

# Anti-LUCIFERASE (GOAT) Antibody Peroxidase Conjugated

Luciferase Antibody Peroxidase Conjugated Catalog # ASR3978

#### **Specification**

## Anti-LUCIFERASE (GOAT) Antibody Peroxidase Conjugated - Product Information

Physical State BufferLyophilized 0.02 M Potassium Phosphate, 0.15 M Sodium Chloride, pH 7.2 Luciferase [Photinus pyralis (Firefly)] 100 μL Restore with deionized water (or equivalent)Stabilizer10 mg/mL Bovine Serum Albumin (BSA) - Immunoglobulin and Protease free 0.01% (w/v) Gentamicin Sulfate. Do NOT add Sodium Azide!	Host Conjugate Clonality Application Application Note	Goat Peroxidase (Horseradish) Polyclonal WB, E, I, LCI Anti-Luciferase Peroxidase conjugated antibody is produced from Photinus pyralis (Firefly) which produces a green light with a wavelength of 562 nm. Anti-Luciferase Peroxidase has been tested by ELISA and western blot. Expect a band at approximately 60.7 kDa in size corresponding to Luciferase by western blotting in the appropriate cell lysate or extract. Anti-Luciferase has been assayed against 1.0 ug of Luciferase [Photinus pyralis (Firefly)] in a standard capture ELISA using ABTS (2,2'-azino-bis-[3-ethylbe nthiazoline-6-sulfonic acid]) code # ABTS-100 as a substrate for 30 minutes at room temperature. A working dilution of 1:50,000 to 1:200,000 of the reconstitution concentration is suggested for this product.
Sodium Chloride, pH 7.2ImmunogenReconstitution VolumeReconstitution BufferStabilizerStabilizerPreservative0.01% (w/v) Gentamicin Sulfate. Do NOT		
Immunogen Reconstitution Volume Reconstitution BufferLuciferase [Photinus pyralis (Firefly)] 100 μL Restore with deionized water (or equivalent)Stabilizer100 μL Restore with deionized water (or equivalent)Stabilizer10 mg/mL Bovine Serum Albumin (BSA) - Immunoglobulin and Protease free 0.01% (w/v) Gentamicin Sulfate. Do NOT	Butter	
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Preservative Immunoglobulin and Protease free 0.01% (w/v) Gentamicin Sulfate. Do NOT	Reconstitution Buffer	Restore with deionized water (or
	Stabilizer	
	Preservative	

### Anti-LUCIFERASE (GOAT) Antibody Peroxidase Conjugated - Additional Information

Purity

This product is an IgG fraction antibody purified from monospecific antiserum by a multi-step process which includes delipidation, salt fractionation and ion exchange chromatography followed by extensive dialysis against the buffer stated above. Assay by immunoelectrophoresis resulted in a single precipitin arc against anti-Peroxidase, anti-Goat Serum as well as purified and partially purified Luciferase [Photinus pyralis (Firefly)]. No reactivity is observed against Sea pansy (Renilla reniformis) luciferase.



#### Storage Condition

Store vial at 4° C prior to restoration. For extended storage aliquot contents and freeze at -20° C or below. Avoid cycles of freezing and thawing. Centrifuge product if not completely clear after standing at room temperature. This product is stable for several weeks at 4° C as an undiluted liquid. Dilute only prior to immediate use.

#### **Precautions Note** This product is for research use only and is not intended for therapeutic or diagnostic applications.

## Anti-LUCIFERASE (GOAT) Antibody Peroxidase Conjugated - Protein Information

## Anti-LUCIFERASE (GOAT) Antibody Peroxidase Conjugated - Protocols

Provided below are standard protocols that you may find useful for product applications.

- <u>Western Blot</u>
- <u>Blocking Peptides</u>
- Dot Blot
- Immunohistochemistry
- Immunofluorescence
- Immunoprecipitation
- Flow Cytomety
- <u>Cell Culture</u>

# Anti-LUCIFERASE (GOAT) Antibody Peroxidase Conjugated - Images

# Anti-LUCIFERASE (GOAT) Antibody Peroxidase Conjugated - Background

Anti luciferase Antibody recognizes luciferase that is commonly used in biological research as a reporter to assess the transcriptional activity in cells that are transfected with a genetic construct containing the luciferase gene under the control of a promoter of interest. Luciferase can also be used to detect the level of cellular ATP in cell viability assays or for kinase activity assays. Additionally proluminescent molecules that are converted to luciferin upon activity of a particular enzyme can be used to detect enzyme activity in coupled or two-step luciferase assays. Such substrates have been used to detect caspase activity and cytochrome P450 activity, among others.