

Research use only. Not for use in diagnostic procedures.

LANCE® Ultra

# Eu-W1024 labeled Anti-phospho Acetyl-CoA carboxylase (Ser79)

## **Antibody**

Product number: TRF0208-D Lot Number: 3254971

**Product Format:** TRF0208-D: 10 μg

TRF0208-M: 100 μg

Manufacturing date: 8/23/2023 Document version: 1

#### **Product Information**

Antibody: Europium-labeled mouse monoclonal antibody recognizing phospho-Ser79 in rat acetyl CoA

carboxylase.

Storage Buffer: 50 mM Tris-HCl (pH 7.4), 0.9% NaCl, 0.1% BSA and 0.05% sodium azide as preservative

Molecular Weight: 160 000

Stability: This product is stable for at least 24 months from the manufacturing date when stored in its

original packaging and at the recommended storage conditions.

**Storage Conditions:** Store at 4°C

## **Quality Control**

The QC release specifications are based on spectrophotometric analysis of the labeled antibody. We certify that results meet our quality release criteria.

Labeling Ratio: 7.5

Concentration:  $100 \mu g/mL (0.625 \mu M)$ 

**Recommended Assay Conditions** 

**AMPKα1 KINASE: ATP TITRATION** 

#### SUGGESTED METHOD:

(Specific applications might require optimization)

#### **Reagent Preparation**

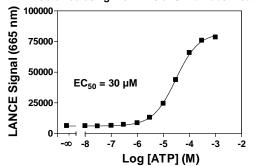
- Prepare 1X Kinase Assay Buffer: 50 mM HEPES pH 7.5, 1 mM EGTA, 10 mM MgCl₂, 2 mM DTT and 0.01% Tween-20.
- Prepare a 2X AMPK 1 solution: dilute enzyme to a concentration of 4 nM in Kinase Assay Buffer. Keep on ice.
- Prepare a 4X ULight-Acetyl CoA Carboxylase (Ser79) peptide solution: dilute ULight-Acetyl CoA carboxylase (Ser79) to a
  concentration of 200 nM in Kinase Assay Buffer.
- Prepare a 4X ATP solution: dilute ATP to concentrations ranging from 40 nM to 4 mM (serial half-log dilutions) in Kinase Assay Buffer. Keep on ice.
- Prepare 1X Detection Buffer: dilute 0.5 mL of 10X Detection Buffer with 4.5 mL of H<sub>2</sub>O.
- Prepare a 4X Stop solution\*: dilute EDTA to a concentration of 40 mM in 1X Detection Buffer.
- Prepare a 4X Detection Mix: dilute Europium-anti-phospho-Acetyl CoA Carboxylase (Ser79) antibody to a concentration of 8 nM in 1X Detection Buffer.

#### Protocol

- Pipet 5 µL of 2X AMPK11 solution into a 384-well white OptiPlate-384 (2 nM final concentration).
- Add 2.5 μL of 4X ULight- Acetyl CoA carboxylase (Ser79) solution (50 nM final concentration).
- Add 2.5 μL of 4X ATP solution (10 nM to 1 mM final concentrations).
- Cover plate with TopSeal-A and incubate 30 min at 23°C.
- Add 5 μL of 4X Stop solution\* and incubate 5 min at 23°C.
- Add 5 μL of 4X Detection Mix (2 nM Eu-anti-phospho-Acetyl CoA Carboxylase (Ser79) final conc.).
- Cover plate with TopSeal-A and incubate 60 min at 23°C.
- Remove TopSeal-A and read in TR-FRET mode at 665 nm (excitation at 320 or 340 nm).

### **Typical Product Data**

AMPKα1 kinase assay using ULight-Acetyl CoA Carboxylase (Ser79) Peptide and Eu-anti-phospho-Acetyl CoA Carboxylase (Ser79) Antibody obtained using the EnVision® Multilabel Reader:



#### **Suggested Materials**

• Substrate: ULight™- Acetyl CoA Carboxylase (Ser79) Pept.

Antibody: Eu- anti-phospho-Acetyl CoA Carboxylase (Ser79)

Kinase: AMPK21

Detection Buffer: LANCE® Detection Buffer, 10X

• Plate: OptiPlate™-384, white

#### **Supplier**

Revvity Inc

Revvity Inc

Carna Biosciences

Revvity Inc

Revvity Inc

<sup>\*</sup>Alternatively, the Stop solution and Detection Mix can be premixed and added together to the kinase reaction.

TopSeal™-A Revvity Inc

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