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## **Revision 3** Cell Signaling Store at -20C SF3B1 (D7L5T) Rabbit mAb H. Orders: 877-616-CELL (2355) orders@cellsignal.com 4434 877-678-TECH (8324) Support: Web: info@cellsignal.com cellsignal.com 3 Trask Lane | Danvers | Massachusetts | 01923 | USA

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Applications: W, W-S	<b>Reactivity:</b> H M R Mk	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 155	<b>Source/Isotype:</b> Rabbit IgG	<b>UniProt ID:</b> #075533	Entrez-Gene Id: 23451	
Product Usage Information		<b>Application</b> Western Blotting Simple Western™			<b>Dilution</b> 1:1000 1:10 - 1:50		
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. <i>Do not aliquot the antibody.</i>					
Specificity/Sensitivity		SF3B1 (D7L5T) Rabbit mAb recognizes endogenous levels of total SF3B1 protein.					
Species predicted to react based on 100% sequence homology		Hamster, Bovine					
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues near the amino terminus of human SF3B1 protein.					
Background		Splicing factor 3b subunit 1 (SF3B1) is an integral component of the U2 small nuclear ribonucleoprotein (U2 snRNP) and plays an important role in the splicing of pre-mRNA that involves the removal of introns and the joining of exons to form mature mRNA (1-3). The assembly and proper recognition of splice sites are driven by sequences at the pre-mRNA intron-exon splice sites. The 5' splice donor site is recognized by the U1 snRNP complex, while U2 snRNP binds to the 3' splice site (branch point), ensuring the anchoring of the spliceosome machinery at the splice sites (3,4). Recent whole exome sequencing studies have demonstrated a high incidence of somatic mutations of <i>SF3B1</i> in patients with various hematological malignancies such as chronic lymphocytic leukemia and myelodysplastic syndromes (2,3,5,6). Misregulation of pre-mRNA splicing arising from mutations of the spliceosome components such as SF3B1 is thought to contribute to changes in the expression patterns of key proteins that are involved in pathways such as cell cycle progression, cell death, and cancer metabolism (2,3).					
Background References		1. Jurica, M.S. and Moore, M.J. (2003) <i>Mol Cell</i> 12, 5-14. 2. Cazzola, M. et al. (2013) <i>Blood</i> 121, 260-9. 3. Bonnal, S. et al. (2012) <i>Nat Rev Drug Discov</i> 11, 847-59. 4. Gozani, O. et al. (1998) <i>Mol Cell Biol</i> 18, 4752-60. 5. Quesada, V. et al. (2012) <i>Nat Genet</i> 44, 47-52. 6. Baliakas, P. et al. (2015) <i>Leukemia</i> 29, 329-36.					
Species Reactivity		Species reactivity is determined by testing in at least one approved application (e.g., western blot).					
Western Blot Buffer		IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.					
Applications Key Your web bro		W: Western Blotting W-S: Simple Western™ wser (Chrome 109) has a serious security vulnerability!					

Cross-Reactivity Key IC DROWS H: Human M: Mouse R: Rat Mk: Monkey Ind the Dest experience on this site.					
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