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| <b>Cat. No:</b>       | ABN19521   |
| <b>Conjugate:</b>     | Unconjugated   |
| <b>Size:</b>          | 100µL  |
| <b>Clone:</b>         | Polyclonal   |
| <b>Concentration:</b> | 1mg/ml   |
| <b>Host:</b>          | Rabbit   |
| <b>Isotype:</b>       | IgG  |
| <b>Immunogen:</b>     | Synthesized peptide derived from the Internal region of human UBE2G1.  |
| <b>Reactivity:</b>    | Human,Mouse,Rat  |
| <b>Applications:</b>  | IHC 1:100-1:300,ICC/IF 1:50-1:200,ELISA 1:20000-1:40000  |
| <b>Purification:</b>  | Affinity purification  |
| <b>Synonyms:</b>      | UBE2G1; UBE2G; Ubiquitin-conjugating enzyme E2 G1; E217K; UBC7; Ubiquitin carrier protein G1; Ubiquitin-protein ligase G1  |
| <b>Background:</b>    | <p>The modification of proteins with ubiquitin is an important cellular mechanism for targeting abnormal or short-lived proteins for degradation. Ubiquitination involves at least three classes of enzymes: ubiquitin-activating enzymes, or E1s, ubiquitin-conjugating enzymes, or E2s, and ubiquitin-protein ligases, or E3s. This gene encodes a member of the E2 ubiquitin-conjugating enzyme family and catalyzes the covalent attachment of ubiquitin to other proteins. The protein may be involved in degradation of muscle-specific proteins. [provided by RefSeq, Jul 2008],catalytic activity:ATP + ubiquitin + protein lysine = AMP + diphosphate + protein N-ubiquityllysine.,function:Catalyzes the covalent attachment of ubiquitin to other proteins. May be involved in degradation of muscle-specific proteins.,pathway:Protein modification; protein ubiquitination.,similarity:Belongs to the ubiquitin-conjugating enzyme family.,tissue specificity:Widely expressed, mainly in skeletal muscle.,</p> |
| <b>Form:</b>          | Liquid   |
| <b>Buffer:</b>        | Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% New type preservative N.   |
| <b>Storage:</b>       | Store at 4°C short term. Aliquot and store at -20°C for 12 months. Avoid freeze/thaw cycles.   |

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