

APC3 (D3I1V) Rabbit mAb



877-616-CELL (2355) Orders:

orders@cellsignal.com

Support: 877-678-TECH (8324)

Web: info@cellsignal.com

cellsignal.com

3 Trask Lane | Danvers | Massachusetts | 01923 | USA

Product Usage Information Application Western Blotting 1:1000 Immunoprecipitation 1:1000 Storage Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA, 50% glycerol and less that 0.02% sodium azide. Store at ~20°C. Do not aliquot the antibody. Specificity/Sensitivity APC/3 (D3I1V) Rabbit mAb recognizes endogenous levels of total APC3 protein. This antibody does recreased on 100% sequence homology Source / Purification Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues near the carboxy terminus of human APC3 protein. Cell proliferation in all eukaryotic cells depends strictly upon the ubiquitin ligase (E3) activity of the anaphase promoting complex/cyclosome (APC/C), whose main function is to trigger the transition the cell cycle from metaphase to anaphase. APC/C performs its various functions by promoting the assembly of polyubiquitin chains on substrate APC/C subunits have been identified. Like E3 enzymes, APC/C utilizes ubiquitin residues that have been activated by E1 enzymes and then transferred to E2 enzymes. Indeed, APC/C has been shown to interact with UBEZS and UBEZC E2 enzymes, in part, via the RING-finger domain-containing subunit, APC11 (3-5), APC/C activity is also strictly dependent upon its association with multiple cofactors. For example, the related proteins, Cd220 and Cd11/FZR1, participate in the recognition of APC/C substrates by interacting with specific recognition elements in these substrates (6), called D-boxes (7) and KEN-boxes (8). Anaphase-promoting complex subunit 3 (APC3), APC8, and APC6 are components of the tetratricopeptide (TPR) APC/C subcomplex (9). The presence of APC3 is required for binding of	For Research Use Only. Not for Use in Diagnostic Procedures.							
Information Western Blotting Immunoprecipitation Storage Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA, 50% glycerol and less that 0.02% sodium azide. Store at ~20°C. Do not aliquot the antibody. Specificity/Sensitivity APC3 (D3I1V) Rabbit mAb recognizes endogenous levels of total APC3 protein. This antibody does not cross-react with either APC8/CDC23 or APC6/CDC16. Species predicted to react based on 100% sequence homology Source / Purification Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues near the carboxy terminus of human APC3 protein. Background Cell proliferation in all eukaryotic cells depends strictly upon the ubiquitin ligase (E3) activity of the anaphase promoting complex/cyclosome (APC/C), whose main function is to trigger the transition the cell cycle from metaphase to anaphase. APC/C performs its various functions by promoting the assembly of polyubiquitin chains on substrate proteins, which targets these proteins for degradation by the 265 proteasome (1,2). In humans, twelve different APC/C subunits have been identified. Like E3 enzymes, APC/C utilizes ubiquitin residue that have been activated by E1 enzymes and then transferred to E2 enzymes. Indeed, APC/C has been shown to interact with UBE2S and UBE2C E2 enzymes, in part, via the RING-finger domain-containing subunit, APC11 (3-5). APC/C activity is also strictly dependent upon its association with multiple cofactors. For example, the related proteins, Cdc20 and Cdh1/FZR1, participate in the recognition of APC/C substrates by interacting with specific recognition elements in these substrates (6), called 0-boxes (7) and KEN-boxes (8). Anaphase-promoting complex subunit 3 (APC3, APC8, and APC6 are components of the tetratricopeptide (TPR) APC/C subcomplex (9). The presence of APC3 is required for binding of							Entrez-Gene Id 996	
O.02% sodium azide. Store at –20°C. Do not aliquot the antibody. APC3 (D3I1V) Rabbit mAb recognizes endogenous levels of total APC3 protein. This antibody does not cross-react with either APC8/CDC23 or APC6/CDC16. Species predicted to react based on 100% sequence homology Source / Purification Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues near the carboxy terminus of human APC3 protein. Cell proliferation in all eukaryotic cells depends strictly upon the ubiquitin ligase (E3) activity of the anaphase promoting complex/cyclosome (APC/C), whose main function is to trigger the transition-the cell cycle from metaphase to anaphase. APC/C performs its various functions by promoting the assembly of polyubiquitin chains on substrate proteins, which targets these proteins for degradatiby the 26S proteasome (1,2). In humans, twelve different APC/C subunits have been identified. Like E3 enzymes, APC/C utilizes ubiquititin residues that have been activated by E1 enzymes and then transferred to E2 enzymes. Indeed, APC/C has been shown to interact with UBE2S and UBE2C E2 enzymes, in part, via the RING-finger domain-containing subunit, APC11 (3-5). APC/C activity is also strictly dependent upon its association with multiple cofactors. For example, the related proteins, Cdc20 and Cdh1/FZR1, participate in the recognition of APC/C substrates by interacting with specific recognition elements in these substrates (6), called D-boxes (7) and KEN-boxes (8). Anaphase-promoting complex subunit 3 (APC3), APC8, and APC6 are components of the tetratricopeptide (TPR) APC/C subcomplex (9). The presence of APC3 is required for binding of			Western Blotting			1:1000		
Species predicted to react based on 100% sequence homology Source / Purification Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues near the carboxy terminus of human APC3 protein. Background Cell proliferation in all eukaryotic cells depends strictly upon the ubiquitin ligase (E3) activity of the anaphase promoting complex/cyclosome (APC/C), whose main function is to trigger the transition of the cell cycle from metaphase to anaphase. APC/C performs its various functions by promoting the assembly of polyubiquitin chains on substrate proteins, which targets these proteins for degradation by the 26S proteasome (1,2). In humans, twelve different APC/C subunits have been identified. Like E3 enzymes, APC/C utilizes ubiquitin residues that have been activated by E1 enzymes and then transferred to E2 enzymes. Indeed, APC/C has been shown to interact with UBE2S and UBE2C E2 enzymes, in part, via the RING-finger domain-containing subunit, APC11 (3-5). APC/C activity is also strictly dependent upon its association with multiple cofactors. For example, the related proteins, Cdc20 and Cdh1/FZR1, participate in the recognition of APC/C substrates by interacting with specific recognition elements in these substrates (6), called D-boxes (7) and KEN-boxes (8). Anaphase-promoting complex subunit 3 (APC3), APC8, and APC6 are components of the tetratricopeptide (TPR) APC/C subcomplex (9). The presence of APC3 is required for binding of	Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.					
Source / Purification Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues near the carboxy terminus of human APC3 protein. Cell proliferation in all eukaryotic cells depends strictly upon the ubiquitin ligase (E3) activity of the anaphase promoting complex/cyclosome (APC/C), whose main function is to trigger the transition the cell cycle from metaphase to anaphase. APC/C performs its various functions by promoting the assembly of polyubiquitin chains on substrate proteins, which targets these proteins for degradative by the 26S proteasome (1,2). In humans, twelve different APC/C subunits have been identified. Like E3 enzymes, APC/C utilizes ubiquitin residues that have been activated by E1 enzymes and then transferred to E2 enzymes. Indeed, APC/C has been shown to interact with UBE2S and UBE2C E2 enzymes, in part, via the RING-finger domain-containing subunit, APC11 (3-5). APC/C activity is also strictly dependent upon its association with multiple cofactors. For example, the related proteins, Cdc20 and Cdh1/FZR1, participate in the recognition of APC/C substrates by interacting with specific recognition elements in these substrates (6), called D-boxes (7) and KEN-boxes (8). Anaphase-promoting complex subunit 3 (APC3), APC8, and APC6 are components of the tetratricopeptide (TPR) APC/C subcomplex (9). The presence of APC3 is required for binding of	Specificity/Sensitivity		APC3 (D3I1V) Rabbit mAb recognizes endogenous levels of total APC3 protein. This antibody does not cross-react with either APC8/CDC23 or APC6/CDC16.					
residues near the carboxy terminus of human APC3 protein. Cell proliferation in all eukaryotic cells depends strictly upon the ubiquitin ligase (E3) activity of the anaphase promoting complex/cyclosome (APC/C), whose main function is to trigger the transition the cell cycle from metaphase to anaphase. APC/C performs its various functions by promoting the assembly of polyubiquitin chains on substrate proteins, which targets these proteins for degradatic by the 26S proteasome (1,2). In humans, twelve different APC/C subunits have been identified. Like E3 enzymes, APC/C utilizes ubiquitin residues that have been activated by E1 enzymes and then transferred to E2 enzymes. Indeed, APC/C has been shown to interact with UBE2S and UBE2C E2 enzymes, in part, via the RING-finger domain-containing subunit, APC11 (3-5). APC/C activity is also strictly dependent upon its association with multiple cofactors. For example, the related proteins, Cdc20 and Cdh1/FZR1, participate in the recognition of APC/C substrates by interacting with specific recognition elements in these substrates (6), called D-boxes (7) and KEN-boxes (8). Anaphase-promoting complex subunit 3 (APC3), APC8, and APC6 are components of the tetratricopeptide (TPR) APC/C subcomplex (9). The presence of APC3 is required for binding of	based on 100% sequence		Hamster, Bovine, Dog, Pig, Horse					
anaphase promoting complex/cyclosome (APC/C), whose main function is to trigger the transition of the cell cycle from metaphase to anaphase. APC/C performs its various functions by promoting the assembly of polyubiquitin chains on substrate proteins, which targets these proteins for degradation by the 26S proteasome (1,2). In humans, twelve different APC/C subunits have been identified. Like E3 enzymes, APC/C utilizes ubiquitin residues that have been activated by E1 enzymes and then transferred to E2 enzymes. Indeed, APC/C has been shown to interact with UBE2S and UBE2C E2 enzymes, in part, via the RING-finger domain-containing subunit, APC11 (3-5). APC/C activity is also strictly dependent upon its association with multiple cofactors. For example, the related proteins, Cdc20 and Cdh1/FZR1, participate in the recognition of APC/C substrates by interacting with specific recognition elements in these substrates (6), called D-boxes (7) and KEN-boxes (8). Anaphase-promoting complex subunit 3 (APC3), APC8, and APC6 are components of the tetratricopeptide (TPR) APC/C subcomplex (9). The presence of APC3 is required for binding of	Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues near the carboxy terminus of human APC3 protein.					
with APC3 that enables APC/C to recognize the D-box of substrates (6,10). APC3 localizes to the centrosome and the mitotic spindle, suggesting that APC3 plays a critical role in the transition from metaphase to anaphase (11). Phosphorylation of APC3 at multiple sites during mitosis likely leads t	Buckground		anaphase promoting complex/cyclosome (APC/C), whose main function is to trigger the transition of the cell cycle from metaphase to anaphase. APC/C performs its various functions by promoting the assembly of polyubiquitin chains on substrate proteins, which targets these proteins for degradation by the 26S proteasome (1,2). In humans, twelve different APC/C subunits have been identified. Like all E3 enzymes, APC/C utilizes ubiquitin residues that have been activated by E1 enzymes and then transferred to E2 enzymes. Indeed, APC/C has been shown to interact with UBE2S and UBE2C E2 enzymes, in part, via the RING-finger domain-containing subunit, APC11 (3-5). APC/C activity is also strictly dependent upon its association with multiple cofactors. For example, the related proteins, Cdc20 and Cdh1/FZR1, participate in the recognition of APC/C substrates by interacting with specific recognition elements in these substrates (6), called D-boxes (7) and KEN-boxes (8). Anaphase-promoting complex subunit 3 (APC3), APC8, and APC6 are components of the tetratricopeptide (TPR) APC/C subcomplex (9). The presence of APC3 is required for binding of Cdh1/FZR1 to the APC/C. This suggests that APC/C is activated by an association between Cdh1/FZR1 with APC3 that enables APC/C to recognize the D-box of substrates (6,10). APC3 localizes to the centrosome and the mitotic spindle, suggesting that APC3 plays a critical role in the transition from metaphase to anaphase (11). Phosphorylation of APC3 at multiple sites during mitosis likely leads to structural changes within the APC/C by altering subunit interactions or changing affinity for molecules					

Background References

- 1. Qiao, X. et al. (2010) *Cell Cycle* 9, 3904-12.
- 2. Harper, J.W. et al. (2002) *Genes Dev* 16, 2179-206.
- 3. Carroll, C.W. and Morgan, D.O. (2002) Nat Cell Biol 4, 880-7.
- 4. Gmachl, M. et al. (2000) Proc Natl Acad Sci U S A 97, 8973-8.
- 5. Leverson, J.D. et al. (2000) *Mol Biol Cell* 11, 2315-25.
- 6. Kraft, C. et al. (2005) Mol Cell 18, 543-53.
- 7. Glotzer, M. et al. (1991) Nature 349, 132-8.
- 8. Pfleger, C.M. and Kirschner, M.W. (2000) *Genes Dev* 14, 655-65.
- 9. Tugendreich, S. et al. (1993) *Proc Natl Acad Sci U S A* 90, 10031-5.
- 10. Vodermaier, H.C. et al. (2003) Curr Biol 13, 1459-68.
- 11. Tugendreich, S. et al. (1995) Cell 81, 261-8.
- 12. Topper, L.M. et al. Cell Cycle 1, 282-92.
- 13. Kraft, C. et al. (2003) EMBO J 22, 6598-609.

Western Blot Buffer IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v nonfat

dry milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

Applications Key W: Western Blotting IP: Immunoprecipitation

Cross-Reactivity Key H: Human M: Mouse R: Rat Mk: Monkey

Trademarks and Patents Cell Signaling Technology is a trademark of Cell Signaling Technology, Inc.

XP is a registered trademark of Cell Signaling Technology, Inc.

All other trademarks are the property of their respective owners. Visit cellsignal.com/trademarks for

more information.

Limited Uses

Except as otherwise expressly agreed in a writing signed by a legally authorized representative of CST, the following terms apply to Products provided by CST, its affiliates or its distributors. Any Customer's terms and conditions that are in addition to, or different from, those contained herein, unless separately accepted in writing by a legally authorized representative of CST, are rejected and are of no force or effect.

Products are labeled with For Research Use Only or a similar labeling statement and have not been approved, cleared, or licensed by the FDA or other regulatory foreign or domestic entity, for any purpose. Customer shall not use any Product for any diagnostic or therapeutic purpose, or otherwise in any manner that conflicts with its labeling statement. Products sold or licensed by CST are provided for Customer as the end-user and solely for research and development uses. Any use of Product for diagnostic, prophylactic or therapeutic purposes, or any purchase of Product for resale (alone or as a component) or other commercial purpose, requires a separate license from CST. Customer shall (a) not sell, license, loan, donate or otherwise transfer or make available any Product to any third party, whether alone or in combination with other materials, or use the Products to manufacture any commercial products, (b) not copy, modify, reverse engineer, decompile, disassemble or otherwise attempt to discover the underlying structure or technology of the Products, or use the Products for the purpose of developing any products or services that would compete with CST products or services, (c) not alter or remove from the Products any trademarks, trade names, logos, patent or copyright notices or markings, (d) use the Products solely in accordance with CST Product Terms of Sale and any applicable documentation, and (e) comply with any license, terms of service or similar agreement with respect to any third party products or services used by Customer in connection with the Products.