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Product datasheet for CF804867

hHR23b (RAD23B) Mouse Monoclonal Antibody [Clone ID: OTI7G10]

Product data:

| Product Type: | Primary Antibodies |
|-------------------------|--|
| Clone Name: | OTI7G10 |
| Applications: | IHC, WB |
| Recommended Dilution: | WB 1:500, IHC 1:150 |
| Reactivity: | Human, Mouse, Rat |
| Host: | Mouse |
| lsotype: | lgG1 |
| Clonality: | Monoclonal |
| Immunogen: | Human recombinant protein fragment corresponding to amino acids 1-253 of human RAD23B (NP_002865) produced in E.coli. |
| Formulation: | Lyophilized powder (original buffer 1X PBS, pH 7.3, 8% trehalose) |
| Reconstitution Method: | For reconstitution, we recommend adding 100uL distilled water to a final antibody concentration of about 1 mg/mL. To use this carrier-free antibody for conjugation experiment, we strongly recommend performing another round of desalting process. (OriGene recommends Zeba Spin Desalting Columns, 7KMWCO from Thermo Scientific) |
| Purification: | Purified from mouse ascites fluids or tissue culture supernatant by affinity chromatography (protein A/G) |
| Conjugation: | Unconjugated |
| Storage: | Store at -20°C as received. |
| Stability: | Stable for 12 months from date of receipt. |
| Predicted Protein Size: | 43 kDa |
| Gene Name: | RAD23 homolog B, nucleotide excision repair protein |
| Database Link: | <u>NP_002865</u> <u>Entrez Gene 19359 MouseEntrez Gene 298012 RatEntrez Gene 5887 Human</u> <u>P54727</u> |



Scheme Grigene hHR23b (RAD23B) Mouse Monoclonal Antibody [Clone ID: OTI7G10] – CF804867

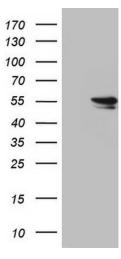
Background:The protein encoded by this gene is one of two human homologs of Saccharomyces
cerevisiae Rad23, a protein involved in the nucleotide excision repair (NER). This protein was
found to be a component of the protein complex that specifically complements the NER
defect of xeroderma pigmentosum group C (XP-c) cell extracts in vitro. This protein was also
shown to interact with, and elevate the nucleotide excision activity of 3-methyladenine-DNA
glycosylase (MPG), which suggested a role in DNA damage recognition in base excision repair.
This protein contains an N-terminal ubiquitin-like domain, which was reported to interact
with 26S proteasome, and thus this protein may be involved in the ubiquitin mediated
proteolytic pathway in cells. Alternative splicing results in multiple transcript variants
encoding distinct isoforms. [provided by RefSeq, Sep 2011]

| Synonyms: | HHR23B; HR23B; P58 |
|-----------|--------------------|
| | |

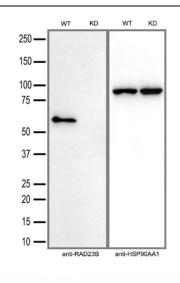
Protein Families: Druggable Genome

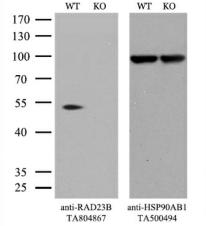
Protein Pathways: Nucleotide excision repair

Product images:



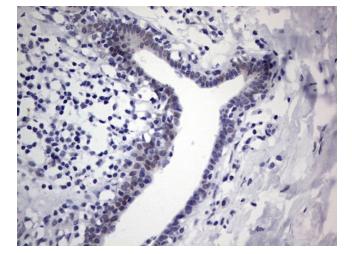
HEK293T cells were transfected with the pCMV6-ENTRY control (Cat# [PS100001], Left lane) or pCMV6-ENTRY RAD23B (Cat# [RC202185], Right lane) cDNA for 48 hrs and lysed. Equivalent amounts of cell lysates (5 ug per lane) were separated by SDS-PAGE and immunoblotted with anti-RAD23B(Cat# [TA804867]). Positive lysates [LY401012] (100ug) and [LC401012] (20ug) can be purchased separately from OriGene.



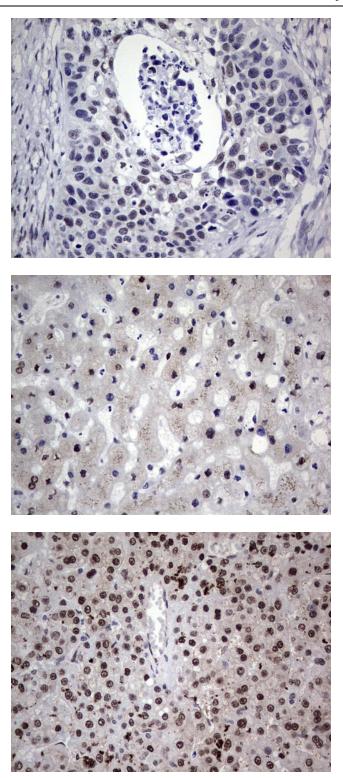


Equivalent amounts of cell lysates (30 ug per lane) of wild-type HT-1080 cells (WT) and RAD23B-Knockdown HT-1080 cells (KD) were separated by SDS-PAGE and immunoblotted with anti-RAD23B monoclonal antibody [TA804867] (1:2500). Then the blotted membrane was stripped and reprobed with anti-HSP90AA1 antibody as a loading control.

Equivalent amounts of cell lysates (10 ug per lane) of wild-type Hela cells (WT, Cat# LC810HELA) and RAD23B-Knockout Hela cells (KO, Cat# [LC810196]) were separated by SDS-PAGE and immunoblotted with anti-RAD23B monoclonal antibody [TA804867]. Then the blotted membrane was stripped and reprobed with anti-HSP90AB1 antibody ([TA500494]) as a loading control (1:500).



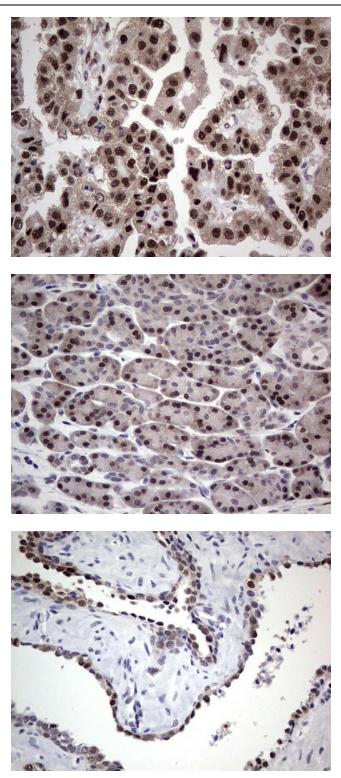
Immunohistochemical staining of paraffinembedded Adenocarcinoma of Human breast tissue using anti-RAD23B mouse monoclonal antibody. Heat-induced epitope retrieval by EDTA solution buffer pH 8.0 at 120°C for 3 min.



Immunohistochemical staining of paraffinembedded Carcinoma of Human kidney tissue using anti-RAD23B mouse monoclonal antibody. Heat-induced epitope retrieval by EDTA solution buffer pH 8.0 at 120°C for 3 min.

Immunohistochemical staining of paraffinembedded Human liver tissue within the normal limits using anti-RAD23B mouse monoclonal antibody. Heat-induced epitope retrieval by EDTA solution buffer pH 8.0 at 120°C for 3 min.

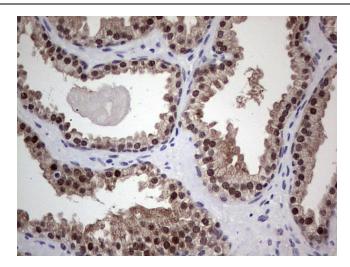
Immunohistochemical staining of paraffinembedded Carcinoma of Human liver tissue using anti-RAD23B mouse monoclonal antibody. Heat-induced epitope retrieval by EDTA solution buffer pH 8.0 at 120°C for 3 min.



Immunohistochemical staining of paraffinembedded Adenocarcinoma of Human ovary tissue using anti-RAD23B mouse monoclonal antibody. Heat-induced epitope retrieval by EDTA solution buffer pH 8.0 at 120°C for 3 min.

Immunohistochemical staining of paraffinembedded Human pancreas tissue within the normal limits using anti-RAD23B mouse monoclonal antibody. Heat-induced epitope retrieval by EDTA solution buffer pH 8.0 at 120°C for 3 min.

Immunohistochemical staining of paraffinembedded Human prostate tissue within the normal limits using anti-RAD23B mouse monoclonal antibody. Heat-induced epitope retrieval by EDTA solution buffer pH 8.0 at 120°C for 3 min.



Immunohistochemical staining of paraffinembedded Carcinoma of Human prostate tissue using anti-RAD23B mouse monoclonal antibody. Heat-induced epitope retrieval by EDTA solution buffer pH 8.0 at 120°C for 3 min.