

PTMScan Discovery Proteomics Services and Kits

UNPARALLELED PRODUCT QUALITY, VALIDATION, AND TECHNICAL SUPPORT



PTMScan[®] Proteomics Services

PTMScan® Proteomics Services from Cell Signaling Technology (CST) employ proprietary methodologies for antibody-based peptide enrichment combined with tandem mass spectrometry for quantitative profiling of post-translational modifications (PTMs), including phosphorylation, ubiquitination, and acetylation.



PTMScan[®] Discovery

A proteomics method for the identification and quantitation of known and novel post-translational modifications (PTMs), including Ser, Thr, and Tyr phosphorylation, as well as ubiquitination and acetylation. The method employs immunoaffinity purification using CST's proprietary motif antibodies coupled with tandem mass spectroscopy for comprehensive profiling of up to thousands of PTMs from cell line and tissue samples. This discovery-mode proteomics technology is useful for the study of PTMs throughout the proteome in a variety of biological model systems and disease states.



PTMScan[®] Direct

A multiplex proteomics assay for quantitative measurement of a defined set of known PTMs on critical signaling nodes within a group of cellular signaling pathways. The method employs an immunoaffinity purification technique using CST's proprietary modification state-specific antibodies coupled with mass spectrometry, allowing for targeted screening of hundreds of defined signaling proteins from cell line and tissue samples. This targeted-mode proteomics technology is used to investigate changes in PTMs to specific or known protein targets in response to a drug treatment or disease state.

PTMScan[®] Inquiries

Applications

Target Validation

PTMScan® Proteomics Services have been used most extensively to discover novel mechanisms of disease based on changes in PTM profiles associated with aberrantly activated kinases in disease models. PTMScan® Technology is equally effective in the identification of drug targets and potential off-target effects, mechanism-of-action studies, and the elucidation of cellular regulatory networks.

Pathway Profiling

PTMScan® Proteomics Services allow monitoring of the activation of up to thousands of cellular signaling proteins across a broad range of pathways.

Biomarker Discovery

PTMScan® Proteomics Services provide a strategy for profiling lead compounds and performing proof-of-mechanism and proof-of-concept studies. This enables the discovery and validation of biomarkers that predict compound activity, disease susceptibility, and drug response.

Patient Stratification

PTMScan® Proteomics Services enable correlation of target enzyme activation and drug sensitivity, which can provide the basis for patient stratification assay development.

Deliverables

Summary Report

The report contains qualitative and quantitative results, analysis, interpretation, and recommendations.

The following is provided in table format with detailed explanation of contents and guidelines for data review:

- Peptide Sequences
- Quantitative changes at sites of PTMs using label-free or SILAC methods
- Parent proteins and functions
- · Classification information
- DTA/mzXML ion spectra for client internal analysis

Consultation

Consultation with CST scientists for discussion and review of the results.



PTMScan[®] Discovery Proteomics Services

Advanced Proteomic Technology for Target and Biomarker Discovery

PTMScan® Proteomics Technology, developed at Cell Signaling Technology (CST), provides comprehensive analytical profiling of post-translational modifications (PTMs), including serine/threonine and tyrosine phosphorylation (PhosphoScan® Technology), ubiquitination (UbiScan® Technology), and acetylation (AcetylScan® Technology).

PTMScan Technology employs CST™ Motif Antibodies for peptide immunoaffinity purification from protease-digested cell extracts combined with LC tandem mass spectrometry (LC-MS/MS) to identify and quantify changes in modification state levels. Motif Antibodies are specific to Ser/Thr/Tyr-phosphorylated peptides, acetylated peptides, or ubiquitin-tagged peptides, enabling the analysis of phosphorylation, acetylation, or ubiquitination status of the cell.

PTMScan Services, performed by CST scientists, deliver hundreds to thousands of protein PTM identifications and quantification of PTM changes in customer samples. PTMScan Services include proprietary sample fractionation, phosphopeptide immunoaffinity purification with our optimized PTMScan Motif Antibodies, and identification using LC-MS/MS. This is followed by extensive informatic data analysis and a consultation between in-house PTMScan Technology experts and the customer.

PTMScan® Discovery Service Options

Standard PTMScan® Discovery Service: Includes a qualitative and quantitative results table, an informatics results table, a summary presentation, and a copy of the raw LC-MS data files in mzXML format.

Premium PTMScan® Discovery Service: Includes everything from the Standard PTMScan® Discovery Service option plus a summary methods report, which includes a formal, written summary of the methods and data interpretation, as well as printouts of the results table.



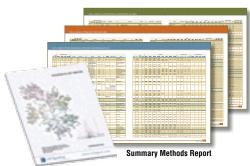
Workflow

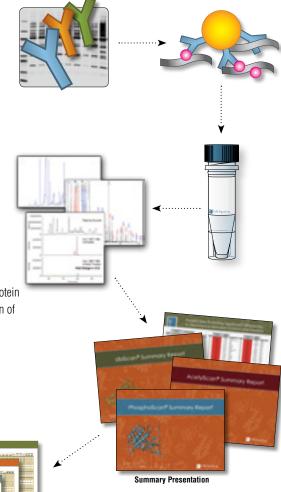
Step 1: Analysis

- Cell line and/or tissue samples ± treatment; Protease digestion.
- **::** Quality assurance of samples by western blot.
- **::** Peptide immunoaffinity purification (IAP).
- Tandem mass spectrometry (LC-MS/MS) analysis of enriched peptides for qualitative sequence and modification site identification.
- Quantitative analysis of peptide fold-change between study samples.

Step 2: Report and Consultation

- :: Report with qualitative and quantitative results.
- Report contains sequence and corresponding protein identification in table format, detailed explanation of table contents, and guidelines for data review.
- Detailed discussion and review of report with PTMScan Technology experts.
- **Typical timeline: approximately 3 to 4 weeks;** timeline will vary with project size.





PTMScan Discovery Proteomics Services

PhosphoScan° (Phosphorylation Proteomics)

Tyrosine PhosphoScan® Services: Tyrosine PhosphoScan® Services provide a comprehensive strategy for global phosphotyrosine profiling. The Cell Signaling Technology (CST) P-Tyr-100 mAb is more broadly reactive with phosphotyrosine peptides than any other antibody. Employing P-Tyr-100 for phosphopeptide enrichment in combination with LC-MS/MS analysis yields quantitative profiles of thousands of non-redundant phosphorylated sequences.

Tyrosine PhosphoScan Proteomics is used routinely and successfully for target discovery, mutant RTK screening, lead compound characterization, and biomarker discovery.

Serine/Threonine PhosphoScan® Proteomics: KinomeView® and PhosphoScan Proteomics Services together are a two-step process enabling kinome-wide quantitative analysis of cellular phosphorylation for the discovery and validation of signaling targets, kinase substrates, and response biomarkers. KinomeView Service is a western blot-based assay that provides a first step to guide the design of PhosphoScan studies by determining which branches of the kinome are responsive to the treatment condition under investigation. KinomeView can identify which Phospho-Motif Antibodies will yield the greatest insights when used in PhosphoScan mass spectrometry analysis. PhosphoScan Services provide in-depth, yet focused phosphoproteomic analysis with patented CST™ Phospho-Motif Antibodies. Each of these antibodies react broadly with diverse sets of serine-, threonine- and tyrosine-phosphorylated sequences.

AcetylScan[®] (Acetylation Proteomics)



AcetylScan® Services are a unique and unparalleled strategy for global analysis of HDAC and HAT activity on protein acetylation. CST employs antibodies optimized for high affinity to acetylated-lysine (Ac-K) peptides. Our AcetylScan Services utilize these antibodies for enrichment of acetylated peptides from protease digested cell samples, followed by LC-MS/MS analysis for quantitative profiles of thousands of non-redundant acetylated sequences.

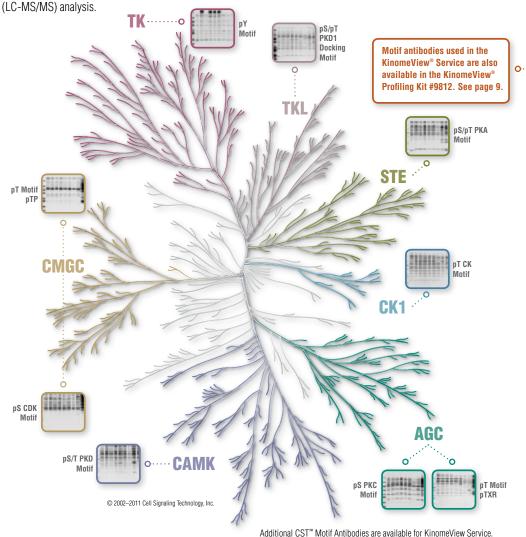
UbiScan° (Ubiquitination Proteomics)



Ubiquitination is an important mechanism for both signal transduction and proteolytic degradation. The UbiScan® Proteomics method is a unique and unparalleled strategy for global analysis of cellular ubiquitination. CST has developed a proprietary ubiquitin branch ("K-GG") XP® monoclonal antibody with specificity for a di-glycine remnant on ubiquitinated lysine residues after trypsin digestion. In UbiScan Services, the K-GG antibody is employed on trypsin digested samples to enrich ubiquitinated peptides, followed by LC-MS/MS analysis for quantitative profiles of thousands of non-redundant ubiquitinated sequences. UbiScan Proteomic Services are applied in comparative studies to identify substrates and biomarkers of ubiquitin pathway targets and compounds.

KinomeView Profiling Service

The serine/threonine phosphoproteome is extremely complex. KinomeView® Service is a western blotting service that employs CST™ Motif Antibodies with broad reactivity for serine, threonine, and tyrosine phosphorylation to provide a kinome-wide view of cellular phosphorylation. Before beginning a PhosphoScan Service, the objective of KinomeView Service is to identify which Phospho-Motif Antibodies detect phosphorylation responsive to the treatment under investigation. The Motif Antibodies that reveal treatment-mediated changes by KinomeView western blotting will be the same antibodies used in subsequent PhosphoScan Services for liquid chromatography — tandem mass spectrometry



Visit our website for a complete listing.

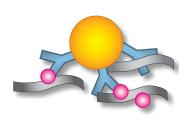
PTMScan Proteomics Product Offering

A collection of products enable you to access and use Cell Signaling Technology's patented PTMScan® Technology, as a complement to PTMScan Services.

PTMScan Discovery Motif Antibodies

Cell Signaling Technology (CST) has developed Motif Antibodies to enable focused analysis of regulatory post-translational modifications (PTMs). It is the specificity of our Motif Antibodies that makes PTMScan Technology the standard for proteomic analysis of regulatory signaling.

These Motif Antibodies are used in our PTMScan Discovery Services. We also provide these same Motif Antibodies as products with a Per-Use License to enable our customers with the capability to perform LC-MS/MS in-house as part of the PTMScan® Proteomics System and PTMScan® Motif Antibody Kits (see page 9).



KinomeView®	and	Serine	Threonine	PhosphoScan®

Target	Antibody	Bead	Kit	Consenus Site
Phospho-Akt Substrate	#9614	#1978	#5561	(RXXS*/T*)
Phospho-Akt Substrate	#10001	#1979	#5563	RXRXX(S*/T*)
Phospho-(Ser/Thr) AMPK Substrate	#5759	#5760	#5564	(L/M)XRXX(S*/T*), RXX(S*/T*)
Phospho-(Ser) ATM/ATR Substrate	#9607	#9884	*	S*Q
Phospho-(Ser/Thr) ATM/ATR Substrate	#6966	#6969	*	(S*/T*)QG, (S*/T*)Q
Phospho-(Ser) CDKs Substrate	#2324	#1981	*	(K/R)S*PX(K/R)
Phospho-(Ser/Thr) CK II Substrate	#5808	#1994	*	(S*/T*)(D/E)X(D/E)
Phospho-MAPK/CDK Substrates	#2325	#1982	#4652	PXS*P, S*PX(K/R)
Phospho-PKA Substrate	#9624	#1984	#5565	(K/R)(K/R)X(S*/T*)
Phospho-(Ser) PKC Substrate	#6967	#6970	*	(K/R)XS*X(K/R)
Phospho-(Ser/Thr) PDK1 Docking Motif	#9634	#1992	*	(F/Y)(S*/T*)(F/Y)
Phospho-(Thr) PLK Binding Motif	#5243	#5756	#5566	ST*P
Phospho-Thr-Pro Motif	#3003	#1996	#5567	T*P, T*PP
Phospho-Thr-Pro-Glu Motif	#3004	#1987	*	T*PE, T*P
Phospho-Thr-X-Arg Motif	#2351	#1988	*	T*XR, T*PR

Tyrosine PhosphoScan®

Target	Antibody	Bead*	Kit	Consenus Site
Phosphotyrosine (P-Tyr-100)	#9411	#1991	#5636	Υ*

UbiScan°

Target	Antibody	Bead	Kit	Consenus Site
Ubiquitin Branch	#3925	#1990	#5562	K-ε-GG

AcetvIScan®

Mootyroodii				
Target	Antibody	Bead	Kit	Consenus Site
Acetylated-Lysine	#9814	1989	*	Ac-K

^{*}Not for individual sale — available only as components of the PTMScan® Proteomics System.

^{*}Visit our website for updates on new products.

PTMScan Motif Antibody Kits

PTMScan® Motif Antibody Kits are a product line that enables the use of PTMScan Technology, a proprietary technology developed at CST. The kits provide a selection of our proprietary Motif Antibodies ready for use in studies that employ PTMScan Technology. These antibodies are identical to those utilized by CST scientists in PTMScan Discovery Services. For our customers with in-house expertise in mass spectrometry proteomics, these kits provide a cost-effective means of accessing a selection of CST™ Motif IAP Beads and a Limited Use License to perform PTMScan Technology.

#5636 PTMScan® Phospho-Tyrosine Mouse mAb (P-Tyr-100) Kit

#5567 PTMScan® Phospho-T*PP Motif (T*PP) XP® Kit

#5564 PTMScan® Phospho-AMPK Substrate Motif (LXRXXS*/T*) Kit

#5561 PTMScan® Phospho-Akt Substrate Motif mAb 1 (RXXS*/T*) Kit

#5563 PTMScan® Phospho-Akt Substrate Motif mAb 2 (RXRXXS*/T*) Kit

#4652 PTMScan® Phospho-MAPK/CDK Substrate Motif
(PXS*P and S*PXK/R) Kit

#5565 PTMScan® Phospho-PKA Substrate Motif (RRXS*/T*) Kit

#5566 PTMScan® Phospho-ST*P Motif (ST*P) XP® Kit

#5562 PTMScan® Ubiquitin Remnant Motif (K-ε-GG) Kit

Reagents for 5 assays as well as detailed protocols are provided in each kit.



PTMScan® Inquiries: Pharma Services Department: ptmscan@cellsignal.com

KinomeView® Profiling Kit #9812

Provides a set of Phospho-Motif Antibodies that covers a large portion of the kinome and reacts broadly with serine, threonine, and tyrosine phosphorylation mediated by diverse kinase families. This kit provides researchers with a western blotting strategy to dissect the complexity of the phosphoproteome and determine the kinase families involved in the regulation of diverse physiological processes.

Products Included	Motif	Isotype
#9614 Phospho-Akt Substrate (110B7E) Rabbit mAb	(RXXS*/T*)	Rabbit IgG
#10001 Phospho-Akt Substrate (23C8D2) Rabbit mAb*	RXRXX(S*/T*)	Rabbit IgG
#5759 Phospho-(Ser/Thr) AMPK Substrate Rabbit mAb*	(L/M)XRXX(S*/T*), RXX(S*/T*)	Rabbit IgG
#9607 Phospho-(Ser) ATM/ATR Substrate Rabbit mAb*	S*Q	Rabbit IgG
#6966 Phospho-(Ser/Thr) ATM/ATR Substrate Rabbit mAb*	(S*/T*)QG, (S*/T*)Q	Rabbit IgG
#2324 Phospho-(Ser) CDK Substrate Antibody	(K/R)S*PX(K/R)	Rabbit IgG
#5808 Phospho-(Ser/Thr) CKII Substrate Antibody*	(S*/T*)(D/E)X(D/E)	Rabbit IgG
#2325 Phospho-MAPK/CDK Substrates (34B2) Rabbit mAb	PXS*P, S*PX(K/R)	Rabbit IgG
#9634 Phospho-(Ser/Thr) PDK1 Docking Motif (18A2) Mouse mAb	(F/Y)(S*/T*)(F/Y)	Mouse IgG2
#9624 Phospho-PKA Substrate (100G7E) Rabbit mAb	$(K/R)(K/R)X(S^*/T^*)$	Rabbit IgG
#6967 Phospho-(Ser) PKC Substrate Rabbit mAb*	(K/R)XS*X(K/R)	Rabbit IgG
#5243 Phospho-(Thr) PLK Binding Motif (D73F6H9) Rabbit mAb*	ST*P	Rabbit IgG
#3003 Phospho-Thr-Pro Motif Antibody*	T*P, T*PP	Rabbit IgG
#3004 Phospho-Thr-Pro-Glu Motif (C32G12) Rabbit mAb*	T*PE, T*P	Rabbit IgG
#2351 Phospho-Thr-X-Arg Motif Antibody	T*XR, T*PR	Rabbit IgG
#9411 Phospho-Tyrosine Mouse mAb (P-Tyr-100)	Y*	Mouse IgG1
#7074 Anti-rabbit IgG, HRP-linked Antibody	_	Goat
#7076 Anti-mouse IgG, HRP-linked Antibody	_	Horse

^{*} Select antibodies contained in the KinomeView Profiling Kit are not available for individual sale

PTMScan[®] Proteomics System

Cell Signaling Technology (CST) offers a Per-Use License with the PTMScan® Proteomics System to provide our customers with in-house expertise in mass spectrometry proteomics with a cost-effective means of accessing CST™ Motif IAP Beads and proprietary methodology. This system provides any of our Motif IAP Beads validated and ready for use in PhosphoScan®, UbiScan®, or AcetylScan® experiments.

Flexibility

System is built-to-order; the Motif Antibodies included and the amount of each can be varied according to your specification and the number of analyses you intend to perform.

Consistent Performance

Each bead-coupled Motif Antibody in the system is custom prepared and validated prior to shipment to ensure consistent and high quality performance.

Support

Scientists with extensive PTMScan Technology experience are available to provide technical support. This system also includes a detailed protocol describing cell sample preparation, IAP enrichment, and peptide sample preparation for LC-MS/MS analysis.

System Components

- Motif IAP Beads One or more available
- IAP Buffer
- PTMScan® Protocol
- PTMScan® Per-Use License



NOTE: The PTMScan® Proteomics System provides a limited Per-Use License that is restricted to allow performance of CST® PTMScan methods with only Motif IAP Beads purchased as part of the CST PTMScan Proteomics System.

Products for Follow-up Validation Studies

Several reagents from Cell Signaling Technology's product offering are readily available for follow-up target validation studies after completion of the PTMScan® Service Project. A selection of these products is high-lighted and hyperlinked from the PTMScan Service Results Summary Reports for our clients' convenience.

Visit www.cellsignal.com for a complete product listing.

AQUA™ Peptide Synthesis

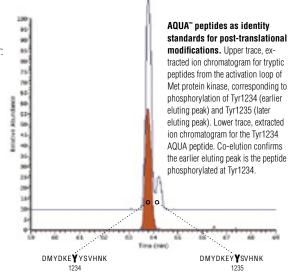
To validate and quantify novel findings through PTMScan Technology, CST offers custom-synthesized, stable isotopelabeled AQUA™ peptides. AQUA peptides enable isotope dilution strategies, which are the gold standard of quantitative mass spectrometry. We specialize in high-quality custom AQUA peptide synthesis, including challenging sequences and unusual post-translational modifications. We have hundreds of synthetic AQUA peptides in stock, and turn-around on new AQUA peptide synthesis and quality control is approximately 4–6 weeks.

This method was developed by scientists at Harvard Medical School. Limited use of this method is permitted under a licensing arrangement with Harvard College.

- Peptides are typically ~20 residues or less in length but longer lengths are available
- Modification options include: phosphorylation; acetylation; ubiquitination; cysteine alkylation; methyl-arginine, methyl-histidine, and methyl-lysine
- ## Heavy residues: leucine, valine, proline, phenylalanine, alanine, and glycine can be incorporated
- ... No minimum order

AQUA[™] Peptide Inquiries:

Sales Department at Cell Signaling Technology: 866-310-9776 • Sales@cellsignal.com



- ** Modification state-specific and total protein antibodies, including our exclusive line of XP® monoclonal antibodies, are produced and validated in-house in a number of applications, including western blotting, immunohistochemistry, and immunofluorescence.
- ** SignalSilence* siRNA duplexes that allow knockdown of specific human or mouse proteins are available for many targets.
- **PathScan® Sandwich ELISA Kits and Antibody Pairs** are available to measure a large selection of intracellular signaling molecules. These products are developed, produced, and validated in-house, ensuring robust, sensitive, and specific assays.

Publications by Cell Signaling Technology Scientists

2011

Gu, T.L. et al. (2011) Survey of tyrosine kinase signaling reveals ROS kinase fusions in human cholangiocarcinoma. *PLoS ONE* 6, e15640.

Gu, T.L. et al. (2011) Survey of activated FLT3 signaling in leukemia. *PLoS ONE* 6, e19169

Kettenbach, A.N. et al. (2011) Absolute quantification of protein and post-translational modification abundance with stable isotope-labeled synthetic peptides. *Nat. Protoc.* 6, 175–86.

Dammer, E.B. et al. (2011) Polyubiquitin linkage profiles in three models of proteolytic stress suggest the etiology of Alzheimer disease. *J. Biol. Chem.* 286, 10457–10465.

Pighi, C. et al. (2011) Phospho-proteomic analysis of mantle cell lymphoma cells suggests a pro-survival role of B-cell receptor signaling. *Cell Oncol. (Dordr)* 34, 141–153.

Brave, S.R. et al. (2011) Assessing the Activity of Cediranib, a VEGFR-2/3 Tyrosine Kinase Inhibitor, against VEGFR-1 and Members of the Structurally Related PDGFR Family. *Mol. Cancer Ther.* 10, 861–873.

2010

Moritz, A. et al. (2010) Akt-RSK-S6 kinase signaling networks activated by oncogenic receptor tyrosine kinases. *Sci. Signal* 3, ra64.

Kang, S. (2010) p90 ribosomal S6 kinase 2 promotes invasion and metastasis of human head and neck squamous cell carcinoma cells. *J. Clin. Invest.*120, 1165–1177.

Bonnette, P.C. et al. (2010) Phosphoproteomic characterization of PYK2 signaling pathways involved in osteogenesis. *J. Proteomics* 73, 1306–1320.

Aguiar, M. et al. (2010) Gas-Phase Rearrangements Do Not Affect Site Localization Reliability in Phosphoproteomics Data Sets. *J. Proteome Res.* 9, 3103–3107.

Carretero, J. et al. (2010) Integrative genomic and proteomic analyses identify targets for Lkb1-deficient metastatic lung tumors. *Cancer Cell* 17, 547–559.

Andersen, J.N. et al. (2010) Pathway-based identification of biomarkers for targeted therapeutics: personalized oncology with PI3K pathway inhibitors. *Sci. Transl. Med.* 2, 43ra55.

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2009

Schwer, B. et al. (2009) Calorie restriction alters mitochondrial protein acetylation. *Aging Cell* 8, 604–606.

Xu, P. et al. (2009) Quantitative proteomics reveals the function of unconventional ubiquitin chains in proteasomal degradation. *Cell* 137, 133–145.

2008

Pan, S. et al. (2008) Application of Targeted Quantitative Proteomics Analysis in Human Cerebrospinal Fluid Using a Liquid Chromatography Matrix-Assisted Laser Desorption/Ionization Time-of-Flight Tandem Mass Spectrometer (LC MALDI TOF/TOF) Platform. *J. Proteome Res.* 7, 720–730.

Guo, A. et al. (2008) Signaling networks assembled by oncogenic EGFR and c-Met. *Proc. Natl. Acad. Sci. USA* 105, 692–697.

2007

Rikova, K. et al. (2007) Global survey of phosphotyrosine signaling identifies oncogenic kinases in lung cancer. *Cell* 131, 1190–1203.

Stokes, MP. et al. (2007) Profiling of UV-induced ATM/ ATR signaling pathways. *Proc. Natl. Acad. Sci. USA* 104, 19855–19860. Gerber, S.A. et al. (2007) The absolute quantification strategy: application to phosphorylation profiling of human separase serine 1126. *Methods Mol. Biol.* 359, 71–86.

Gu, T.L. et al. (2007) Phosphoproteomic analysis identifies the M0-91 cell line as a cellular model for the study of TEL-TRKC fusion-associated leukemia. *Leukemia* 21, 563–566.

Gu, T.L. et al. (2007) A novel fusion of RBM6 to CSF1R in acute megakaryoblastic leukemia. *Blood* 110, 323–333.

2006

Mercher, T. et al. (2006) JAK2T875N is a novel activating mutation that results in myeloproliferative disease with features of megakaryoblastic leukemia in a murine bone marrow transplantation model. *Blood* 108, 2770–2779.

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Griswold, I.J. et al. (2006) Kinase domain mutants of Bcr-Abl exhibit altered transformation potency, kinase activity, and substrate utilization, irrespective of sensitivity to imatinib. *Mol. Cell. Biol.* 26, 6082–6093.

Gu, T.L. et al. (2006) Phosphotyrosine profiling identifies the KG-1 cell line as a model for the study of FGFR1 fusions in acute myeloid leukemia. *Blood* 108, 4202–4204.

Goss, V.L. et al. (2006) A common phosphotyrosine signature for the Bcr-Abl kinase. *Blood* 107, 4888–4897.

2005

Zhang, Y. et al. (2005) Time-resolved mass spectrometry of tyrosine phosphorylation sites in the epidermal growth factor receptor signaling network reveals dynamic modules. *Mol. Cell Proteomics* 4, 1240–1250.

Rush, J. et al. (2005) Immunoaffinity profiling of tyrosine phosphorylation in cancer cells. *Nat. Biotechnol.* 23, 94–101.

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