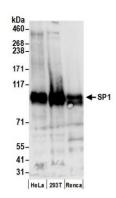
## Sp1 Antibody

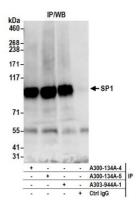
Rabbit Polyclonal Antigen Affinity Purified			Protein ID	XP_028606.4	
Catalog No. A300–134A		GenelD	6667		
Lot No.	A300-	134A-5			LABORATORIES, INC
APPLICATIONS		WB, IP, IHC, ChIP–Seq			
SPECIES REACTIVITY		Human, Mouse			
AMOUNT		100 µl			
CONCENTRATION		1000 μg/ml			
STORAGE/SHELF LIFE		2 - 8° C / 1 year from date of receipt			
PHYSICAL STATE		Liquid			
BUFFER		Tris-citrate/phosphate buffer, pH 7 to 8 containing 0.09% Sodium Azide			
ISOTYPE		IgG			
ORIGIN		USA			
PRODUCTION PROCEDURES		Antibody was affinity purified using the antigen immobilized on solid support.			
		The epitope recognized by A300–134A maps to a region between residue 750 and the C- terminus (residue 785) using the numbering given in entry XP_028606.4 (GeneID 6667).			
		Immunoglobulin concentration was determined by extinction coefficient: absorbance at 280 nm of 1.4 equals 1.0 mg of IgG.			
		Centrifuge tube to remove product from lid. Optimal working dilutions should be determined experimentally by the investigator. Prepare working dilution immediately before use.			
		Western Blot	1:2	2,000 - 1:10,000	
		Immunoprecipit	ation 2 -	- 10 µg/mg lysate	
		Immunohistoch	,	2,000 – 1:10,000. Epitope retrieval with commended for FFPE tissue sections.	a citrate buffer pH 6.0 is
APPLICATION NOTES		ChIP–Seq	4 µ	ıg/30 μg chromatin	
		Western blot of immunoprecipitates performed using Normal Pig Serum (Cat. No. S100-020), Goat anti-Rabbit Light Chain HRP Conjugate (Cat. No. A120-113P) and 4-8% SDS-PAGE (link to IP-western blot protocol in Additional Info section below).			
IHC HUMAN CONTROLS		Western blot of lysates performed using standard western blot reagents and 4–8% SDS-PAGE. Breast Carcinoma, Ovarian Carcinoma			
IHC MOUSE CONTROLS Renal Cell Card			noma		
ADDITIONAL INFO		https://www.bethyl.com/product/A300-134A Use the link above to view SDS, a current list of citations, and other product specific information. IP-western blot protocol: https://www.bethyl.com/content/protocol_IP_WB			

This document certifies that this product has met all of the quality control standards defined by Bethyl Laboratories, Inc. Eric McIntush, PhD | Chief Scientific Officer Date: June 21, 2019

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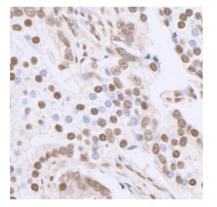


Detection of human and mouse SP1 by western blot. Samples: Whole cell lysate (50  $\mu$ g) from HeLa, HEK293T, and mouse Renca cells prepared using NETN lysis buffer. Antibody: Affinity purified rabbit anti–SP1 antibody A300–134A (lot A300–134A–5) used for WB at 0.1  $\mu$ g/ml. Detection: Chemiluminescence with an exposure time of 10 seconds.

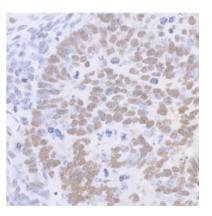


Detection of human SP1 by western blot of

**immunoprecipitates.** Samples: Whole cell lysate (0.5 or 1.0 mg per IP reaction; 20% of IP loaded) from HeLa cells prepared using NETN lysis buffer. Antibodies: Affinity purified rabbit anti–SP1 antibody A300–134A (lot A300–134A–5) used for IP at 6  $\mu$ g per reaction. SP1 was also immunoprecipitated by a previous lot of this antibody (A300–134A–4) and rabbit anti–SP1 antibody A303–944A. For blotting immunoprecipitated SP1, A300–134A was used at 1  $\mu$ g/ml. Detection: Chemiluminescence with an exposure time of 10 seconds.

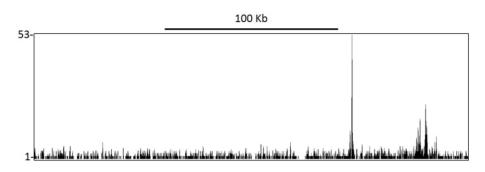


Detection of human SP1 by immunohistochemistry. Sample: FFPE section of human breast carcinoma. Antibody: Affinity purified rabbit anti-SP1 (Cat. No. A300-134A Lot5) used at a dilution of 1:5,000 (0.2µg/ml). Detection: DAB



**Detection of mouse SP1 by immunohistochemistry.** *Sample:* FFPE section of mouse renal cell carcinoma. *Antibody:* Affinity purified rabbit anti-SP1 (Cat. No. A300-134A Lot5) used at a dilution of 1:5,000 (0.2µg/ml). *Detection:* DAB

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**Localization of Sp1 Binding Sites by ChIP-sequencing.** Chromatin from acute myeloid leukemia cell line was immunoprecipitated with anti-Sp1 antibody A300-134A and analyzed by DNA sequencing. The figure illustrates the peak distribution of Sp1 binding within a 250 Kb region of chromosome 6 as detected using anti-Sp1 A300-134A. ChIP-seq validation performed by Active Motif, Carlsbad, CA.