



## Lipolysis (Adipocyte) Colorimetric/Fluorometric Assay Kit

### Product Information

---

**Cat**

Kit-1111

---

**Cat.No.**

Kit-1111

---

**Description**

Lipolysis is the intracellular hydrolysis of triglycerides into glycerol and free fatty acids, which are then released into the bloodstream or culture media. Lipolysis occurs in essentially all cells, but is most abundant in white and brown adipose tissue. Deficiencies in lipolysis lead to increased intracellular lipid accumulation, resulting in abnormal cellular physiology, hyperlipidemia, and insulin resistance. Lipolysis can be induced by hormones or catecholamine, which binds to  $\beta$ -adrenergic receptors leading to activation of adenylate cyclase, forming cAMP from ATP. cAMP serves as a second messenger to activate hormone-sensitive lipase, which hydrolyzes the triglycerides. This pathway is inhibited by insulin. Adipocyte Lipolysis Colorimetric/Fluorometric Assay kit is a simple, facile way to monitor lipolysis. The kit provides all reagents necessary to isolate adipocytes from up to 5 g of mouse or rat adipose tissue and measure glycerol release from primary adipocytes after induction of lipolysis. It also includes the synthetic catecholamine, Isoproterenol, to stimulate the cAMP-mediated pathway. The signal intensity is directly proportional to the amount of glycerol present. This assay kit can detect less than 200 pmol glycerol in the colorimetric assay and less than 20 pmol in the fluorometric assay.

---

**Applications**

Isolation of adipocytes from rat/mouse tissue

Measurement of lipolysis by adipocytes

Screening compounds that influence lipolysis, mechanistic studies and studies on metabolic dysfunction

---

**Storage**

-20°C



## Lipolysis (Adipocyte) Colorimetric/Fluorometric Assay Kit

### Shipping

---

Gel Pack

---

### Size

---

For 5 g Tissue

---

### Kit Components

---

Collagenase (0.2%); Collagenase Stop Buffer; Adipocyte Wash Buffer; Adipocyte Lipolysis Buffer; Glycerol Assay Buffer; Glycerol Probe (in DMSO, anhydrous); Glycerol Enzyme Mix (Lyophilized); Glycerol Standard (100 mM); Isoproterenol (10 mM); Cell strainer (100  $\mu$ m)

---

**Detection method** Absorbance (OD 570 nm) and Fluorescence (Ex/Em = 535/587 nm)

---

### Features & Benefits

---

Simple, rapid & convenient method to isolate adipocytes from adipose tissue & measure lipolysis. The assay kit can detect less than 200 pmol glycerol in the colorimetric assay and less than 20 pmol in the fluorometric assay.

---