

## **Product Data Sheet**

## **Anti-BWR1A Antibody**

Catalog # Source Reactivity Applications

CPA6133 Rabbit H, M, R, Mk WB, IF/IC

**Description** Rabbit polyclonal antibody to BWR1A

Immunogen KLH-conjugated synthetic peptide encompassing a sequence within the C-term

region of human BWR1A. The exact sequence is proprietary.

**Purification** The antibody was purified by immunogen affinity chromatography.

**Specificity** Recognizes endogenous levels of BWR1A protein.

**Clonality** Polyclonal

Conjugation

Form Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,

and 0.01% sodium azide.

**Dilution** WB (1/500 - 1/1000), IF/IC (1/100 - 1/500)

Gene Symbol SLC22A18

Alternative Names BWR1A; BWSCR1A; HET; IMPT1; ITM; ORCTL2; SLC22A1L; TSSC5; Solute carrier

family 22 member 18; Beckwith-Wiedemann syndrome chromosomal region 1

candidate gene A protein; Efflux transporter-like protein; Imprinted

multi-membrane-spanning polyspecific transporter-related protein 1; Organic cation

transporter-like protein 2; ORCTL-2; Solute carrier family 22 member 1-like;

Tumor-suppressing STF cDNA 5 protein; Tumor-suppressing subchromosomal

transferable fragment candidate gene 5 protein; p45-Beckwith-Wiedemann region 1

A; p45-BWR1A

Entrez Gene 5002 (Human); 18400 (Mouse); 309131 (Rat)

SwissProt Q96BI1 (Human); Q78KK3 (Mouse); Q6AY78 (Rat)

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility

Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference

Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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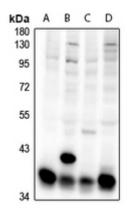
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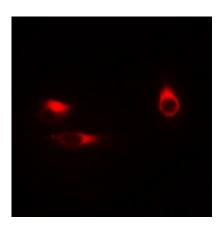
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Storage/Stability

Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid freeze/thaw cycles.



Western blot analysis of BWR1A expression in C6 (A), CT26 (B), Hela (C), A2780 (D) whole cell lysates. (Predicted band size: 44 kD; Observed band size: 40; 44 kD)



Immunofluorescent analysis of BWR1A staining in HeLa cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a Alexa Fluor 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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