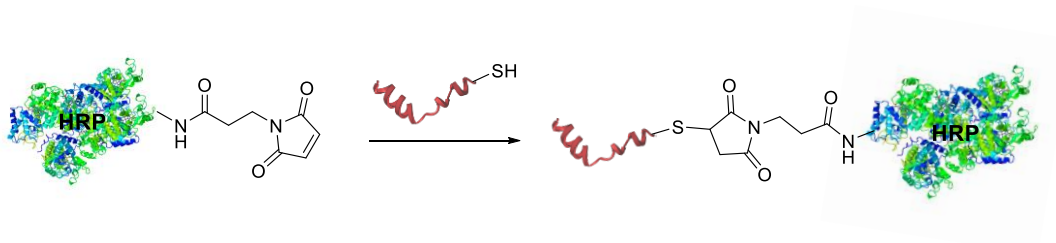
CellMosaic® has designed this personalized conjugation kit to work with any peptide containing a reactive Cys group. Using the kit components, the customer prepares the HRP-peptide conjugate by reacting their peptide (customer supplied) with activated HRP. One step purification typically provides the resulting HRP-peptide at greater than 95% purity.

This kit provides materials to label one to three peptides. Total amount of activated HRP included in a reaction: 2 mg.

Key features of this HRP-peptide conjugation kit:

- High quality maleimide-activated HRP for the conjugation: >99% purity and >200 units/mg protein activity
- Optimal maleimide groups per HRP for single-label peptide: 1.2 for typical batch
- A single purification affords over 95% of the majority of single-labeled HRP-peptide conjugates
- Fast preparation: less than 1 h hands-on time
- All reagents included, from preparation to purification
- Options to choose tailored services at CellMosaic prior to and after conjugation

- Prior to conjugation you can supply your peptide information when you place your order, and CellMosaic will give recommendations for the conjugation if your peptide has special features
- After conjugation you can choose to send your sample to CellMosaic for HPLC analysis of the conjugates



Requirement for Cys-peptide:

1. Amount: 0.5 μmol
2. HPLC purified and lyophilized: please ensure no reducing reagents, such as DTT, are present
3. The Cys peptide should be stored at -80°C
4. HPLC purity: >85% for *N*-terminal Cys peptide and >90% for *C*-terminal Cys peptide

Potential interfering compounds for labeling and conjugation reactions:

Thiols: e.g., DTT and mercaptoethanol

Protocol

1. Lab Instrumentation Needed

- Vortex mixer, centrifuge (preferably refrigerated)
- Pipettes and tips
- Timer
- Incubator or shaker set at 25°C or RT
- Balance

2. HRP Conjugation with Peptide

A1. Add 400 µL of **Solution A** (green label) to a tube containing **Activated HRP** (red insert). Vortex for 30 seconds to 1 minute to dissolve the HRP.

A2. Weigh 0.5 µmol of **Cys-peptide** into a 0.5 mL microcentrifuge tube. Note, the peptide is static charged. Use the tip of a glass Pasteur pipet to weigh the peptide if possible. It may be difficult to obtain the exact weight. Any amount between 0.5 µmol and 0.8 µmol (160%) is acceptable (no need to adjust the volume of the dissolving solution).

Calculation for 0.5 µmol:

Amount of Cys-peptide (mg) = Molecular weight of Cys-peptide x 0.0005

A3. Add 25 µL of **Solution B** (purple label) to the centrifuge tube containing Cys-peptide from **Step A2**. Vortex for 30 seconds and centrifuge the tube to get all of the liquid down to the bottom. Open the cap and add 75 µL of **Solution A** (green label) to the tube. Vortex for 30 seconds or sonicate for a few minutes to ensure all of the solid is dissolved. Discard any unused **Solution B** as hazardous chemical waste **until the experiments are done**.

Tip for solubility check (Step A1): It may take a while for your peptide to fully dissolve. In general, most of the peptide should be able to dissolve in this mixed solution system. Check the bottom of the micro-centrifuge tube to ensure the solution is clear and free of any solid residue. If after a few minutes some solid remains, centrifuge the tube and pipette the supernatant for the next step.

Tip for opening centrifuge tube after vortex: Always centrifuge the tube to ensure no liquid is in the cap.

A4. Transfer the entire solution from **Step A2** to the tube containing **Activated HRP** from **Step A5**. Pipette the solution up and down in the tube three times to mix. Incubate at room temperature for 2 hours.

3. Purification to Remove Excess Peptide

B1. Place the centrifuge tube from **Step A5** containing the reaction mixture into the centrifuge rotor and counterbalance with a similar device. Spin **the centrifuge tube** at 10,000 x g for 2 minutes.

B2. Transfer and divide the supernatant from **Step B1** into two **Centrifuge Filter Devices**. Add 250 μL of **PBS Buffer** to make up the total volume to 500 μL in each filter device and cap it.

B3. Place the capped **Filter Devices** into the centrifuge rotor, aligning the cap straps toward the center of the rotor. Spin the **Filter Devices** at 14,000 x g for 8 minutes.

B4. Remove the **Filter Devices** from the centrifuge. **Save the filtrate until the experiments are done.**

B5. Add 400 μL of **PBS Buffer** to make the total volume 500 μL . Place the two capped **Filter Devices** into the centrifuge rotor, aligning the cap strap toward the center of the rotor. Spin the **Filter Devices** at 14,000 x g for 8 minutes.

B6. If the MW of your peptide is <3000 Da, repeat **Step B5** two times

If the MW of your peptide is between 3000-5000 Da, repeat **Step B5** three times

If the MW of your peptide is >5000 Da, repeat **Step B5** four times

B7. To recover the conjugates, place the **Filter Device** upside down in a clean **Collection Tube**. Place in the centrifuge, aligning the open cap towards the center of the rotor. Spin for 2 minutes at 1,000 x g to transfer the conjugates from the **Filter Device** to the **Collection Tube**.

B8. Transfer the conjugates from the two **Collection Tubes** to a 1.5 mL micro-centrifuge tube.

B9. Rinse each **Collection Tube** with 50 μL of **PBS Buffer** and transfer the entire contents to the 1.5 mL micro-centrifuge tube from **Step B7**.

B10. Add 100 μL of **PBS Buffer** to each **Filter Device** to rinse. Stir it gently with a pipet tip, then transfer the entire contents to the 1.5 mL micro-centrifuge tube from **Step B7**. Add **PBS Buffer** to make the total volume of the sample 400 μL and cap it (use the pipetman to measure the total volume).

B11. Vortex the combined protein sample for 30 seconds and then centrifuge to ensure no liquid is in the cap.

HRP-Peptide is Ready for Your Experiment

Tip: The approximate concentration of **the HRP-peptide conjugate** is 100 μM in 400 μL of PBS buffer (4 mg/mL, assuming 80% recovery). You can also determine the concentration using a UV/Vis spectrophotometer.

Other Considerations

1. Concentration Determination

To determine the concentration, dilute your HRP-peptide from **Step B11** with 1x PBS buffer. Measure the UV absorbance of the HRP-peptide at 403 nm (A_{403}) using a UV spectrometer and calculate the concentration based on the following formula:

$$\text{Concentration } (\mu\text{M}) \text{ of the dilute sample} = (A_{403}) \times 10 / (L \times 1.02)$$

Where L is the UV cell path length (cm). If you are using a 1 cm UV cell, you can dilute HRP-peptide 20-40 times to get a good reading.

2. MW Calculation

Calculation of the MW of the conjugate:

$$\text{MW}(\text{conjugate}) = n \times \text{MW}(\text{peptide}) + 40150$$

Where n is the average molar ratio of peptide per HRP. Use 1.0 if you don't have the SEC profile of your conjugates.

3. Recommended Storage Conditions

For long-term storage, HRP-peptide conjugates can be lyophilized and stored as lyophilized powder at -20°C for 1 year.