

p53 Monoclonal Antibody (DO-7)

Catalog Number **MA5-12557**

Product data sheet

Details		Species Reactivity	
Size	500 µL	Species reactivity	Bovine, Human
Host/Isotope	Mouse / IgG2b, kappa	Published species	Rat, Human, Mouse, Not Applicable
Class	Monoclonal	Tested Applications	
Type	Antibody	ChIP assay (ChIP)	Dilution *
Clone	DO-7	Flow Cytometry (Flow)	Assay-dependent
Immunogen	Recombinant human wild-type p53 protein expressed in E. coli	Immunohistochemistry (Paraffin) (IHC (P))	1:50
Conjugate	Unconjugated	Immunoprecipitation (IP)	1:100-1:200
Form	Liquid	Western Blot (WB)	1:50-1:100
Concentration	0.05 mg/mL	Immunocytochemistry (ICC/IF)	1:500-1:1,000
Purification	Protein A		1:20-1:200
Storage buffer	PBS, pH 7.4	Published Applications	
Contains	0.05% sodium azide	Western Blot (WB)	See 35 publications below
Storage Conditions	-20°C	Immunohistochemistry (IHC)	See 97 publications below
		Immunoprecipitation (IP)	See 2 publications below
		Immunocytochemistry (ICC/IF)	See 4 publications below
		Immunohistochemistry (Frozen) (IHC (F))	See 1 publications below
		Flow Cytometry (Flow)	See 1 publications below
		Miscellaneous PubMed (Misc)	See 5 publications below
		Immunohistochemistry (Paraffin) (IHC (P))	See 5 publications below

* Suggested working dilutions are given as a guide only. It is recommended that the user titrate the product for use in their own experiment using appropriate negative and positive controls.

Product specific information

MA5-12557 targets p53 in FACS, ICC/IF, IHC (P), IP, ChIP, and WB applications and shows reactivity with Human and bovine samples. This antibody is not suitable for mouse colon tissue in IHC applications. The MA5-12557 immunogen is recombinant human wild-type p53 protein expressed in E. coli. This antibody was originally validated as part of a Thermo Scientific Cellomics High Content Screening Kit. The antibody sold separately may have slightly different performance and may need to be further optimized for the best results.

Background/Target Information

The tumor suppressor protein, p53, is a sequence specific transcription factor that is activated by cellular stress. p53 mediates cell cycle arrest or apoptosis in response to DNA damage or starvation for pyrimidine nucleotides. p53 is up-regulated in response to stress signals and stimulated to activate transcription of specific genes, resulting in expression of p21waf1 and other proteins involved in G1 or G2/M arrest. The structure of p53 comprises an N-terminal transactivation domain, a central DNA-binding domain, an oligomerisation domain, and a C-terminal regulatory domain. There are various phosphorylation sites on p53, of which the phosphorylation at Ser15 is important for p53 activation and stabilization. p53 has been characterized to play a role in blocking the proliferative action of damaged cells and act as an anticancer agent. Phosphorylation of Ser392 in p53 has been shown to associate with the formation of human tumors. In addition, p53 has also been linked to the effects of aging and oxidative stress and an increase in p53 has been linked to deficits in LTP (Long Term Potentiation) in learning and memory. p53 is found in very low levels in normal cells, however, in a variety of transformed cell lines, it is expressed in high amounts, and believed to contribute to transformation and malignancy. Mutants of p53 that frequently occur in a number of different human cancers fail to bind the consensus DNA binding site, and cause the loss of tumor suppressor activity. Alterations of the TP53 gene occur not only as somatic mutations in human malignancies, but also as germline mutations in some cancer-prone families such as Li-Fraumeni syndrome.

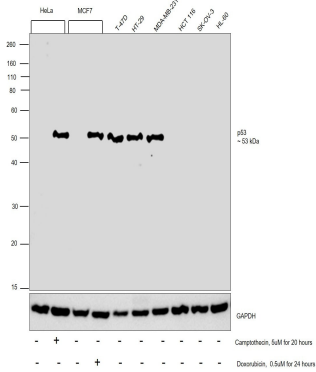
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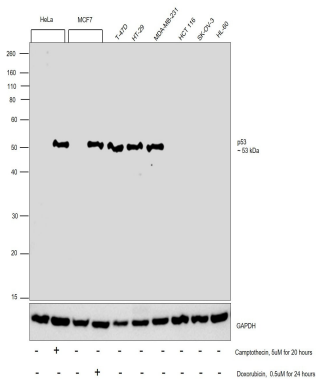
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Product Images For p53 Monoclonal Antibody (DO-7)



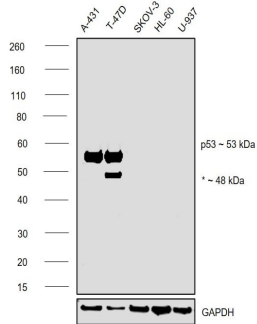
p53 Antibody (MA5-12557)

Antibody specificity was demonstrated by detection of differential basal expression of the target across cell lines owing to their inherent genetic constitution. Relative expression of Cellular tumor antigen p53 was observed in T-47D, HT-29 and MDA-MB-231 which are reported to be positive and not in other cell lines like HCT 116, SK-OV-3 and HL-60 using Anti-p53 Monoclonal Antibody (DO-7) (Product # MA5-12557) in Western Blot. {RE}



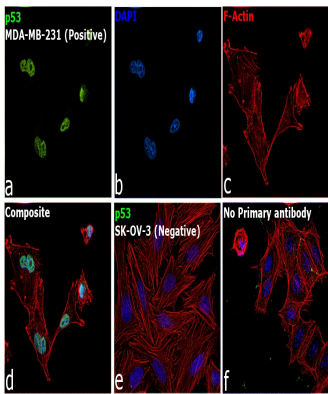
p53 Antibody (MA5-12557) in WB

Western blot was performed using Anti-p53 Monoclonal Antibody (DO-7) (Product # MA5-12557) and a 53 kDa band corresponding to Cellular tumor antigen p53 was observed across cell lines except HCT 116, SK-OV-3 and HL-60 which are reported to be negative, also it was observed to be induced upon Camptothecin and Doxorubicin treatment in HeLa and MCF7 respectively. Modified whole cell extracts (1%SDS) (30 µg lysate) of HeLa (Lane 1), HeLa treated with Camptothecin (5uM for 20 hours) (Lane 2), MCF7 (Lane 3), MCF7 treated with Doxorubicin (0.5uM for 24 hours) (Lane 4), T-47D (Lane 5), HT-29 (Lane 6), MDA-MB-231 (Lane 7), HCT 116 (Lane 8), SK-O-V3 (Lane 9) and HL-60 (Lane 10) were electrophoresed using NuPAGE™ 10% Bis-Tris Protein Gel (Product # NP0302BOX). Resolved proteins were then transferred onto a Nitrocellulose membrane (Product # IB23001) by iBlot® 2 Dry Blotting System (Product # IB21001). The blot was probed with the primary antibody (1:1000 dilution) and detected by chemiluminescence with Goat anti-Mouse IgG (H+L) Superclonal™ Recombinant Secondary Antibody, HRP (Product # A28177, 1:4000 dilution) using the iBright FL 1000 (Product # A32752). Chemiluminescent detection was performed using Novex® ECL Chemiluminescent Substrate Reagent Kit (Product # WP20005).



p53 Antibody (MA5-12557) in WB

Western blot was performed using Anti-p53 Monoclonal Antibody (DO-7) (Product # MA5-12557) and a ~53 kDa band corresponding to TP53 was observed across cell lines tested . Whole cell extracts (30 µg lysate) of A-431 (Lane 1), T-47D (Lane 2), SK-O-V3 (Lane 3), HL-60 (Lane 4), U-937 (Lane 5) were electrophoresed using NuPAGE™ 4-12% Bis-Tris Protein Gel (Product # NP0321BOX). Resolved proteins were then transferred onto a nitrocellulose membrane (Product # IB23001) by iBlot® 2 Dry Blotting System (Product # IB21001). The blot was probed with the primary antibody (1:1000) and detected by chemiluminescence with Goat anti-Mouse IgG (H+L) Superclonal™ Recombinant Secondary Antibody, HRP (Product # A28177,1:20000) using the iBright™ FL1500 Imaging System (Product # A44115). Chemiluminescent detection was performed using SuperSignal™ West Atto Ultimate Sensitivity Substrate (Product # A38556).



p53 Antibody (MA5-12557)

Antibody specificity was demonstrated by detection of differential basal expression of the target across cell models owing to their inherent genetic constitution. Immunofluorescence analysis using Anti-p53 Monoclonal Antibody (DO-7) (Product # MA5-12557), shows expression of p53 in MDA-MB-231 when compared to SK-OV-3. {RE}

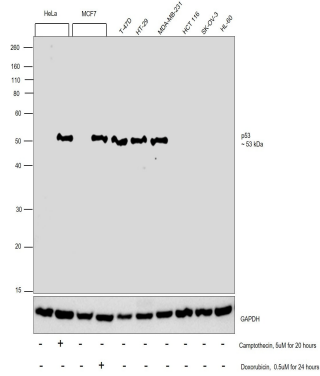
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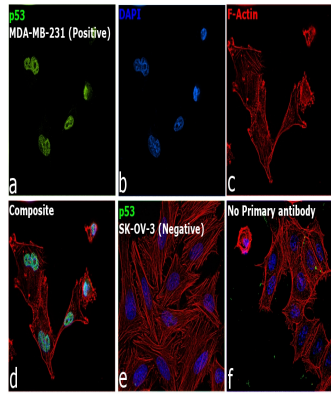
p53 Antibody (MA5-12557)

Altered expression of proteins upon cell treatment demonstrates antibody specificity. Western blot using p53 Monoclonal Antibody (DO-7) (Product # MA5-12557), shows induction of proteins in HeLa and MCF7 on Camptothecin and Doxorubicin treatments respectively. {TM}



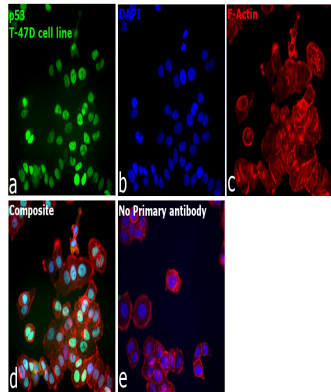
p53 Antibody (MA5-12557) in ICC/IF

Immunofluorescence analysis of Cellular tumor antigen p53 was performed using 70% confluent log phase MDA-MB-231 cells. The cells were fixed with 4% paraformaldehyde for 10 minutes, permeabilized with 0.1% Triton™ X-100 for 15 minutes, and blocked with 2% BSA for 45 minutes at room temperature. The cells were labeled with p53 Monoclonal Antibody (DO-7) (Product # MA5-12557) at 1:100 dilution in 0.1% BSA, incubated at 4 degree celsius overnight and then labeled with Donkey anti-Mouse IgG (H+L) Highly Cross-Adsorbed Secondary Antibody, Alexa Fluor Plus 488 (Product # A32766), (1:2000 dilution), for 45 minutes at room temperature (Panel a: Green). Nuclei (Panel b: Blue) were stained with ProLong™ Diamond Antifade Mountant with DAPI (Product # P36962). F-actin (Panel c: Red) was stained with Rhodamine Phalloidin (Product # R415, 1:300). Panel d represents the merged image showing Nuclear localization. Panel e represents SK-OV-3 cells having no expression of p53. Panel f represents control cells with no primary antibody to assess background. The images were captured at 60X magnification.



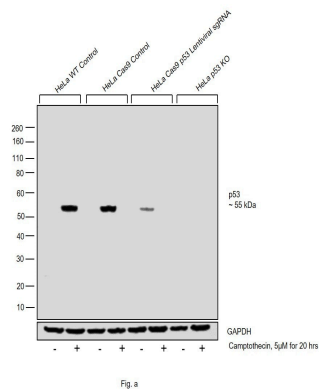
p53 Antibody (MA5-12557) in ICC/IF

Immunofluorescence analysis of TP53 was performed using 70% confluent log phase T-47D cells. The cells were fixed with 4% paraformaldehyde for 10 minutes, permeabilized with 0.1% Triton™ X-100 for 15 minutes, and blocked with 2% BSA for 45 minutes at room temperature. The cells were labeled with p53 Monoclonal Antibody (DO-7) (Product # MA5-12557) at 1:100 dilution in 0.1% BSA, incubated at 4 degree celsius overnight and then labeled with Donkey anti-Mouse IgG (H+L) Highly Cross-Adsorbed Secondary Antibody, Alexa Fluor Plus 488 (Product # A32766), (1:2000), for 45 minutes at room temperature (Panel a: Green). Nuclei (Panel b: Blue) were stained with ProLong™ Diamond Antifade Mountant with DAPI (Product # P36962). F-actin (Panel c: Red) was stained with Rhodamine Phalloidin (Product # R415, 1:300). Panel d represents the merged image showing nuclear localization. Panel e represents control cells with no primary antibody to assess background. The images were captured at 40X magnification.



p53 Antibody (MA5-12557)

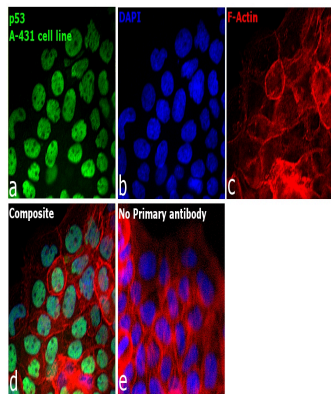
Antibody specificity was demonstrated by CRISPR-Cas9 mediated knockout of target protein. A loss of signal was observed for target protein in p53 KO cell line compared to control cell line using Anti-p53 Monoclonal Antibody (DO-7) (Product # MA5-12557). {KO}



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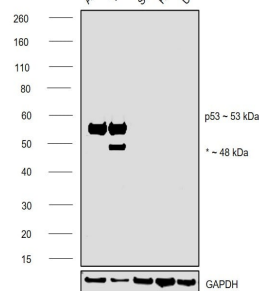


p53 Antibody (MA5-12557) in ICC/IF

Immunofluorescence analysis of TP53 was performed using 70% confluent log phase A-431 cells. The cells were fixed with 4% paraformaldehyde for 10 minutes, permeabilized with 0.1% Triton™ X-100 for 15 minutes, and blocked with 2% BSA for 45 minutes at room temperature. The cells were labeled with p53 Monoclonal Antibody (DO-7) (Product # MA5-12557) at 1:100 dilution in 0.1% BSA, incubated at 4 degree celsius overnight and then labeled with Donkey anti-Mouse IgG (H+L) Highly Cross-Adsorbed Secondary Antibody, Alexa Fluor Plus 488 (Product # A32766), (1:2000), for 45 minutes at room temperature (Panel a: Green). Nuclei (Panel b:Blue) were stained with ProLong™ Diamond Antifade Mountant with DAPI (Product # P36962). F-actin (Panel c: Red) was stained with Rhodamine Phalloidin (Product # R415, 1:300). Panel d represents the merged image showing nuclear localization. Panel e represents control cells with no primary antibody to assess background. The images were captured at 40X magnification.

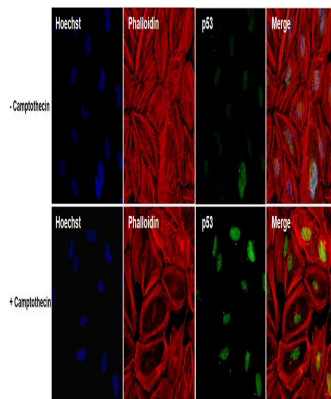
p53 Antibody (MA5-12557)

Antibody specificity was demonstrated by detection of differential basal expression of the target across cell lines owing to their inherent genetic constitution. Relative expression of TP53 was observed as increased expression in A-431 and T-47D as compared to other cell lines using Anti-p53 Monoclonal Antibody (DO-7) (Product # MA5-12557) in Western Blot. {RE}



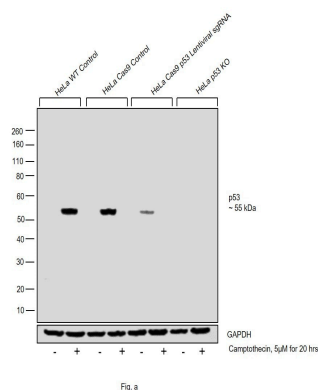
p53 Antibody (MA5-12557) in ICC/IF

Immunofluorescent analysis of p53 (green) in untreated and 5 μ M 20 hours Camptothecin treated HeLa cells. The cells were fixed with 4% paraformaldehyde for 15 minutes at -20c, permeabilized with 0.1% Triton X-100 for 15 minutes, and blocked with 3% BSA for 30 minutes at room temperature. Cells were stained with a p53 mouse monoclonal antibody (Product # MA5-12557) at a dilution of 1:200 in blocking buffer for 1 hour at room temperature, and then incubated with a Goat anti-Mouse IgG (H+L) Secondary Antibody, Alexa Fluor Plus 488 conjugate (Product # A32731) at a dilution of 1:500 for at least 30 minutes at a room temperature in the dark (green). F-actin (red) was stained by Dylight 554 Phalloidin (Product #21834) and nuclei (blue) were stained with Hoechst 33342 (Product # 62249). Images were taken on a Thermo Scientific ToxInsight Instrument at 20X magnification.



p53 Antibody (MA5-12557) in WB

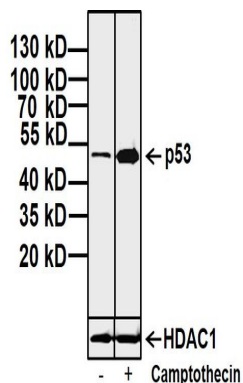
Knockout of p53 was achieved by CRISPR-Cas9 genome editing using LentiArray™ Lentiviral sgRNA (Product # A32042, Assay ID CRISPR718517_LV) and LentiArray Cas9 Lentivirus (Product # A32064). Western blot analysis of p53 was performed by loading 30 μ g of HeLa Wild Type (Lane 1), Treated HeLa Wild type (Lane 2), HeLa Cas9 (Lane 3), Treated HeLa Cas9 (Lane 4), HeLa Cas9 cells transduced with p53 Lentiviral sgRNA (Lane 5), Treated HeLa Cas9 cells transduced with p53 Lentiviral sgRNA (Lane 6), HeLa p53 KO (Lane 7) and Treated HeLa p53 KO (Lane 8) whole cell extracts. The samples were electrophoresed using NuPAGE™ Novex™ 4-12% Bis-Tris Protein Gel (Product # NP0322BOX). Resolved proteins were then transferred onto a nitrocellulose membrane (Product # IB23001) by iBlot® 2 Dry Blotting System (Product # IB21001). The blot was probed with Anti-p53 Monoclonal Antibody (DO-7) (Product # MA5-12557, 1:1,000 dilution) and Goat anti-Mouse IgG (H+L) Superclonal™ Recombinant Secondary Antibody, HRP (Product # A28177, 1:4,000 dilution) using the iBright FL 1000 (Product # A32752). Chemiluminescent detection was performed using SuperSignal™ West Dura Extended Duration Substrate (Product # 34076). Loss of signal upon CRISPR mediated knockout (KO) using the LentiArray™ CRISPR product line confirms that antibody is specific to p53. *Treatment done with 5 μ M Camptothecin for 20 hrs.



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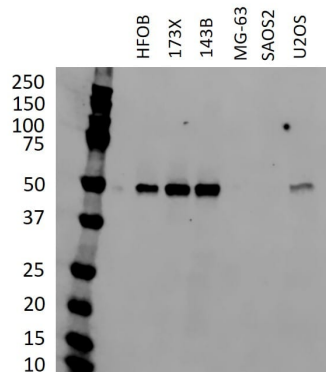
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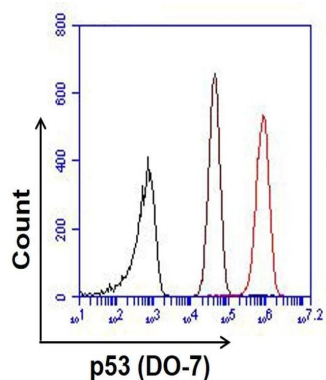
p53 Antibody (MA5-12557) in WB

Western blot analysis of p53 was performed by loading 20 µg HeLa nuclear cell lysate per lane (untreated or treated with 20 µM Camptothecin for 20 hours) and 7 µl of PageRuler Plus Prestained Protein Ladder (Product # 26619) per well onto a 4-20% Tris-Glycine polyacrylamide gel (Product # WT4202BX10). Proteins were transferred to a nitrocellulose membrane using the G2 Blotter (Product # 62288), and blocked with 5% Milk in TBST for 1 hour at room temperature. p53 was detected at ~53 kDa using a p53 mouse monoclonal antibody (Product # MA5-12557) at a dilution of 1:1000 in blocking buffer overnight at 4°C on a rocking platform, followed by a Goat anti-Mouse IgG (H+L) Secondary Antibody, HRP conjugate (Product # 31430) at a dilution of 1:20,000 for at least 30 minutes at room temperature. HDAC1 was detected using a HDAC1 rabbit polyclonal antibody (Product # PA1-860) at a concentration of 1 µg/mL. Chemiluminescent detection was performed using SuperSignal West Pico (Product # 34078).



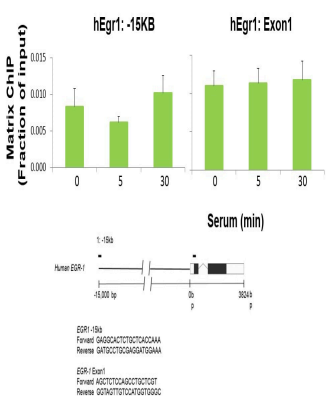
p53 Antibody (MA5-12557) in WB

Western blot analysis of p53 was performed by loading whole-cell lysates of five human osteosarcoma cell lines and non-transformed human osteoblasts (HFOB). p53 was detected at approximately 53 kDa using a p53 monoclonal antibody (Product # MA5-12557) at a dilution of 1:1000 in Blocking Buffer (Product # 37543) overnight at 4°C on a rocking platform, followed by a 1-hour room-temperature incubation of a goat anti-mouse IgG antibody at a dilution of 1:5000. Data courtesy of Thermo Scientific KOL Program.



p53 Antibody (MA5-12557) in Flow

Flow cytometry analysis of p53 was done on HeLa cells. The cells were fixed, permeabilized and stained with a p53 mouse monoclonal antibody (Product # MA5-12557, red histogram) or Mouse IgG2b isotype control (Product # MA5-14447, blue histogram) at a concentration of 10 µg/mL. Black histogram represents negative control unstained cell population. After incubation of the primary antibody on ice for an hour, the cells were stained with a Goat anti-Mouse IgG (H+L) Secondary Antibody, Alexa Fluor Plus 680 conjugate (Product # A32734) at a dilution of 1:50 for at least 30 minutes on ice. A representative 10,000 cells were acquired for each sample.



p53 Antibody (MA5-12557) in ChIP

Multiplex microplate Matrix ChIP has been described in detail (<http://www.ncbi.nlm.nih.gov/pubmed/25959381>). Briefly HTC116 cells were starved followed by addition of serum and samples of cells were cross-linked with formaldehyde after the time points indicated on the x-axis (0, 5 and 30 min). Chromatin was sheared using a Bioruptor and ChIP assays were performed using protein A-coated 96-well polypropylene microplates with 1 µL/100uL well volume of p53 monoclonal antibody (Product # MA5-12557). Quantitative real-time PCRs were performed in quadruplicate using 1-2 µL of DNA with primers to -15kb downstream of Egr1 and exon 1 of Egr1. PCR calibration curves were generated for each primer pair from a dilution series of total human genomic DNA. The PCR primer efficiency curve was fit to cycle threshold (Ct) versus log [genomic DNA concentration] by using an r2 best fit. DNA concentration values for each ChIP and input DNA sample were calculated from their respective average Ct values. Final results are expressed as fraction of input DNA. Schematic representations of Egr1 loci is shown where boxes represent exons (black boxes = translated regions, white boxes = untranslated regions), the zigzag line represents an intron, and the straight line represents upstream sequence. Regions amplified by the primers are represented by black bars. Data courtesy of Dr. Karol Bomsztyk's laboratory.

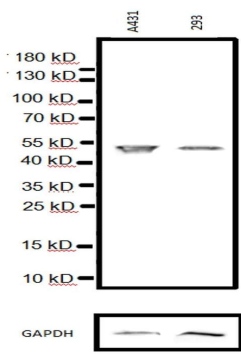
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p53 Antibody (MA5-12557) in WB

Western blot analysis of p53 was performed by loading 20 µg of indicated whole cell lysates per lane and 7 µl of PageRuler Plus Prestained Protein Ladder (Product # 26619) per well onto a 4-20% Tris-Glycine polyacrylamide gel (Product # WT4202BX10). Proteins were transferred to a nitrocellulose membrane using the G2 Blotter (Product # 62288), and blocked with 5% Milk in TBST for 1 hour at room temperature. p53 was detected at ~53 kDa using a p53 mouse monoclonal antibody (Product # MA5-12557) at a dilution of 1:1000 in blocking buffer overnight at 4°C on a rocking platform, followed by a Goat anti-Mouse IgG (H+L) Secondary Antibody, HRP conjugate (Product # 31430) at a dilution of 1:20,000 for at least 30 minutes at room temperature. GAPDH was detected using a GAPDH rabbit polyclonal antibody, HRP conjugate (Product # PA1-987-HRP) at a dilution of 1:1000 overnight at 4°C on a rocking platform. Chemiluminescent detection was performed using SuperSignal West Pico (Product # 34078).



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PubMed References For p53 Monoclonal Antibody (DO-7)

35 Western Blot References

Species / Dilution	Summary
Human / Not Cited	<p>MA5-12557 was used in Western Blotting to show the effect of structurally diverse BET inhibitors on SIRT1 levels is divergent, and the responses might also be cell type-dependent.</p> <p>Gene (May 2020; 741:) "Impact of structurally diverse BET inhibitors on SIRT1." Author(s):Tenhunen J,Kokkola T,Huovinen M,Rahnasto-Rilla M,Lahtela-Kakkonen M PubMed Article URL:http://dx.doi.org/10.1016/j.gene.2020.144558</p>
Mouse / 1:1000	<p>MA5-12557 was used in western blot to investigate the effect of glutathione on chromosome aberrations, cell cycle kinetics, and cell cycle regulator proteins</p> <p>International journal of radiation biology (May 2007; 83: 347) "Influence of glutathione on the induction of chromosome aberrations, delay in cell cycle kinetics and cell cycle regulator proteins in irradiated mouse bone marrow cells." Author(s):Ray S,Chatterjee A PubMed Article URL:http://dx.doi.org/10.1080/09553000701317887</p>
Mouse / 1:500	<p>MA5-12557 was used in western blot to study the role of TGF-beta inhibitors in abolishing the resistance of glioblastoma to ionizing radiation therapy</p> <p>Cancer research (Aug 2012; 72: 4119) "Resistance of glioblastoma-initiating cells to radiation mediated by the tumor microenvironment can be abolished by inhibiting transforming growth factor-." Author(s):Hardee ME,Marciscano AE,Medina-Ramirez CM,Zagzag D,Narayana A,Lonning SM,Barcellos-Hoff MH PubMed Article URL:http://dx.doi.org/10.1158/0008-5472.CAN-12-0546</p>
Human / 1:1000	<p>MA5-12557 was used in Western Blot to indicate that GAS5 participated in the development of OA by regulating the biological behavior of chondrocytes via the miR,34a/Bcl2 axis.</p> <p>Molecular medicine reports (Mar 2020; 21: 1310) "Silencing of longchain noncoding RNA GAS5 in osteoarthritic chondrocytes is mediated by targeting the miR34a/Bcl2 axis." Author(s):Ji Q,Qiao X,Liu Y,Wang D,Yan J PubMed Article URL:http://dx.doi.org/10.3892/mmr.2019.10900</p>
Human / 1:200	<p>MA5-12557 was used in Western Blotting to investigate snake-venom-induced adaptive response at a relatively noncytotoxic concentration (0.01 µg/ml) in two human fibroblast cell lines of different origin, namely WI-38 fetal lung fibroblasts and BJ foreskin fibroblasts.</p> <p>Journal of cellular physiology (May 2019; 234: 6147) "Snake venoms promote stress-induced senescence in human fibroblasts." Author(s):Lewinska A,Bocian A,Petrilla V,Adamczyk-Grochala J,Szymura K,Hendzel W,Kaleniuk E,Hus KK,Petrillova M,Wnuk M PubMed Article URL:http://dx.doi.org/10.1002/jcp.27382</p>
Not Applicable / 1:1000	<p>MA5-12557 was used in western blot to isolate ubiquitin functional conformations that act as potent and selective USP7 inhibitors</p> <p>Nature chemical biology (Jan 2013; 9: 51) "Conformational stabilization of ubiquitin yields potent and selective inhibitors of USP7." Author(s):Zhang Y,Zhou L,Rouge L,Phillips AH,Lam C,Liu P,Sandoval W,Helgason E,Murray JM,Wertz IE,Corn JE PubMed Article URL:http://dx.doi.org/10.1038/nchembio.1134</p>
Human / 1:500	<p>MA5-12557 was used in western blot to study the effects of 2-methoxyestradiol on the growth and apoptosis of human hepatocellular carcinoma cells</p> <p>Journal of gastroenterology and hepatology (Jul 2006; 21: 1207) "A very low toxic agent induces apoptosis and reduces growth of human hepatocellular carcinoma cells." Author(s):Schumacher G,Scheunert S,Rueggeberg A,Bachem MG,Nussler AK,Spinelli A,Mukhopadhyay T,Pratschke J,Neuhaus P PubMed Article URL:http://dx.doi.org/10.1111/j.1440-1746.2006.04327.x</p>
Mouse / 1:200	<p>MA5-12557 was used in Western Blotting to study the effect of AZD7762 on mammary tumour cells and bone cells.</p> <p>International journal of oncology (Sep 2018; 53: 1001) "Effects of a checkpoint kinase inhibitor, AZD7762, on tumor suppression and bone remodeling." Author(s):Wang L,Wang Y,Chen A,Jalali A,Liu S,Guo Y,Na S,Nakshatri H,Li BY,Yokota H PubMed Article URL:http://dx.doi.org/10.3892/ijo.2018.4481</p>

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	<p>MA5-12557 was used in western blot to study the enhancement of cisplatin-induced cytotoxicity and apoptosis in head and neck cancer by a p53-reactivating small-molecule</p>
Human / Not Cited	<p>Cancer letters (Dec 2012; 325: 35) "The p53-reactivating small-molecule RITA enhances cisplatin-induced cytotoxicity and apoptosis in head and neck cancer." Author(s):Roh JL,Ko JH,Moon SJ,Ryu CH,Choi JY,Koch WM PubMed Article URL:http://dx.doi.org/10.1016/j.canlet.2012.05.020</p>
	<p>MA5-12557 was used in western blot to study the mechanism by which DeltaNp63alpha is targeted for proteasomal degradation in head and neck squamous cell carcinoma cells</p>
Human / Not Cited	<p>Cell cycle (Georgetown, Tex.) (Oct 2004; 3: 1285) "RACK1 and stratifin target DeltaNp63alpha for a proteasome degradation in head and neck squamous cell carcinoma cells upon DNA damage." Author(s):Fomenkov A,Zangen R,Huang YP,Osada M,Guo Z,Fomenkov T,Trink B,Sidransky D,Ratovitski EA PubMed Article URL:http://dx.doi.org/10.4161/cc.3.10.1155</p>
	<p>MA5-12557 was used in western blot to study the effect of p53 status on the sensitization of human tumour cells to hyperthermia by a plant flavonol</p>
Human / Not Cited	<p>International journal of hyperthermia : the official journal of European Society for Hyperthermic Oncology, North American Hyperthermia Group (Aug 2008; 24: 415) "p53 status-dependent sensitization of human tumour cells to hyperthermia by plant flavonol." Author(s):Hamamoto T,Suzuki K,Yamauchi M,Kodama S,Sasaki H,Watanabe M PubMed Article URL:http://dx.doi.org/10.1080/02656730802064613</p>
	<p>MA5-12557 was used in western blot to study the mechanism by which axin downregulatesTCF-4 transcription and inhibits lung cancer cell proliferation and invasiveness</p>
Human / 1:200	<p>Molecular cancer (Feb 2010; 9:) "Axin downregulates TCF-4 transcription via beta-catenin, but not p53, and inhibits the proliferation and invasion of lung cancer cells." Author(s):Yang LH,Xu HT,Han Y,Li QC,Liu Y,Zhao Y,Yang ZQ,Dong QZ,Miao Y,Dai SD,Wang EH PubMed Article URL:http://dx.doi.org/10.1186/1476-4598-9-25</p>
	<p>MA5-12557 was used in western blot to study the growth factor requirements and basal phenotype of an immortalized mammary epithelial cell line</p>
Human / Not Cited	<p>Cancer research (Jan 2002; 62: 89) "Growth factor requirements and basal phenotype of an immortalized mammary epithelial cell line." Author(s):DiRenzo J,Signoretti S,Nakamura N,Rivera-Gonzalez R,Sellers W,Loda M,Brown M PubMed Article URL:http://www.ncbi.nlm.nih.gov/pubmed/11782364</p>
	<p>MA5-12557 was used in western blot to study the molecular and cellular biology of high nitric oxide-adapted human tongue squamous cell carcinoma cell lines</p>
Human / Not Cited	<p>Tumour biology : the journal of the International Society for Oncodevelopmental Biology and Medicine (Feb 2011; 32: 87) "Part II. Initial molecular and cellular characterization of high nitric oxide-adapted human tongue squamous cell carcinoma cell lines." Author(s):Tarjan G,Haines GK,Vesper BJ,Xue J,Altman MB,Yarmolyuk YR,Khurram H,Elseth KM,Roeske JC,Aydogan B,Radosevich JA PubMed Article URL:http://dx.doi.org/10.1007/s13277-010-0102-0</p>
	<p>MA5-12557 was used in western blot to study the role of MDM2 antagonists in activating p53 and synergizing with genotoxic drugs in B-cell chronic lymphocytic leukemia cells</p>
Human / Not Cited	<p>Blood (May 2006; 107: 4109) "MDM2 antagonists activate p53 and synergize with genotoxic drugs in B-cell chronic lymphocytic leukemia cells." Author(s):Coll-Mulet L,Iglesias-Serret D,Santidrián AF,Cosíalls AM,de Frias M,Castaño E,Campàs C,Barragán M,de Sevilla AF,Domingo A,Vassilev LT,Pons G,Gil J PubMed Article URL:http://dx.doi.org/10.1182/blood-2005-08-3273</p>
	<p>MA5-12557 was used in western blot to study the different molecular phenotype and tumorigenicity of 5 breast cancer cell lines derived from a single patient</p>
Human / Not Cited	<p>PloS one (Jul 2013; 8:) "Multiple breast cancer cell-lines derived from a single tumor differ in their molecular characteristics and tumorigenic potential." Author(s):Mosoyan G,Nagi C,Marukian S,Teixeira A,Simonian A,Resnick-Silverman L,DiFeo A,Johnston D,Reynolds SR,Roses DF,Mosoian A PubMed Article URL:http://dx.doi.org/10.1371/journal.pone.0055145</p>

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	MA5-12557 was used in Western Blot to study the effect of combined inhibition of AXL and HDACs in diffuse intrinsic pontine glioma cells.
Mouse / 1:1000	<p>Clinical cancer research : an official journal of the American Association for Cancer Research (Jul 2020; 26: 3319)</p> <p>"Combined Therapy of AXL and HDAC Inhibition Reverses Mesenchymal Transition in Diffuse Intrinsic Pontine Glioma."</p> <p>Author(s):Meel MH,de Gooijer MC,Metselaar DS,Sewing ACP,Zwaan K,Waranecki P,Breur M,Buil LCM,Lagerweij T,Wedekind LE,Twisk JWR,Koster J,Hashizume R,Raabe EH,Montero Carcaboso Á,Bugiani M,Phoenix TN,van Tellingen O,van Vuurden DG,Kaspers GJL,Hulleman E</p> <p>PubMed Article URL:http://dx.doi.org/10.1158/1078-0432.CCR-19-3538</p>
Human / 1:500	<p>MA5-12557 was used in Western Blotting to demonstrate a new approach for identifying tumorigenic drivers from genomic regions highly susceptible to ER-related chromothripsis.</p> <p>BMC medical genomics (May 2020; 13:)</p> <p>"ER-related chromothripsis enhances concordant gene transcription on chromosome 17q11.1-q24.1 in luminal breast cancer."</p> <p>Author(s):Lin CL,Tan X,Chen M,Kusi M,Hung CN,Chou CW,Hsu YT,Wang CM,Kirma N,Chen CL,Lin CH,Lathrop KI,Elledge R,Kaklamani VG,Mitsuya K,Huang TH</p> <p>PubMed Article URL:http://dx.doi.org/10.1186/s12920-020-0729-7</p>
Human / 1:1000	<p>MA5-12557 was used in Western Blotting to investigate the effects of a rotating magnetic field (RMF; 0.2 T, 4 Hz) on the growth of human umbilical vein endothelial cells (HUVECs) and Caenorhabditis elegans.</p> <p>Aging (Nov 2019; 11: 10385)</p> <p>"Rotating magnetic field delays human umbilical vein endothelial cell aging and prolongs the lifespan of Caenorhabditis elegans."</p> <p>Author(s):Xu J,Liu K,Chen T,Zhan T,Ouyang Z,Wang Y,Liu W,Zhang X,Sun Y,Xu G,Wang X</p> <p>PubMed Article URL:http://dx.doi.org/10.18632/aging.102466</p>
Mouse / Not Cited	<p>MA5-12557 was used in western blot to study the mechanism by which p53 loss enhances IKKbeta catalytic activity</p> <p>Proceedings of the National Academy of Sciences of the United States of America (Mar 2009; 106: 3431)</p> <p>"Loss of p53 enhances catalytic activity of IKKbeta through O-linked beta-N-acetyl glucosamine modification."</p> <p>Author(s):Kawauchi K,Araki K,Tobiome K,Tanaka N</p> <p>PubMed Article URL:http://dx.doi.org/10.1073/pnas.0813210106</p>
Mouse / Not Cited	<p>MA5-12557 was used in western blot to study the role of glutathione levels in the genotoxic effects of raw betel-nut extract in mammalian cells</p> <p>Mutation research (Jul 2003; 538: 1)</p> <p>"Genotoxic effect of raw betel-nut extract in relation to endogenous glutathione levels and its mechanism of action in mammalian cells."</p> <p>Author(s):Kumpawat K,Deb S,Ray S,Chatterjee A</p> <p>PubMed Article URL:http://dx.doi.org/10.1016/s1383-5718(03)00048-2</p>
Human / 1:1000	<p>MA5-12557 was used in western blot to study the proteomics of selected prognostic factors in breast cancer</p> <p>Proteomics (Mar 2004; 4: 784)</p> <p>"Proteomic analysis of selected prognostic factors of breast cancer."</p> <p>Author(s):Roberts K,Bhatia K,Stanton P,Lord R</p> <p>PubMed Article URL:http://dx.doi.org/10.1002/pmic.200300633</p>
Human / Not Cited	<p>MA5-12557 was used in western blot to study the ability of prodigiosin to induce apoptosis in primary B- and T-cells from patients with B-cell chronic lymphocytic leukemia</p> <p>Leukemia (Apr 2003; 17: 746)</p> <p>"Prodigiosin induces apoptosis of B and T cells from B-cell chronic lymphocytic leukemia."</p> <p>Author(s):Campàs C,Dalmáu M,Montaner B,Barragán M,Bellosillo B,Colomer D,Pons G,Pérez-Tomás R,Gil J</p> <p>PubMed Article URL:http://dx.doi.org/10.1038/sj.leu.2402860</p>
Human / Not Cited	<p>MA5-12557 was used in Western Blotting to investigate whether the anti-HIV effect of lactobacilli is mediated by extracellular vesicles released by these bacteria.</p> <p>Nature communications (Dec 2019; 10:)</p> <p>"Extracellular vesicles from symbiotic vaginal lactobacilli inhibit HIV-1 infection of human tissues."</p> <p>Author(s):Ñahui Palomino RA,Vanpouille C,Laghi L,Parolin C,Melikov K,Backlund P,Vitali B,Margolis L</p> <p>PubMed Article URL:http://dx.doi.org/10.1038/s41467-019-13468-9</p>
Human / 1:400	<p>MA5-12557 was used in western blot to study the mechanism by which Delta-like 1 contributes to cell growth in hepatocellular carcinoma</p> <p>Liver international : official journal of the International Association for the Study of the Liver (May 2010; 30: 703)</p> <p>"Delta-like 1 contributes to cell growth by increasing the interferon-inducible protein 16 expression in hepatocellular carcinoma."</p> <p>Author(s):Yu F,Hao X,Zhao H,Ge C,Yao M,Yang S,Li J</p> <p>PubMed Article URL:http://dx.doi.org/10.1111/j.1478-3231.2010.02214.x</p>

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Human / Not Cited	Haematologica (Dec 2009; 94: 1698) "Akt inhibitors induce apoptosis in chronic lymphocytic leukemia cells." Author(s):de Frias M,Iglesias-Serret D,Cosialls AM,Coll-Mulet L,Santidrián AF,González-Gironès DM,de la Banda E,Pons G,Gil J PubMed Article URL: http://dx.doi.org/10.3324/haematol.2008.004028
	MA5-12557 was used in western blot to investigate the role of CDK2 in the MYCN over-expressing cancer cells
Human / Not Cited	Proceedings of the National Academy of Sciences of the United States of America (Aug 2009; 106: 12968) "Inactivation of CDK2 is synthetically lethal to MYCN over-expressing cancer cells." Author(s):Molenaar JJ,Ebus ME,Geerts D,Koster J,Lamers F,Valentijn LJ,Westerhout EM,Versteeg R,Caron HN PubMed Article URL: http://dx.doi.org/10.1073/pnas.0901418106
	MA5-12557 was used in western blot to study the mechanism by which Smad3 deletion confers radioprotection in primary dermal fibroblasts
Mouse / Not Cited	Journal of dermatological science (Oct 2007; 48: 35) "Absence of Smad3 confers radioprotection through modulation of ERK-MAPK in primary dermal fibroblasts." Author(s):Arany PR,Flanders KC,DeGraff W,Cook J,Mitchell JB,Roberts AB PubMed Article URL: http://dx.doi.org/10.1016/j.jdermsci.2007.05.012
	MA512557 was used in western blot to develop a method to examine protein oligomerization in cells using a single electrophoresis gel
Human / Not Cited	Analytical and bioanalytical chemistry (Feb 2016; 408: 1715) "Tris-acetate polyacrylamide gradient gel electrophoresis for the analysis of protein oligomerization." Author(s):Cubillos-Rojas M,Schneider T,Sánchez-Tena S,Bartrons R,Ventura F,Rosa JL PubMed Article URL: http://dx.doi.org/10.1007/s00216-015-9283-0
	MA5-12557 was used in western blot to assess links between telomere-dysfunction and centrosome defects in early breast carcinogenesis
Not Applicable / 1:1000	Oncotarget (Sep 2015; 6: 28238) "Centrosome aberrations in human mammary epithelial cells driven by cooperative interactions between p16INK4a deficiency and telomere-dependent genotoxic stress." Author(s):Dominguez D,Feijoo P,Bernal A,Ercilla A,Agell N,Genescà A,Tusell L PubMed Article URL: http://dx.doi.org/10.18632/oncotarget.4958
	MA5-12557 was used in western blot to report that p53 and DeltaNp63alpha are transcriptional partners for SMAD proteins.
Human / Not Cited	Molecular cancer research : MCR (Apr 2015; 13: 732) "p53 and Np63 Coregulate the Transcriptional and Cellular Response to TGF and BMP Signals." Author(s):Balboni AL,Cherukuri P,Ung M,DeCastro AJ,Cheng C,DiRenzo J PubMed Article URL: http://dx.doi.org/10.1158/1541-7786.MCR-14-0152-T
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Human / Not Cited	The EMBO journal (May 2008; 27: 1368) "A synthetic lethal siRNA screen identifying genes mediating sensitivity to a PARP inhibitor." Author(s):Turner NC,Lord CJ,Iorns E,Brough R,Swift S,Elliott R,Rayter S,Tutt AN,Ashworth A PubMed Article URL: http://dx.doi.org/10.1038/emboj.2008.61
	MA5-12557 was used in western blot to study the ability of acadesine to activate AMPK and promote apoptosis in B-cells but not in T-cells from patients with B-cell chronic lymphocytic leukemia
Human / Not Cited	Blood (May 2003; 101: 3674) "Acadesine activates AMPK and induces apoptosis in B-cell chronic lymphocytic leukemia cells but not in T lymphocytes." Author(s):Campàs C,Lopez JM,Santidrián AF,Barragán M,Bellosillo B,Colomer D,Gil J PubMed Article URL: http://dx.doi.org/10.1182/blood-2002-07-2339
	MA5-12557 was used in western blot to study the feasibility of using a random peptide microarray to map antibody epitopes
Human / Not Cited	Molecular & cellular proteomics : MCP (Mar 2011; 10:) "Exploring antibody recognition of sequence space through random-sequence peptide microarrays." Author(s):Halperin RF,Stafford P,Johnston SA PubMed Article URL: http://dx.doi.org/10.1074/mcp.M110.000786

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	MA5-12557 was used in western blot to study the interaction of cyclin G2 with DNA damage responses and the role of cyclin G2 in robust cell cycle arrest following doxorubicin treatment
Human / 1:1000	The Journal of biological chemistry (Jun 2012; 287: 22838) "Elevated cyclin G2 expression intersects with DNA damage checkpoint signaling and is required for a potent G2 /M checkpoint arrest response to doxorubicin." Author(s): Zimmermann M, Arachchige-Don AS, Donaldson MS, Dallapiazza RF, Cowan CE, Horne MC PubMed Article URL: http://dx.doi.org/10.1074/jbc.M112.376855
97 Immunohistochemistry References	
Species / Dilution	Summary
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Human / 1:50	Anticancer research (Sep 2005; 25: 2385) "Prognostic role of p53 in stage III non-small cell lung cancer." Author(s): Berghmans T, Mascaux C, Martin B, Ninane V, Sculier JP PubMed Article URL: http://www.ncbi.nlm.nih.gov/pubmed/16080465
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Human / 1:300	Archives of gynecology and obstetrics (Nov 2011; 284: 1231) "Investigation of galectin-3 and heparanase in endometrioid and serous carcinomas of the endometrium and correlation with known predictors of survival." Author(s): Ege CB, Akbulut M, Zekiolu O, Ozdemir N PubMed Article URL: http://dx.doi.org/10.1007/s00404-010-1766-9
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Human / 1:1000	Human pathology (Oct 2010; 41: 1475) "High levels of p53 expression correlate with DNA aneuploidy in (pre)malignancies of the vulva." Author(s): van der Avoort IA, van de Nieuwenhof HP, Otte-Höller I, Nirmala E, Bulten J, Massuger LF, van der Laak JA, Slootweg PJ, de Hullu JA, van Kempen LC PubMed Article URL: http://dx.doi.org/10.1016/j.humpath.2009.12.015
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Human / Not Cited	Pathology, research and practice (Aug 2008; 204: 367) "Apocrine carcinomas of the breast in Turkish women: hormone receptors, c-erbB-2 and p53 immunoexpression." Author(s): Kaya H, Bozkurt SU, Erbarut I, Djamgoz MB PubMed Article URL: http://dx.doi.org/10.1016/j.prp.2008.01.012
	MA5-12557 was used in immunohistochemistry to report on a case of myointimoma of the glans penis
Human / Not Cited	Pathology international (Mar 2007; 57: 158) "Myointimoma of the glans penis." Author(s): Vardar E, Gunlusoy B, Arslan M, Kececi S PubMed Article URL: http://dx.doi.org/10.1111/j.1440-1827.2006.02074.x

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Human / 1:50	Yonsei medical journal (Aug 2007; 48: 694) "Prognostic implications of cyclin B1, p34cdc2, p27(Kip1) and p53 expression in gastric cancer." Author(s):Kim DH PubMed Article URL: http://dx.doi.org/10.3349/ymj.2007.48.4.694
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Human / 1:1000	Virchows Archiv : an international journal of pathology (Jul 2016; 469: 61) "Identical TP53 mutations in pelvic carcinosarcomas and associated serous tubal intraepithelial carcinomas provide evidence of their clonal relationship." Author(s):Ardighieri L,Mori L,Conzadori S,Bugatti M,Falchetti M,Donzelli CM,Ravaggi A,Odicino FE,Facchetti F PubMed Article URL: http://dx.doi.org/10.1007/s00428-016-1933-x
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Human / 1:100	Pathology international (Jul 2002; 52: 463) "Impact of p53 and Ki-67 in predicting recurrence and progression of superficial (pTa and pT1) urothelial cell carcinomas of urinary bladder." Author(s):Kilicli-Camur N,Kilicaslan I,Gulluoglu MG,Esen T,Uysal V PubMed Article URL: http://dx.doi.org/10.1046/j.1440-1827.2002.01371.x

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Human / 1:500	<p>Neuropathology and applied neurobiology (Feb 2014; 40: 205)</p> <p>"Loss of FUBP1 expression in gliomas predicts FUBP1 mutation and is associated with oligodendroglial differentiation, IDH1 mutation and 1p/19q loss of heterozygosity."</p> <p>Author(s):Baumgarten P,Harter PN,Tönjes M,Capper D,Blank AE,Sahm F,von Deimling A,Kolluru V,Schwamb B,Rabenhorst U,Starzetz T,Kögel D,Rieker RJ,Plate KH,Ohgaki H,Radlwimmer B,Zörnig M,Mittelbronn M</p> <p>PubMed Article URL:http://dx.doi.org/10.1111/nan.12088</p>
Human / 1:250	<p>MA5-12557 was used in immunohistochemistry to investigate the relation between endometrial intraepithelial carcinoma and serous ovarian carcinoma</p> <p>International journal of gynecological cancer : official journal of the International Gynecological Cancer Society (Mar 2012; 22: 457)</p> <p>"Concurrent endometrial intraepithelial carcinoma (EIC) and serous ovarian cancer: can EIC be seen as the precursor lesion?"</p> <p>Author(s):Roelofsen T,van Kempen LC,van der Laak JA,van Ham MA,Bulten J,Massuger LF</p> <p>PubMed Article URL:http://dx.doi.org/10.1097/IGC.0b013e3182434a81</p>
Human / 1:2000	<p>MA5-12557 was used in immunohistochemistry to study the correlation of GABA-A subunit expression with tumor histology and clinical outcome in human glioma</p> <p>PloS one (Jan 2013; 7:)</p> <p>"GABA-A channel subunit expression in human glioma correlates with tumor histology and clinical outcome."</p> <p>Author(s):Smits A,Jin Z,Elsir T,Pedder H,Nistér M,Alafuzoff I,Dimberg A,Edqvist PH,Pontén F,Aronica E,Birnir B</p> <p>PubMed Article URL:http://dx.doi.org/10.1371/journal.pone.0037041</p>
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Human / Not Cited	<p>MA5-12557 was used in immunohistochemistry to investigate esophageal function and its relationship with gastroesophageal reflux disease and Barrett esophagus in patients with cervical inlet patch</p> <p>Diseases of the esophagus : official journal of the International Society for Diseases of the Esophagus (Aug 2012; 25: 498)</p> <p>"Detailed esophageal function and morphological analysis shows high prevalence of gastroesophageal reflux disease and Barrett's esophagus in patients with cervical inlet patch."</p> <p>Author(s):Rosztóczy A,Izbéki F,Németh IB,Dulic S,Vadászi K,Róka R,Gecse K,Gyökeres T,Lázár G,Tiszlavicz L,Wittmann T</p> <p>PubMed Article URL:http://dx.doi.org/10.1111/j.1442-2050.2011.01281.x</p>
Human / 1:1000	<p>MA5-12557 was used in immunohistochemistry to study p53 and a number of other potential markers at the genetic and immunohistochemical levels in ovarian high grade adenocarcinoma patients</p> <p>Pathology international (May 2013; 63: 252)</p> <p>"Genetic alteration and immunohistochemical staining patterns of ovarian high-grade serous adenocarcinoma with special emphasis on p53 immnnostaining pattern."</p> <p>Author(s):Lee SH,Kim H,Kim WY,Han HS,Lim SD,Kim WS,Kim S,Hwang TS</p> <p>PubMed Article URL:http://dx.doi.org/10.1111/pin.12060</p>
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	MA5-12557 was used in immunohistochemistry to study tumor heterogeneity and progression in early rectal cancer
Human / 1:1000	Clinical cancer research : an official journal of the American Association for Cancer Research (Feb 2008; 14: 772) "Progression and tumor heterogeneity analysis in early rectal cancer." Author(s):Lips EH,van Eijk R,de Graaf EJ,Doornebosch PG,de Miranda NF,Oosting J,Karsten T,Eilers PH,Tollenaar RA, van Wezel T,Morreau H PubMed Article URL: http://dx.doi.org/10.1158/1078-0432.CCR-07-2052
Human / 1:50	MA5-12557 was used in immunohistochemistry to investigate the expression of some histopathological and immunohistochemical markers in sudden cardiac death
	Romanian journal of morphology and embryology = Revue roumaine de morphologie et embryologie (Dec 2008; 49: 315) "Immunophenotypical pleomorphism expression in sudden cardiac death." Author(s):Ceauu M,Curc C,Ardeleanu C,Dermengiu D PubMed Article URL: http://www.ncbi.nlm.nih.gov/pubmed/18758635
Human / 1:250	MA5-12557 was used in immunohistochemistry to perform a detailed analysis of the presence of lesions in the Müllerian ducts of patients with serous ovarian cancer
	Modern pathology : an official journal of the United States and Canadian Academy of Pathology, Inc (Jul 2014; 27: 1002) "Müllerian precursor lesions in serous ovarian cancer patients: using the SEE-Fim and SEE-End protocol." Author(s):Mingels MJ,van Ham MA,de Kievit IM,Snijders MP,van Tilborg AA,Bulten J,Massuger LF PubMed Article URL: http://dx.doi.org/10.1038/modpathol.2013.212
Human / 1:50	MA5-12557 was used in immunohistochemistry to study the expression of p53 and Mdm2 in the skin of patients with vitiligo and the significance for non-melanoma skin cancer
	Acta dermatovenerologica Croatica : ADC (Jun 2014; 21: 71) "Vitiligo: is it grace or curse?" Author(s):Bakry OA,Hammam MA,Abdel Wahed MM PubMed Article URL: http://www.ncbi.nlm.nih.gov/pubmed/24001413
Human / 1:25	MA5-12557 was used in immunohistochemistry to report on a case of primary squamous carcinoma of the salpinx associated with serous carcinoma of the omentum
	International journal of gynecological pathology : official journal of the International Society of Gynecological Pathologists (Nov 2010; 29: 533) "A Unique case of primary squamous carcinoma of the salpinx associated with serous carcinoma of the omentum: a pathological and molecular study." Author(s):Giordano G,D'Adda T,Brigati F,Lombardi M,Raboni S PubMed Article URL: http://dx.doi.org/10.1097/PGP.0b013e3181e8ae3d
Human / 1:150	MA5-12557 was used in immunohistochemistry to study the association between p16, p53 and Ki-67 protein expression and progression of epithelial dysplasia of the oral cavity
	Anticancer research (Dec 2008; 28: 2535) "Expression of p16, p53 and Ki-67 proteins in the progression of epithelial dysplasia of the oral cavity." Author(s):Angiero F,Berenzi A,Benetti A,Rossi E,Del Sordo R,Sidoni A,Stefani M,Dessy E PubMed Article URL: http://www.ncbi.nlm.nih.gov/pubmed/19035275
Human / Not Cited	MA5-12557 was used in immunohistochemistry to report on a case of cerebellar liponeurocytoma/lipidized medulloblastoma
	Journal of neuro-oncology (Jan 2005; 71: 53) "Cerebellar liponeurocytoma/lipidized medulloblastoma." Author(s):Aker FV,Ozkara S,Eren P,Peker O,Armaan S,Hakan T PubMed Article URL: http://dx.doi.org/10.1007/s11060-004-9172-4
Human / 1:50	MA5-12557 was used in immunohistochemistry to examine the diagnostic significance of p53 protein in the tumorigenesis in biliopancreatic tree
	Digestive diseases and sciences (Apr 2009; 54: 789) "Immunocytochemical assessment of p53 protein to detect malignancy in increased cell-yield brush cytology from the biliopancreatic tree." Author(s):Villanacci V,Cestari R,Giulini S,Cengia P,Missale G,Berenzi A,Rossi E,Bonardi M,Baiocchi L,Bassotti G PubMed Article URL: http://dx.doi.org/10.1007/s10620-008-0431-7
Human / Not Cited	MA5-12557 was used in immunohistochemistry to study the prognostic value of different PIK3CA gene mutations in breast cancer
	Clinical cancer research : an official journal of the American Association for Cancer Research (Oct 2007; 13: 6064) "Different prognostic roles of mutations in the helical and kinase domains of the PIK3CA gene in breast carcinomas." Author(s):Barbareschi M,Buttitta F,Felicioni L,Cotrupi S,Barassi F,Del Grammastro M,Ferro A,Dalla Palma P,Galligioni E, Marchetti A PubMed Article URL: http://dx.doi.org/10.1158/1078-0432.CCR-07-0266

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	MA5-12557 was used in immunohistochemistry to compare expression of Maspin, VEGF, and p53 between carcinomas
Human / 1:100	<p>Pathology oncology research : POR (Dec 2010; 16: 553)</p> <p>"Maspin, VEGF and p53 expression in small biopsies of primary advanced lung cancer and relationship with clinicopathologic parameters."</p> <p>Author(s):Bircan A,Bircan S,Kapucuoglu N,Songur N,Ozturk O,Akkaya A</p> <p>PubMed Article URL:http://dx.doi.org/10.1007/s12253-010-9259-5</p>
Human / Not Cited	<p>MA5-12557 was used in immunohistochemistry to study the inhibitory effect of 5-FU and selenium on the growth of colorectal cancer cells and the role of caspases</p> <p>Anticancer research (Mar 2009; 28: 3579)</p> <p>"Role of caspases in 5-FU and selenium-induced growth inhibition of colorectal cancer cells."</p> <p>Author(s):Thant AA,Wu Y,Lee J,Mishra DK,Garcia H,Koeffler HP,Vadgama JV</p> <p>PubMed Article URL:http://www.ncbi.nlm.nih.gov/pubmed/19189638</p>
Human / Not Cited	<p>MA5-12557 was used in immunohistochemistry to investigate the efficacy of p53 therapy in Li-Fraumeni syndrome</p> <p>Molecular cancer therapeutics (May 2007; 6: 1478)</p> <p>"p53 therapy in a patient with Li-Fraumeni syndrome."</p> <p>Author(s):Senzer N,Nemunaitis J,Nemunaitis M,Lamont J,Gore M,Gabra H,Eeles R,Sodha N,Lynch FJ,Zumstein LA, Menander KB,Sobol RE,Chada S</p> <p>PubMed Article URL:http://dx.doi.org/10.1158/1535-7163.MCT-07-0125</p>
Human / Not Cited	<p>MA5-12557 was used in immunohistochemistry to study the role of fragile histidine triad protein in non-small cell lung cancer and its correlation with tumor proliferation</p> <p>The European respiratory journal (May 2003; 21: 753)</p> <p>"Fragile histidine triad protein expression in nonsmall cell lung cancer and correlation with Ki-67 and with p53."</p> <p>Author(s):Mascaux C,Martin B,Verdebout JM,Meert AP,Ninane V,Sculier JP</p> <p>PubMed Article URL:http://dx.doi.org/10.1183/09031936.03.00090202</p>
Human / Not Cited	<p>MA5-12557 was used in immunohistochemistry to study the molecular determinants of the response to radiotherapy in oral carcinoma</p> <p>International journal of cancer (May 2004; 109: 710)</p> <p>"p53, p16 and cyclin D1: molecular determinants of radiotherapy treatment response in oral carcinoma."</p> <p>Author(s):Jayasurya R,Francis G,Kannan S,Lekshminarayanan K,Nalinakumari KR,Abraham T,Abraham EK,Nair MK</p> <p>PubMed Article URL:http://dx.doi.org/10.1002/ijc.20042</p>
Human / 1:50	<p>MA5-12557 was used in immunohistochemistry to report on four cases of chordoid glioma of the third ventricle</p> <p>Neuropathology : official journal of the Japanese Society of Neuropathology (Apr 2013; 33: 134)</p> <p>"Chordoid glioma of the third ventricle: four cases including one case with papillary features."</p> <p>Author(s):Ni HC,Piao YS,Lu DH,Fu YJ,Ma XL,Zhang XJ</p> <p>PubMed Article URL:http://dx.doi.org/10.1111/j.1440-1789.2012.01333.x</p>
Human / 1:100	<p>MA5-12557 was used in immunohistochemistry to study the value of glutamate synthetase expression as both a marker of epilepsy and a prognostic indicator in patients with glioblastoma multiforme</p> <p>Neuro-oncology (May 2013; 15: 618)</p> <p>"Glutamine synthetase expression as a valuable marker of epilepsy and longer survival in newly diagnosed glioblastoma multiforme."</p> <p>Author(s):Rosati A,Poliani PL,Todeschini A,Cominelli M,Medicina D,Cenzato M,Simoncini EL,Magrini SM,Buglione M, Grisanti S,Padovani A</p> <p>PubMed Article URL:http://dx.doi.org/10.1093/neuonc/nos338</p>
Human / 1:100	<p>MA5-12557 was used in immunohistochemistry to study the altered distribution of metaplastic cell types in atrophic gastritic mucosa with endocrine cell lesions</p> <p>The Tohoku journal of experimental medicine (Jan 2004; 202: 13)</p> <p>"Altered distribution of metaplastic Paneth, gastrin and pancreatic acinar cells in atrophic gastritic mucosa with endocrine cell lesions."</p> <p>Author(s):Deveci MS,Deveci G</p> <p>PubMed Article URL:http://dx.doi.org/10.1620/tjem.202.13</p>
Human / Not Cited	<p>MA5-12557 was used in immunohistochemistry to study the diagnostic value of p53, bcl-2 and nm23 in serous ovarian tumors</p> <p>Turk patoloji dergisi (Jan 2011; 27: 38)</p> <p>"P53, bcl-2, and nm23 expressions in serous ovarian tumors: correlation with the clinical and histopathological parameters."</p> <p>Author(s):Arik D,Kulaçolu S</p> <p>PubMed Article URL:http://dx.doi.org/10.5146/tjpath.2010.01045</p>

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	MA512557 was used in immunohistochemistry to describe the mechanism behind the association between single nucleotide polymorphism rs55705857 and glioma development.
Human / Not Cited	Scientific reports (Jun 2016; 6:) "IDH-mutant glioma specific association of rs55705857 located at 8q24.21 involves MYC deregulation." Author(s): Oktay Y, Ülgen E, Can Ö, Akyerli CB, Yüksel , Erdemgil Y, Duras IM, Henegariu OI, Nanni EP, Selevsek N, Grossmann J, Erson-Omay EZ, Bai H, Gupta M, Lee W, Turcan , Özpınar A, Huse JT, Sav MA, Flanagan A, Günel M, Sezerman OU, Yakcer MC, Pamir MN, Özdoğan K PubMed Article URL: http://dx.doi.org/10.1038/srep27569
Human / Not Cited	MA5-12557 was used in immunohistochemistry to report on a case of interdigitating dendritic cell tumor with breast and cervical lymph-node involvement Virchows Archiv : an international journal of pathology (May 2005; 446: 546) "Interdigitating dendritic cell tumor with breast and cervical lymph-node involvement: a case report and review of the literature." Author(s): Uluolu O, Akyürek N, Uner A, Cokun U, Ozdemir A, Gökçora N PubMed Article URL: http://dx.doi.org/10.1007/s00428-005-1209-3
Human / 1:50	MA5-12557 was used in immunohistochemistry to report on a morphological and molecular analysis of two cases of Villoglandular adenocarcinoma of the cervix International journal of gynecological pathology : official journal of the International Society of Gynecological Pathologists (Apr 2007; 26: 199) "Villoglandular adenocarcinoma of the cervix: two new cases with morphological and molecular study." Author(s): Giordano G, D'Adda T, Gnetti L, Merisio C, Gabrielli M, Melpignano M PubMed Article URL: http://dx.doi.org/10.1097/01.pgp.0000228141.01964.e7
Human / Not Cited	MA5-12557 was used in immunohistochemistry to study the utility of transient inhibition of hedgehog/smoothed signaling in vivo for selectively and efficiently induce tumor cell differentiation and apoptosis European journal of dermatology : EJD (Jul 2004; 14: 96) "Induction of the differentiation and apoptosis of tumor cells in vivo with efficiency and selectivity." Author(s): Tabs S, Avci O PubMed Article URL: http://www.ncbi.nlm.nih.gov/pubmed/15196999
Human / 1:200	MA5-12557 was used in immunohistochemistry to investigate the association of Axin expression with non-small-cell lung cancer apoptosis induced by X-radiation International journal of radiation oncology, biology, physics (Oct 2009; 75: 518) "X-radiation induces non-small-cell lung cancer apoptosis by upregulation of Axin expression." Author(s): Han Y, Wang Y, Xu HT, Yang LH, Wei Q, Liu Y, Zhang Y, Zhao Y, Dai SD, Miao Y, Yu JH, Zhang JY, Li G, Yuan XM, Wang EH PubMed Article URL: http://dx.doi.org/10.1016/j.ijrobp.2009.05.040

2 Immunoprecipitation References

Species / Dilution	Summary
Human / Not Cited	MA5-12557 was used in immunoprecipitation and western blot to study the modulation of p53 activity and oligomerization by the E3 ubiquitin ligase HERC2 The Journal of biological chemistry (May 2014; 289: 14782) "The E3 ubiquitin protein ligase HERC2 modulates the activity of tumor protein p53 by regulating its oligomerization." Author(s): Cubillos-Rojas M, Amair-Pinedo F, Peiró-Jordán R, Bartrons R, Ventura F, Rosa JL PubMed Article URL: http://dx.doi.org/10.1074/jbc.M113.527978
Human / 2 µg/mL	MA5-12557 was used in immunoprecipitation to study the role of the p53 target gene FLJ11259/DRAM encoding a novel transmembrane protein Biochimica et biophysica acta (Apr 2007; 1769: 209) "The direct p53 target gene, FLJ11259/DRAM, is a member of a novel family of transmembrane proteins." Author(s): Kerley-Hamilton JS, Pike AM, Hutchinson JA, Freemantle SJ, Spinella MJ PubMed Article URL: http://dx.doi.org/10.1016/j.bbaexp.2007.02.002

4 Immunocytochemistry References

Species / Dilution	Summary
Not Applicable / 1:200	MA5-12557 was used in immunocytochemistry and western blot to analyze the activity of p53 after stimulation with p53-dependent and -independent DNA damaging agents during human herpesvirus-infection PLoS one (Sep 2013; 8:) "Inhibition of p53-dependent, but not p53-independent, cell death by U19 protein from human herpesvirus 6B." Author(s): Kofod-Olsen E, Møller JM, Schleimann MH, Bundgaard B, Bak RO, Øster B, Mikkelsen JG, Hupp T, Höllsberg P PubMed Article URL: http://dx.doi.org/10.1371/journal.pone.0059223

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	MA5-12557 was used in Immunocytochemistry to develop a novel network pharmacology approach based on multi-omics functional data integration to predict how stingray venom disrupts the physiological systems of target animals.
Human / Not Cited	Marine drugs (Dec 2021; 20:) "Stingray Venom Proteins: Mechanisms of Action Revealed Using a Novel Network Pharmacology Approach." Author(s):Kirchhoff KN,Billion A,Voolstra CR,Kremb S,Wilke T,Vilcinskas A PubMed Article URL: http://dx.doi.org/10.3390/md20010027
	MA5-12557 was used in immunocytochemistry to derive and characterize a Wilms' tumour cell line
Mouse / 1:100	International journal of cancer (Nov 2003; 107: 365) "Derivation and characterization of a Wilms' tumour cell line, WiT 49." Author(s):Alami J,Williams BR,Yeger H PubMed Article URL: http://dx.doi.org/10.1002/ijc.11429
	MA5-12557 was used in immunocytochemistry to study the mechanism by which apigenin induces apoptosis in human neuroblastoma cells
Human / Not Cited	Molecular cancer therapeutics (Jan 2005; 4: 1) "Induction of caspase-dependent, p53-mediated apoptosis by apigenin in human neuroblastoma." Author(s):Torkin R,Lavoie JF,Kaplan DR,Yeger H PubMed Article URL: http://www.ncbi.nlm.nih.gov/pubmed/15657348
1 Immunohistochemistry (Frozen) References	
Species / Dilution	Summary
	MA5-12557 was used in Immunohistochemistry (Frozen) to lay the foundation for accurate prognostic and therapeutic stratification of SOC.
Human / Not Cited	Cancer cell (Feb 2020; 37: 226) "The Repertoire of Serous Ovarian Cancer Non-genetic Heterogeneity Revealed by Single-Cell Sequencing of Normal Fallopian Tube Epithelial Cells." Author(s):Hu Z,Artibani M,Alsaadi A,Wietek N,Morotti M,Shi T,Zhong Z,Santana Gonzalez L,El-Sahhar S,Carrami EM,Mallett G,Feng Y,Masuda K,Zheng Y,Chong K,Damato S,Dhar S,Campo L,Garruto Campanile R,Soleymani Majd H,Rai V,Maldonado-Perez D,Jones S,Cerundolo V,Sauka-Spengler T,Yau C,Ahmed AA PubMed Article URL: http://dx.doi.org/10.1016/j.ccell.2020.01.003
1 Flow Cytometry References	
Species / Dilution	Summary
	MA5-12557 was used in Flow Cytometry to reveal that P1C1TM distinguishes between mutant and wild-type p53 expressing HLA-A24+ cells and mediates antibody dependent cellular cytotoxicity of mutant p53 expressing cells in vitro.
Human / Not Cited	Nature communications (Nov 2019; 10:) "Targeting mutant p53-expressing tumours with a T cell receptor-like antibody specific for a wild-type antigen." Author(s):Low L,Goh A,Koh J,Lim S,Wang CI PubMed Article URL: http://dx.doi.org/10.1038/s41467-019-13305-z
5 Miscellaneous PubMed References	
Species / Dilution	Summary
	MA5-12557 was used in western blot to study APC/C(Cdh1) function
Human / 1:1000	Nucleic acids research (Jun 2016; 44: 4745) "New origin firing is inhibited by APC/CCdh1 activation in S-phase after severe replication stress." Author(s):Ercilla A,Llopis A,Feu S,Aranda S,Ernfors P,Freire R,Agell N PubMed Article URL: http://dx.doi.org/10.1093/nar/gkw132
	MA5-12557 was used in immunohistochemistry (paraffin) to test if Ki-67, proliferating cell nuclear antigen, silver-staining nucleolar organizer regions and p53 could differentiate spontaneous abortions from subtypes of gestational trophoblastic diseases.
Human / 1:50	American journal of obstetrics and gynecology (Mar 2001; 184: 567) "Expressions of proliferation markers (Ki-67, proliferating cell nuclear antigen, and silver-staining nucleolar organizer regions) and of p53 tumor protein in gestational trophoblastic disease." Author(s):Kale A,Söylemez F,Ensari A PubMed Article URL: http://dx.doi.org/10.1067/mob.2001.111243
	MA5-12557 was used in immunohistochemistry to report and characterize two cases of accessory breast cancer occurring concurrently with primary invasive breast cancer.
Human / Not Cited	Cancer biology & medicine (Sep 2012; 9: 197) "Accessory breast cancer occurring concurrently with bilateral primary invasive breast carcinomas: a report of two cases and literature review." Author(s):Hao JY,Yang CC,Liu FF,Yang YL,Li S,Li WD,Li YQ,Lang RG,Fan Y,Paulos E,Zhang XM,Fu L PubMed Article URL: http://dx.doi.org/10.7497/j.issn.2095-3941.2012.03.008

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	MA5-12557 was used in immunohistochemistry - paraffin section to assess antibodies to use for the diagnosis of metastatic adenocarcinomas
Human / 1:100	<p>Diagnostic cytopathology (Dec 2011; 39: 900)</p> <p>"Cytological differential diagnosis among adenocarcinoma, epithelial mesothelioma, and reactive mesothelial cells in serous effusions by immunocytochemistry."</p> <p>Author(s):Su XY,Li GD,Liu WP,Xie B,Jiang YH</p> <p>PubMed Article URL:http://dx.doi.org/10.1002/dc.21489</p>
	MA5-12557 was used in immunohistochemistry (paraffin) to determine the roles of p53 and hMSH2 proteins in oral squamous cell carcinoma and oral dysplastic lesions.
Human / Not Cited	<p>Oral oncology (Feb 2012; 48: 120)</p> <p>"Immunoexpression of p53 and hMSH2 in oral squamous cell carcinoma and oral dysplastic lesions in Yemen: relationship to oral risk habits and prognostic factors."</p> <p>Author(s):Helal Tel A,Fadel MT,El-Thobhani AK,El-Sarhi AM</p> <p>PubMed Article URL:http://dx.doi.org/10.1016/j.oraloncology.2011.08.024</p>
5 Immunohistochemistry (Paraffin) References	
Species / Dilution	Summary
	MA5-12557 was used in immunohistochemistry - paraffin section to study the expression of Ki-67, tumor protein P53, P21, and P27 in 8 paired WHO grade II astrocytoma samples
Not Applicable / Not Cited	<p>Chinese journal of cancer (Mar 2012; 31: 159)</p> <p>"Long-term molecular changes in WHO grade II astrocytomas following radiotherapy."</p> <p>Author(s):Yue WY,Sai K,Wu QL,Xia YF,Yu SH,Chen ZP</p> <p>PubMed Article URL:http://dx.doi.org/10.5732/cjc.011.10149</p>
	MA512557 was used in immunohistochemistry - paraffin section to evaluate the diagnostic value of stathmin expression in samples of vulvar intraepithelial neoplastic lesions
Human / 1:2000	<p>Journal of clinical pathology (Dec 2016; 69: 1070)</p> <p>"Stathmin is a highly sensitive and specific biomarker for vulvar high-grade squamous intraepithelial lesions."</p> <p>Author(s):Nooij LS,Dreef EJ,Smit VT,van Poelgeest MI,Bosse T</p> <p>PubMed Article URL:http://dx.doi.org/10.1136/jclinpath-2016-203676</p>
	MA5-12557 was used in immunohistochemistry - paraffin section find prognostic clusters for breast cancer
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Not Applicable / 1:30	<p>International journal of oncology (Apr 2012; 40: 1122)</p> <p>"Identification of prognostic biomarkers for glioblastomas using protein expression profiling."</p> <p>Author(s):Jung Y,Joo KM,Seong DH,Choi YL,Kong DS,Kim Y,Kim MH,Jin J,Suh YL,Seol HJ,Shin CS,Lee JI,Kim JH,Song SY,Nam DH</p> <p>PubMed Article URL:http://dx.doi.org/10.3892/ijo.2011.1302</p>

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