

## Product datasheet

# Anti-Phospho SQ/TQ ATM/ATR Substrate antibody ab130947

[3 References](#) [1 Image](#)

### Overview

|                            |   |
|----------------------------|---|
| <b>Product name</b>        | Anti-Phospho SQ/TQ ATM/ATR Substrate antibody   |
| <b>Description</b>         | Rabbit polyclonal to Phospho SQ/TQ ATM/ATR Substrate  |
| <b>Host species</b>        | Rabbit  |
| <b>Specificity</b>         | Ab130947 recognizes proteins phosphorylated on SQ/TQ motifs (ATM/ATR kinase consensus phosphorylation site motif)   |
| <b>Tested applications</b> | <b>Suitable for:</b> WB   |
| <b>Species reactivity</b>  | <b>Reacts with:</b> Human   |
| <b>Immunogen</b>           | Synthetic peptide corresponding to Human Phospho SQ/TQ ATM/ATR Substrate aa 2600-2700 (phospho S2609 + T2609) conjugated to keyhole limpet haemocyanin.<br>Database link: <a href="#">P78527</a>  |
| <b>Positive control</b>    | This antibody gave a positive signal in Western Blot in the following cell line: HeLa untreated and UV treated whole cell   |
| <b>General notes</b>       | <p>The Life Science industry has been in the grips of a reproducibility crisis for a number of years. Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets your needs before purchasing.</p> <p>If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be found below, along with publications, customer reviews and Q&amp;As</p> |

### Properties

|                             |  |
|-----------------------------|--|
| <b>Form</b>                 | Liquid   |
| <b>Storage instructions</b> | Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C or -80°C. Avoid freeze / thaw cycle.   |
| <b>Storage buffer</b>       | <p>pH: 7.40</p> <p>Preservative: 0.02% Sodium azide</p> <p>Constituent: PBS</p> <p>Batches of this product that have a concentration &lt; 1mg/ml may have BSA added as a stabilising</p> |

agent. If you would like information about the formulation of a specific lot, please contact our scientific support team who will be happy to help.

|                  |                             |
|------------------|-----------------------------|
| <b>Purity</b>    | Immunogen affinity purified |
| <b>Clonality</b> | Polyclonal                  |
| <b>Isotype</b>   | IgG                         |

## Applications

**The Abpromise guarantee** Our **Abpromise guarantee** covers the use of ab130947 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

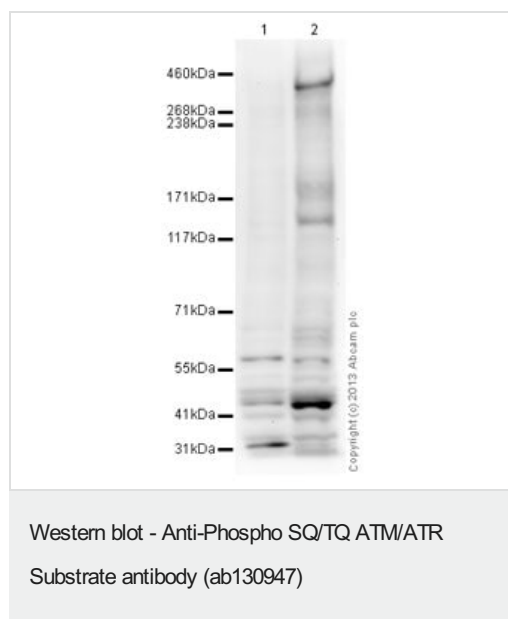
| Application | Abreviews | Notes   |
|-------------|-----------|---|
| <b>WB</b>   |           | Use a concentration of 1 µg/ml. Detects a band of approximately 450 kDa (predicted molecular weight: 469 kDa).<br>An increase in signal is observed upon UV treatment |

## Target

### Relevance

Several protein kinases mediate cellular responses to DNA damage. This includes the serine/threonine kinases ATM (ataxia telangiectasia mutated) and ATR (ATM and Rad3-related). In response to DNA damage ATM and ATR phosphorylate a large range of proteins on Serine (S) or Threonine (T) residues next to a Glutamine (Q) residue, so called SQ or TQ consensus sites.

## Images



**All lanes :** Anti-Phospho SQ/TQ ATM/ATR Substrate antibody (ab130947) at 1 µg/ml

**Lane 1 :** HeLa (Human epithelial carcinoma cell line) Whole Cell Lysate

**Lane 2 :** HeLa Whole Cell Lysate - UV Irradiated

Lysates/proteins at 25 µg per lane.

### Secondary

**All lanes :** Goat Anti-Rabbit IgG H&L (HRP) (**ab97051**) at 1/10000 dilution

Developed using the ECL technique.

Performed under reducing conditions.

**Predicted band size:** 469 kDa

**Observed band size:** 450 kDa

**Exposure time:** 8 minutes

This blot was produced using a 3-8% Tris Acetate gel under the TA buffer system. The gel was run at 150V for 60 minutes before being transferred onto a Nitrocellulose membrane at 30V for 70 minutes. The membrane was then blocked for an hour using 5% Bovine Serum Albumin before being incubated with ab130947 overnight at 4°C. Antibody binding was detected using an anti-rabbit antibody conjugated to HRP, and visualised using ECL development solution.

**Please note:** All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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