MSD[®] Mouse/Rat C-peptide Kit

For quantitative determination in mouse and rat serum and plasma

Alzheimer's Disease MSD MULTI-SPOT® **BioProcess** 96-Well 4-Spot Plate Cardiac SULFO-TAG[™] labeled Cell Signaling C-peptide 1. Detection Antibody Clinical Immunology 2. BSA blocked Analyte Cytokines 3. BSA blocked **Growth Factors** Δ BSA blocked Capture Antibody Hypoxia Immunogenicity Working Electrode Inflammation

Connecting peptide, or C-peptide, is a 31 amino acid protein constituent of proinsulin produced by pancreatic β -cells in the lslets of Langerhans. C-peptide connects the A-chain and the B-chain of the proinsulin polypeptide and facilitates proper processing of proinsulin in the endoplasmic reticulum. Proinsulin is cleaved by Ca2+-dependent endopeptidases; equimolar quantities of active insulin and C-peptide are stored in secretory granules, released into the portal vein, enter circulation, and are excreted by the kidneys.¹ Circulating C-peptide binds multiple cell types, including endothelial cells, fibroblasts, neuronal cells, and renal tubular cells. Bound C-peptide activates Ca2+-dependent signaling pathways, including MAPK, PLC_{γ}, and PKC, resulting in both increased eNOS and Na+K+ATPase activity and upregulation of transcription factors.²

Plasma levels of C-peptide have traditionally been used to distinguish between Type 1 and Type 2 diabetes.³ Type 1 diabetics fail to produce proinsulin so that C-peptide levels are decreased, while Type II diabetics produce increased quantities of proinsulin so that C-peptide levels are increased. C-peptide is an active biomolecule of interest in the study of hormone-dependent conditions, including gastromas associated with multiple endocrine neoplasms syndrome, polycystic ovarian syndrome, metabolic syndrome X, and sub-sets of renal disease.⁴ The MSD Mouse/Rat C-peptide assay is available on 96-well, 4-spot plates. Representative data from the assay is presented below.

Assay Sensitivity

The following standard curve illustrates the dynamic range of the Mouse/Rat C-peptide assay.



	C-peptide
Average LLOD (pg/mL)	25
LLOD Range (pg/mL)	18–32

The lower limit of detection (LLOD) is a calculated concentration based on a signal 2.5 standard deviations above the background (zero calibrator blank). The LLOD shown above was calculated based on 3 runs.

MSD Advantage

- **Multiplexing:** Multiple analytes can be measured in one well using typical sample volumes of 25 μL or less without compromising speed or performance
- Large dynamic range: Linear range of up to five logs enables the measurement of native levels of biomarkers in normal and diseased samples without multiple dilutions
- Minimal background: The stimulation mechanism (electricity) is decoupled from the response (light signal), minimizing matrix interference
- > Simple protocols: Only labels bound near the electrode surface are excited, enabling assays with fewer washes
- > Flexibility: Labels are stable, non-radioactive, and conveniently conjugated to biological molecules
- > High sensitivity and precision: Multiple rounds of label excitation and emission enhance light levels and improve sensitivity

For a complete list of products, please visit our website at www.mesoscale.com.







Ordering Information

MSD Customer Service

Metabolic Oncology

Toxicology

Catalog Numbers

Mouse/Rat C-peptide Kit

Catalog #

K1500ID-1

K1500ID-2

K1500ID-4

Vascular

Kit Size

1 plate

5 plates

25 plates

Scientific Support

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Tested Samples

Normal mouse plasma and rat serum samples were collected and tested with the Mouse/Rat C-peptide Kit. Median and range of concentrations for each sample set are displayed below.

Sample Type	Statistic	C-peptide
Mouse EDTA plasma	Median (pg/mL)	335
	Range (pg/mL)	284–391
	Number of Samples	4
	Samples in Quantitative Range	4
Rat serum	Median (pg/mL)	209
	Range (pg/mL)	108–693
	Number of Samples	3
	Samples in Quantitative Range	3

References

- 1. Steiner DF, et al. Insulin biosynthesis: evidence for a precursor. Science. 1967 Aug 11;157(3789):697-700.
- 2. Hills CE, et al. Intracellular signaling by C-peptide. Exp Diabetes Res. 2008;2008:635158.
- 3. Vasic C, et al. C-peptide: a new mediator of artherosclerosis in diabetes. Mediators Inflamm. 2012;2012:858692.
- 4. Wahren J. C-peptide: new findings and therapeutic implications in diabetes. Clin Physiol Funct Imaging. 2004 Jul;24(4):180-9.

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