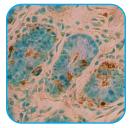
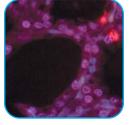
# MILLIPORE

# Apoptosis Antibodies, Reagents and Kits









now part of Millipore

## Introduction

Apoptosis is a critical process for homeostasis ... disruption of an appropriate apoptotic response is implicated in the development of many disease states.

# Upstate<sup>®</sup>, Chemicon<sup>®</sup> and Linco<sup>®</sup> are now part of Millipore

The goal of this combined company is to provide more innovative tools, services and application expertise that will improve your productivity. Our first priority is to support your work as a valued partner. Together we now offer:

- A broader range of products and services for markets including drug discovery, protein identification, antibody purification, molecular biology, stem cell research, water purification and general filtration.
- An increased number of applications and protocols that facilitate your development process.
- Extended global expertise, leadership and presence to serve you more effectively.
- Larger support and research development teams to deliver innovative products that cost-effectively impact your business.

# Apoptosis Antibodies, Reagents and Kits

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# Apoptosis Programmed Cell Death

## **A Brief Review**

Apoptosis, or programmed cell death, is a process for the orderly disposal of unwanted cells without the development of an inflammatory response, which is often associated with necrotic cell death. Table 1 highlights the differences between apoptosis and necrosis.

Apoptosis is required for many normal physiological functions. For example, interdigital apoptosis is a critical developmental process that contributes to the formation of fingers and toes; disruptions in this process lead to severely debilitating congenital birth defects. In addition, normal resolution of inflammation requires that inflammatory cells undergo apoptosis in order to prevent chronic inflammation.

Apoptosis is also linked to cancer. Normal cells exhibit adhesion-dependent growth; they must be attached (to other cells or an extracellular matrix) for survival. Cell adhesion transmits survival signals, and loss of these signals triggers apoptosis. Cells must not survive should they migrate to a new tissue, where they may exhibit unregulated growth. In fact, most cancer cells have lost adhesion-dependence. Growth in soft agar and "piling", characteristics of cells which have lost adhesion-dependence, are also behaviors observed in cells isolated from aggressive and metastatic cancer. Vast effort has been expended to identify steps in the apoptotic cascade that may serve as targets for cancer therapy.

"Extrinsic" versus "Intrinsic" Apoptosis Induction

At least two pathways lead to apoptosis, the "extrinsic" and "intrinsic" pathways. The extrinsic pathway is initiated by extracellular determinants through "death receptors". Binding of Fas ligand to its receptor results in receptor trimerization and the formation of an intracellular complex composed of Fas "death domains", the protein FADD and pro-Caspase 8 (an initiator Caspase; others include Caspases 9, 10 and 12). Pro-Caspase 8 undergoes proteolytic processing to form active Caspase 8. Activated Caspase 8 activates pro-Caspase 3; active Caspase 3 is a principle effector caspase of apoptosis. Caspase 8 also activates the protein Bid, which activates Bak and Bax. Similar processes are initiated from the TNF receptor, another death receptor family member, which recruits TRADD, and pro-Caspase 8 and so on.

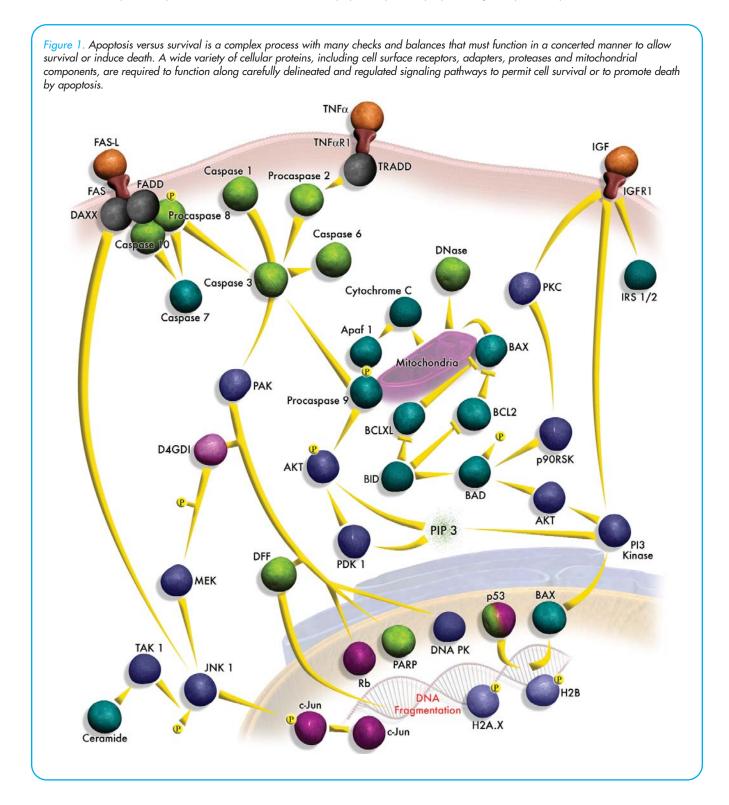
The intrinsic pathway begins when an injury occurs within the cell as a result of drug treatment,

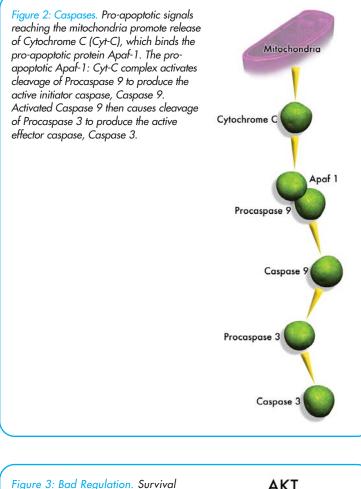
	Apoptosis	Necrosis	
Incidence	Scattered	Localized	
Cell Volume	Shrunken	Swollen	gical es
Cell Surface	Sealed	Leaky	holoç erenc
Chromatin	Condensed	Clumped	Morphological Differences
Response	Phagocytosis	Inflammation	
Onset	Regulated	Accidental	
Enzyme Cascade	Complete	Truncated	<b>Biochemical</b> Differences
Biosynthesis	Viable	Non-viable	ochei iffere
DNA Fragmentation	Non-random (Laddered)	Random (Smeared)	iii iii

#### Table 1: Apoptosis vs. Necrosis

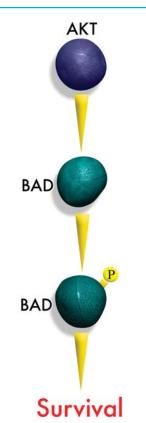
radiation (i.e., radiation-induced DNA damage) or detoxification-induced redox damage. Mitochondria are pivotal in orchestrating events that keep a cell alive or commit it to death. The key event seems to be mitochondrial membrane permeabilization and release of Cytochrome C (Cyt-C). Cytoplasmic Cyt-C binds to and activates the protein Apaf-1, which activates pro-Caspase 9. Active Caspase 9 then activates Caspase 3 (see Figure 2: Caspases on page 6).

This process is complicated as mitochondrial membrane permeability (and Cyt-c release) is induced by the proteins Bak and Bax, which are activated by BID (Figure 1). The pro-survival protein Bcl-2 prevents apoptosis, probably by binding to Cyt-C or Apaf-1,





rigure 3: bad Regulation. Survival signals result in activation of Akt, which phosphorylates the Bcl-family protein Bad, and promotes survival. Unphosphorylated Bad dimerizes with Bcl-2 and Bcl-XL, preventing them from performing their pro-survival function, and thus promoting cell death. Phosphorylated Bad is unable to associate with Bcl-2 and Bcl-XL, allowing them to perform their pro-survival function.



or through binding to Bak and Bax, preventing release of Cyt-C. The pro-apoptotic protein Bad binds to Bcl-2 and Bcl-XL, preventing their pro-survival function. Survival signals, however, activate the kinase Akt, which phosphorylates Bad and prevents binding to Bcl-2, promoting survival (see Figure 3: Bad Regulation). Inhibitor of Apoptosis Proteins (IAPs) normally bind and prevent activation of Caspase 3. Mitochondrial proteins released by Bak and Bax, such as Smac/DIABLO and Omi/HtrA2, counteract the effect of IAPs and promote death. The abundance, location and phosphorylation state of all of these proteins are essential in deciding between life and death; disruptions in any of them produce aberrant apoptosis. These checks and balances provide assurance that apoptosis occurs only when necessary.

Both pathways result in activation of effector Caspases, such as Caspase 3. Effector Caspases (Caspases 3, 6 and 7) are the executioners of apoptosis, as they are responsible for the degradation of other cellular proteins (i.e., cytoskeletal proteins) prior to packaging and disposal. Subsequent processes result in many events, including phosphorylation of histone H2A.X (Ser149) and H2B (Ser14), cleavage of PARP, chromatin condensation, inter-nucleosomal degradation of DNA and packaging of the cell into "apoptotic bodies" that are taken up by phagocytes.

## Loss of Membrane Asymmetry

A very early event in apoptosis is loss of membrane asymmetry (see Figure 5 on page 10). This occurs as the membrane phospholipid phosphatidylserine (PS), normally present on the inner leaflet of cell membranes, redistributes to both the inner and outer leaflets. PS translocation is a key event, as cells lacking PS are not phagocytized. It seems that PS on the outside of a cell is the trigger for recognition by a PS receptor on phagocytic cells, which recognize and remove dying cells. PS receptor deficient mice accumulate dead cells, resulting in embryonic lethality.

#### Ubiquitin-Mediated Regulation of Apoptosis

Previous work has shown the extensive role of phosphorylation in regulating Apoptosis. What is becoming more evident is apoptosis regulation mediated by ubiquitin and ubiquitin-like proteins. Like cell division and DNA Damage repair, apoptosis requires extremely tight regulation to be initiated in the proper context. Mostly, addition of ubiquitin to target proteins results in rapid and irreversible protein degradation via the proteosome. Protein degradation is an efficient and quick mechanism to eliminate proteins that have crucial activities. For example, pro-survival Bcl-2 family members are specifically degraded in response to TNF- $\alpha$  mediated apoptosis, thereby eliminating its "survival" ability. This is a key event in TNF $\alpha$  mediated apoptosis. There are many more examples of ubiquitin mediated degradation of key apoptotic proteins, this is just one. However, degradation is not the only outcome. Ubiqutin and other ubiquitin-like proteins can regulate other cellular events like trafficking, proteinprotein, interactions and activation/deactivation. Active research in these areas will soon provide details on the regulatory ability of ubiquitin on the apoptosis pathway.

There are many more proteins and pathways involved in apoptosis than could be mentioned here. Indeed, there have been nearly 80,000 publications on the subject, including over 13,000 reviews. Apoptosis is a critical process during development and is fundamental to homeostasis in nearly all multicellular organisms. Disruption of an appropriate apoptotic response is implicated in the development of many disease states, including cancer (lack of apoptosis) or one of several degenerative diseases (enhanced apoptosis).

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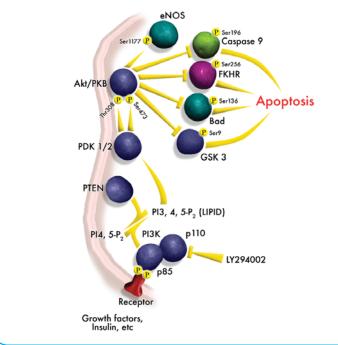
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# Akt/PKB A Key Molecule in Apoptosis Regulation

# Background

The Ser/Thr kinase Akt is a major known effecter of the PI 3-Kinase pathway and is involved in multiple signaling pathways that relate to many biological processes including cell survival, cell cycle control, cell growth and cell metabolism. The activity of Akt is regulated by its phosphorylation status, primarily on two residues, Thr308 and Ser473. A better understanding of its activation/deactivation was only recently further elucidated by the discovery of the TORC2 kinase complex and PHLPP1 & 2 phosphatases. The regulation of Akt activity and downstream targets greatly affect the cell's ability to enter into apoptosis. With Akt having so many signaling partners and its involvement in multiple signaling pathways and cellular mechanisms, it is

Figure 4. The direct interaction of activated PDK1 with Akt results in the recruitment of Akt to the plasma membrane and the subsequent phosphorylation and activation of Akt. Activated AKT is then released from the plasma membrane into the cytosol and nucleus where it interacts with and phosphorylates multiple binding partners including Caspase 9, Bad, and GSK3 resulting in the inhibition of apoptosis and the promotion of cell survival.



no wonder why Akt is so well studied and a highly sought after drug target.

The activation and regulation of Akt is dependent on a dual regulatory mechanism that requires both its translocation to the plasma membrane and dual phosphorylation on Thr308 and Ser473 by PDK1 and the TORC2 complex, respectively (Figure 5). This is accomplished by the generation and build-up of PIP3 by PI3K in conjunction with reduced PTEN function, which results in the activation of PDK1 (3-phosphoinositide-dependent protein kinase-1) and the recruitment of Akt to the plasma membrane by direct interaction with its PH domain. PDK1 then in turn phosphorylates Akt on Thr 308 in its activation loop. This phosphorylation is necessary and sufficient for Akt activation; however maximal activation requires the additional phosphorylation at Ser473. Another kinase complex, recently determined as TORC2 that is composed of the mTOR/Rictor heteroduplex (previously referred to as the unidentified kinase PDK2) phosphorylates Akt on Ser473 in the hydrophobic motif. This latter phosphorylation/activation is not fully understood. Following its activation, Akt is released from the plasma membrane into the cytosol and nucleus where it interacts and phosphorylates multiple binding partners including FKHR, BAD and GSK. These two phosphorylation sites are regulated by two different phosphatases. The dephosphorylation event results in the termination of Akt activity which is much less understood than its activation. This process was less understood until the recent discovery of two new phosphatases, PHLPP1 and PHLPP2 (PH domain leucine-rich repeat protein phosphatase), which dephosphorylates Akt in the cytoplasm and nucleus, respectively (specifically on the hydrophobic phosphorylation site of Ser473, but not the Thr308 site). Interestingly, dephosphorylation on this site has been shown to trigger apoptosis and the suppression of tumor growth. Another phosphatase, PP2A, that is much precarious, is now believed to dephosphorylate Akt on the PDK1 phosphorylation site of Thr308. Together, they help regulate the activity of Akt.

Akt functions in many anti-apoptotic pathways through its binding of many apoptotic mediators that become either activated or de-activated through their Akt phosphorylation. The Akt cell survival mechanism via cell cycle, growth translation and apoptosis is likely to involve many targets that it phosphorylates. These downstream mediators of apoptosis can be divided into two separate groups: ones that Akt activates via phosphorylation, such as Caspase 9, IKK, MDM2,  $\mathsf{NF}\kappa\mathsf{B}$  and mTOR and ones that it inhibits via phosphorylation, such as GSK3 $\beta$ , FKHR and BAD. Akt activates many targets that inhibit various apoptosis pathways. The phosphorylation of Caspase-9 on Ser196 by Akt results in the inhibition of its catalytic activity. In the NF $\kappa$ B pathway, which promotes survival in response to pro-apoptotic stimuli, Akt phosphorylates IKK ( $I\kappa B$  Kinase) that results in the phosphorylation and subsequent degradation of the NF $\kappa$ B inhibitor I $\kappa$ B. This frees the NF $\kappa$ B p50/p65 complex to translocate from the cytoplasm to the nucleus where it activates its target genes. Akt also helps mediate the proapoptotic tumor suppressor/transcription factor p53 by phosphorylating its negative regulating binding protein MDM2. Akt phosphorylation of MDM2 allows it to more efficiently translocate to the nucleus where it binds to p53 and target it for degradation by the proteosome through E3 ubiquitin ligase activity. For others, Akt phosphorylation results in its inactivation. For the pro-apoptotic member of the BCL2 family, BAD, phosphorylation on Ser136 prevents BAD from interacting with BCL-XL, thus enabling the anti-apoptotic function of BCL-XL. Akt also phosphorylates the FOXO family of transcription factors. For FKHR, it is on sites Thr24/ Ser256/Ser319. This phosphorylation results in it bidding to 14-3-3 and ultimately its inactivation of cytoplasm/nuclear shuttling and prevents the activation of the FKHR target genes that include many pro-apoptotic proteins, such as BIM and FAS ligand. Finally, as cell cycle control is very important is growth and apoptosis, the levels of Cyclin D1 are very important as they are important in the G1/S phase transition. Akt aids in the role of preventing the degradation of CyclinD1 by regulating the activity of GSK3 $\beta$  (Glycogen Synthase Kinase- $3\beta$ ) by its phosphorylation, thus de-activating it. GSK3<sup>β</sup> phosphorylates CyclinD1 and marks it for degradation. If its activity is blocked, then CyclinD1 will not be marked for degradation and will accumulate in the cell. In addition to CyclinD1, GSK also phosphorylates many other downstream targets that

affect apoptosis and cell cycle regulation, including c-Myc and Cyclin E and the transcription factors c-Jun, β-catenin, GLI, Notch, Snail and SREBP1. Millipore offers a complete Akt/PKB solution with Upstate antibodies, kinases, phosphatases, substrates and siRNA for targets upstream and downstream of Akt. This includes mouse and rabbit monoclonal and polyclonal antibodies directed against total and phospho-specific variants Akt, as well as Alexa Fluor® conjugates. High quality active and inactive Upstate kinases of both upstream activators and downstream substrates are available. Activation of Akt can be measured directly by immunoprecipitation followed by phosphorylation of a known substrate with radio labeled ATP or with many of our phospho-specific antibodies such as AS160, PRAS40, cardiac PFK2, BAD, FKHR and GSK3. Millipore has what you need for your Akt research.

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# Anti-Phosphatidlyserine Clone 1H6, Alexa Fluor 488 conjugate

Anti-Phosphatidylserine, now conjugated with Alexa Fluor 488, is perfect for your flow cytometry and immunofluorescence applications.

Antibody-based detection of early apoptotic events is now possible for less cost than Annexin V assays. Upstate's Anti-Phosphatidylserine, clone 1H6, Alexa Fluor 488 from Millipore, a monoclonal antibody that detects the initial cellular changes of apoptosis, is sensitive, specific and useful for flow cytometry studies for only \$0.68 per assay. It can also be used for immunocytochemistry.

Apoptosis, or programmed cell death, is a physiological mechanism for the neat and tidy elimination of unnecessary or unwanted cells. One well-documented event that occurs very early in apoptosis is the loss of membrane phospholipid – asymmetry [the "phosphatidylserine (PS) flip" – see Figure 5]. Following the initiation of apoptosis, PS flips from the cytoplasmic to the extracellular side of the still-intact cell membrane. The PS flip occurs well before any other apoptosis events, such as caspase activation, PARP cleavage or DNA fragmentation.

Cell-surface PS is thought by some to be a marker for recognition and phagocytosis of apoptotic cells.

PS is recognized by a PS Receptor (PSR) on phagocytes and PSR gene knockout has been shown to be an embryonic lethal mutation. This seems to be critical in prevention of an inflammatory response to cell death. Other reports have suggested, on the other hand, that the PSR is not as critical in clearance of apoptotic cells.

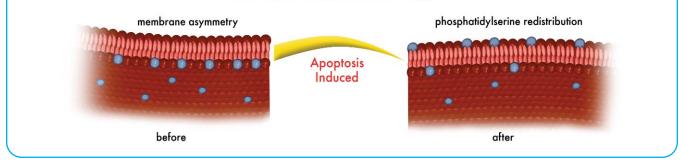
Anti-Phosphatidylserine, clone 1H6, Alexa Fluor 488 delivers long-term product consistency, high specificity and is more affordably priced than Annexin V recombinant protein-based assays. For more information on Anti-Phosphatidylserine, clone 1H6, Alexa Fluor 488 and our other products for the study of apoptosis, visit our website at www.millipore.com/apoptosis.

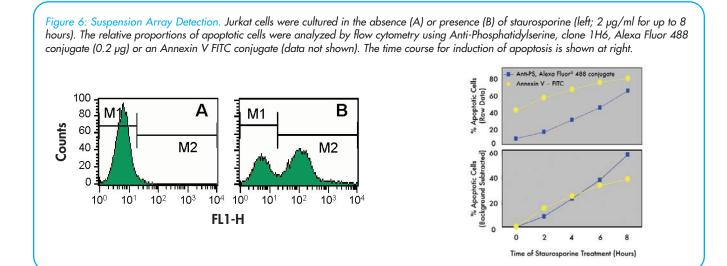
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Figure 5: Phosphatidylserine "Flip". PS is a membrane phospholipid that is normally localized to the inner surface of cell membranes. During apoptosis, PS redistributes from the inner membrane leaflet to both the inner and outer membrane leaflets. "Loss of membrane asymmetry" is a key marker for apoptosis.

# Phosphatidylserine "Flip"





## Products for Phosphatidylserine Detection

Description	Quantity	Cat. No.	Price
Anti-Phosphatidylserine, clone 1H6, Alexa Fluor 488 conjugate	100 ha	16-256	\$339
Unconjugated Anti-Phosphatidylserine Antibody	200 µg	05-719	\$339
Related Products			
Propidium Iodide Solution	1 mL	S7109	\$60
Propidium Iodide/Antifade Solution	1 mL	S7112	\$60

# **Other Apoptosis Detection Products**

## **TUNEL Based Apoptosis Detection Kits**

1 1			
ApopTag Fluorescein In Situ Apoptosis Detection Kit	40 assays	S7110	\$405
ApopTag Plus Fluorescein In Situ Apoptosis Detection Kit	40 assays	S7111	\$415
ApopTag Red In Situ Apoptosis Detection Kit	40 assays	S7165	\$390

#### **Active Caspase Detection Kits**

CaspScreen Flow Cytometric Apoptosis Detection Kit	25 assays	APT105	\$395
CaspaTag Pan-Caspase In Situ Assay Kit	100 assays	APT400	\$435
CaspaTag Caspase-3,7 In Situ Assay Kit	100 assays	APT403	\$435

## Antibodies to Apoptosis Specific Cleavage Products

Anti-Fractin (32 kDa Fragment of $\beta$ -Actin)	100 µL	AB3150	\$265
Anti-PARP (Inactive and Cleaved)	100 µL	AB16661	\$330

# Apoptosis Detection Kits

# Millipore's Offering

Millipore offers Apoptosis Detection Kits for researchers to determine if apoptosis is occurring in cells or tissue of interest. The broad variety of choices allows the researcher to choose the kit that best fits their needs. The kits are shown below based on their detection options.

### Table 2:

#### Recommended Sample Types for Kits

Sample Type	ApopTag	CaspaTag	MitoLight	Glutathione	DNA Ladder	CaspSCREEN	Anti-PS*
Adherent Cells	•	•	•	•	•		•
Suspended Cells	•	•	•	•	•	•	•
Tissues	•				•		

\*Anti-Phosphatidlyserine (Anti-PS)

#### Apoptosis Detection Kits by Hallmark Apoptotic Event

Detection Method	Extracellular Phosphatidylserine Exposure	Caspase Activity	GSH Level Changes	Mitochondrial Permeabilization	DNA Fragmentation
Flow Cytometry	Anti-PS Page 11 Cat. Nos. 16-256, 05-719	CaspaTag Page 13 Cat. Nos. APT400, APT403, APT420, APT423, APT408, APT428, APT409, APT429		MitoLight Page 15 Cat. Nos. APT142, APT242	<b>ApopTag</b> Page 17 <b>Cat. Nos.</b> S7110, S7111, S7165, S7160
		CaspSCREEN Page 17 Cat. No. APT105			Apo-Direct Page 16 Cat. No. APT110
Fluorescence Microscopy	Anti-PS Page 11 Cat. Nos. 16-256, 05-719	CaspaTag Page 13 Cat. Nos. APT400, APT403, APT500, APT503, APT420, APT423, APT408, APT428, APT409, APT429, APT520, APT523	Glutathione Detection Kit Page 14 Cat. No. APT250	MitoLight Page 15 Cat. Nos. APT142, APT242	<b>ApopTag</b> Page 17 <b>Cat. Nos.</b> S7110, S7111, S7165
Fluorescence Reader		CaspaTag Page 13 Cat. Nos. APT400, APT403, APT500, APT503, APT420, APT423, APT408, APT428, APT409, APT429, APT520, APT523		Mitolight Page 15 Cat. Nos. APT142, APT242	ApopTag Pages 16 & 17 Cat. Nos. S7100, S7101, S7200
Light Microscopy	Anti-PS Page 11 Cat. Nos. 16-256, 05-719				
Agarose Gel					DNA Ladder Page 15 Cat. No. APT151

# CaspaTag In Situ Apoptosis Detection Kits

Chemicon CaspaTag *In Situ* Apoptosis Detection Kits from Millipore detect caspases in living cells undergoing apoptosis. Fluorescently labeled inhibitors of caspase activity irreversibly bind to active caspases, allowing apoptotic cells to be identified by their fluorescence using flow cytometry, fluorescence microscopy or fluorescence reader.

CaspaTag Kits utilize carboxyfluorescein (FAM) or sulforhodamine (SR) labeled fluoromethyl ketone (FMK)

inhibitor probes. These probes are derivatives of benzyloxycarbonyl-peptide (caspase recognition sequence)-FMK caspase inhibitors. They enter the cell and irreversibly bind to caspase 3, 7 (DEVD), caspase 8 (LETD), caspase 9 (LEHD) or all caspases (VAD).

#### CaspaTag Advantages

Fast: Assay completed in less than 90 minutes Simple: One-step assay in live cells Effective: Irreversibly binds to active caspases with high affinity

Figure 7: Suspension Array Detection. Jurkat cells were treated with camptothecin to induce apoptosis. Cells were then labeled with SR-VAD-FMK and Hoescht stain and visualized by fluorescence microscopy. Cells with caspase activity are shown in red (right) while nuclear staining of all cells is shown in blue (left).



Description	Quantity	Cat. No.	Price
CaspaTag Pan-Caspase In Situ Assay Kit, Fluorescein	25 assays	APT420	\$155
CaspaTag Pan-Caspase In Situ Assay Kit, Fluorescein	100 assays	APT400	\$435
CaspaTag Pan-Caspase In Situ Assay Kit , Sulforhodamine	25 assays	APT520	\$185
CaspaTag Pan-Caspase In Situ Assay Kit, Sulforhodamine	100 assays	APT500	\$470
CaspaTag Caspase-3,7 In Situ Assay Kit, Fluorescein	25 assays	APT423	\$155
CaspaTag Caspase-3,7 In Situ Assay Kit, Fluorescein	100 assays	APT403	\$435
CaspaTag Caspase-3,7 In Situ Assay Kit, Sulforhodamine	25 assays	APT523	\$185
CaspaTag Caspase-3,7 In Situ Assay Kit, Sulforhodamine	100 assays	APT503	\$490
CaspaTag Caspase-8 In Situ Assay Kit, Fluorescein	25 assays	APT428	\$155
CaspaTag Caspase-8 In Situ Assay Kit, Fluorescein	100 assays	APT408	\$440
CaspaTag Caspase-9 In Situ Assay Kit, Fluorescein	25 assays	APT429	\$155
CaspaTagCaspase-9 In Situ Assay Kit, Fluorescein	100 assays	APT409	\$440

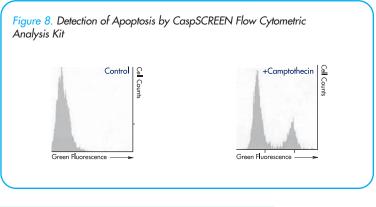
# CaspSCREEN Flow Cytometric Apoptosis Detection Kit

The Chemicon CaspSCREEN Flow Cytometric Apoptosis Detection Kit from Millipore provides a simple and convenient means for detecting active caspases by flow cytometry in non-adherent cells. The assay utilizes a molecule that contains rhodamine 110 linked to two aspartate residues (D2R), a reported substrate for the caspase protease family. The intact substrate is nonfluorescent; however, upon cleavage by caspases, the released rhodamine 110 gives rise to fluorescence that can be measured at Ex = 488 nm and Em = 530 nm. The D2R substrate is more cell-permeable than other fluorometric caspase substrates to effectively and easily detect caspase activity, and therefore, apoptosis.

## **CaspSCREEN** Advantages

Fast: Flow cytometry assay to detect all caspase activity Simple: One-step assay in live cells

**Effective:** Cell-permeable caspase substrate for use in live cells



APT105

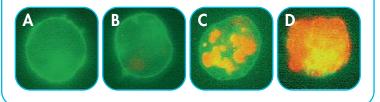
25 assays

\$395

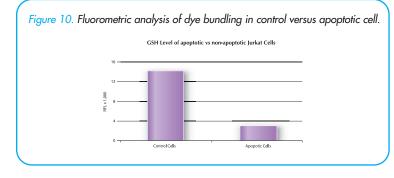
## **ApopNexin Apoptosis Detection Kits**

Chemicon ApopNexin Apoptosis Detection Kits from Millipore detect the appearance of PS in the outer leaflet of the membrane of the cell. The system uses Annexin V conjugated to either FITC or Biotin to allow for

Figure 9. Human peripheral blood lymphocytes induced with an anti-Fas antibody for 3-4 hours and then assayed with ApopNexin FITC and propidium iodide. Results obtained by fluorescence microscopy show that apoptotic cells have ApopNexin FITC bound to the PS on the outer membrane surface (Panel A). As apoptosis progressed, the plasma membrane becomes permeable, and the propidium iodide is able to enter into the cell and bind to cellular DNA (Panels B and C). Necrotic cells are detected by intense propidium iodide staining of the cytoplasm, due to complete disruption of the plasma membrane (Panel D).

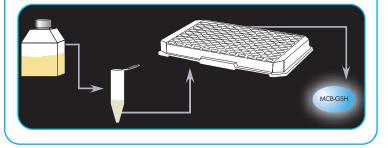


Description ApopNexin Biotin Apoptosis Detection Kit ApopNexin FITC Apoptosis Detection Kit



#### Figure 11: Glutathione Detection Kit – Protocol.

- Wash pellet and discard supernatant
- Resuspend pellet in lysis buffer
- Centrifuge and collect lysate
- Add MCB fluorochrome
- Fluorochrome bound to GSH fluoresces blue



convenient quantitative assays. The counterstain Propidium lodide is included to differentiate the apoptotic and necrotic cells.

ApopNexin FITC binding can be analyzed by flow cytometry or fluorescence microscopy in less than one hour. It may be used in conjunction with other fluorescent cell surface markers. ApopNexin Biotin can be used for chromogenic or fluorescence detection using a streptavidin conjugate of choice. This versatile approach also allows bicolor or tricolor fluorescence experiments for mutiliparameter cell surface analysis. Fixation can be performed after Annexin V binding for further analysis of other cellular antigens.

#### ApopNexin Advantages

Sensitive: Highly sensitive and specific biochemical results Fast: Quantitative results using flow cytometry Effective: Two labeling options

	Quantity	Cat. No.	Price
ptosis Detection Kit	100 tests	APT700	\$380
tosis Detection Kit	100 tests	APT750	\$380

# **Glutathione Detection Kit**

Reduced glutathione is an antioxidant in human tissues that removes reactive oxygen species from cells. The oxidized form of glutathione is returned to its reduced form by glutathione reductase. In cells undergoing apoptosis, the cytosolic level of reduced glutathione is decreased. Chemicon's Glutathione Detection Kit from Millipore measures the level of reduced glutathione (GSH) in the cytosol using a fluorometric indicator dye which binds preferentially to GSH. The unbound dye shows virtually no fluorescence, while the dye bound to GSH fluoresces blue. The blue fluorescence is detected using a standard fluorometer or fluorescence plate reader.

## **Glutathione Advantages**

Fast: Assay completed in a little over an hour Simple: Fluorescence detection using a standard fluorometer or fluorescence plate reader.

APT250

Glutathione Detection Kit

# MitoLight Mitochondrial Permeabilization Detection Kits

The Chemicon MitoLight Mitochondrial Apoptosis Detection Kit from Millipore is a quick and easy method for detecting changes in the mitochondrial membrane potential in live cells. Mitochondrial depolarization is a well-characterized event in apoptosis that can be detected using MitoLight. This lipophilic cationic dye stains living cell mitochondria in a membrane potential-dependent fashion. In healthy cells, the dye accumulates and aggregates in the mitochondria and fluoresces red (I em = 585-590 nm). During apoptosis, the electrochemical gradient across the mitochondrial membrane breaks down (presumably due to the formation of additional pores in the membrane), and the dye remains in the cytosol in its monomeric form and fluoresces green (I em = 527-530 nm).

The MitoLight Mitochondrial Apoptosis Detection Kit will work with adherent cells and cells in suspension. With cells in suspension, the analysis can be performed by fluorescence microscopy or by flow cytometry; with adherent cells, the analysis must be performed using fluorescence microscopy.

## **MitoLight Advantages**

Fast: Assay complete in 20 minutes

Simple: Incubate the cells with the MitoLight dye, wash and analyze

**Effective:** Accurate detection of changes in mitochondrial membrane potential

Description	Quantity	Cat. No.	Price
MitoLight Mitochondrial Apoptosis Detection Kit	25 assays	APT142	\$170
MitoLight Mitochondrial Apoptosis Detection Kit	100 assays	APT242	\$305

# **DNA Ladder Detection Kit**

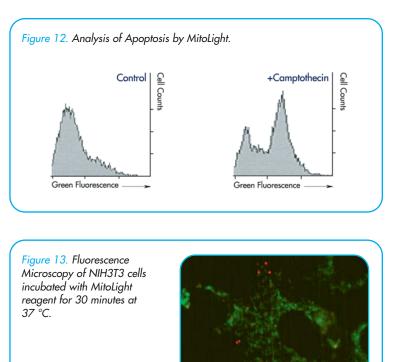
The Chemicon DNA Ladder Kit from Millipore provides a fast, simple method for the detection of apoptotic cells. The procedure involves lysis of the cells followed by sequential digestion of RNA and proteins. The genomic DNA is then precipitated, resuspended and separated on an agarose gel. Unlike other kits, our method requires less than 90 minutes to prepare DNA in a single tube without the need for extraction or expensive columns. DNA fragmentation is easily visualized by agarose gel electrophoresis. The assay procedure also increases recovery of small fragmented DNA, resulting in greater sensitivity.

APT151

## **DNA Ladder Advantages**

**Fast:** Requires less than 90 minutes **Simple:** Single tube, straightforward detection of apoptotic cells by agarose gel

\$280



# ApopTag Peroxidase ISOL Apoptosis Detection Kit

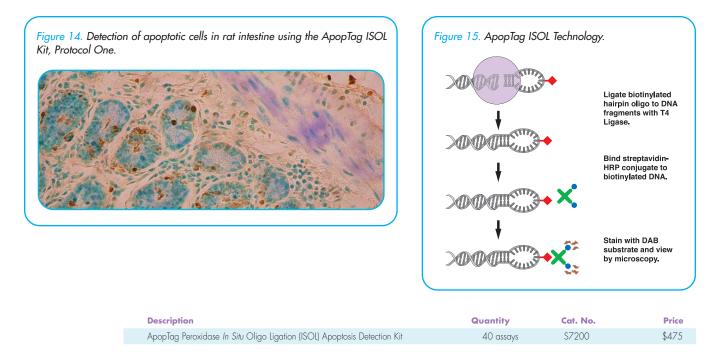
The Chemicon ApopTag Peroxidase ISOL Apoptosis Detection Kit from Millipore is the newest member of the ApopTag family of Apoptosis Detection Kits. Like the original ApopTag kits, the ApopTag ISOL Kit detects DNA fragmentation by end-labeling the DNA. However, this kit uses a novel "In Situ Oligo Ligation (ISOL)" method to label the ends of DNA fragments that takes advantage of the prevalence of blunt-ends and 3' single-base overhangs characteristic only to apoptotic cells. This allows the ApopTag ISOL Kit to clearly differentiate cells undergoing apoptosis from cells undergoing necrosis (see Figure 14).

The ApopTag ISOL Kit procedure (shown in Figure 15) attaches a specific biotinylated, synthetic oligonucleotide to the fragmented DNA *in situ*, provided that the DNA has a blunt end or a single base overhang on its 3' end. Either Oligo A (3' overhang) or Oligo B (blunt end) will bind to an exact complementary, doublestranded DNA end in a reaction catalyzed by T4 DNA Ligase. Then, a streptavidin-peroxidase conjugate is added that binds tightly to the biotin on the oligo. Lastly, the addition of DAB (diaminobenzidine), a chromogenic peroxidase substrate, causes a brown precipitate that can be clearly visualized using brightfield microscopy.

## ApopTag Peroxidase ISOL Advantages

Innovative: Novel labeling method that clearly differentiates apoptotic from necrotic cells Sensitive: Minimizes background through high specificity of *in situ* labeling

Accurate: Reduces the incidence of false positives Flexible: Detects apoptotic cells in paraffin-embedded tissues, fixed adherent cells, frozen tissues and cell suspensions



# Apo-Direct Flow Cytometry Apoptosis Detection Kit

The Chemicon Apo-Direct Kit from Millipore is a single-step staining method for labeling DNA breaks to detect apoptotic cells by flow cytometry. The kit contains all the reagents required for measuring apoptosis in cells including positive and negative control cells; washing, reaction and rinsing buffers; and propidium iodide/RNase A solution for counter staining the total DNA.

# ApopTag TUNEL Apoptosis Detection Kits

One of the hallmarks of apoptosis is DNA fragmentation. Chemicon's ApopTag Apoptosis Detection Kits from Millipore detect apoptotic cells in situ by specific end-labeling of those DNA fragments using the TUNEL method. Nucleotides labeled with either digoxigenin or fluorescein (see Figure 16) are enzymatically added to 3'-hydroxyl DNA ends by terminal deoxynucleotidyl transferase (TdT). This enzyme is more selective for DNA fragments from apoptotic rather than necrotic cells and is more specific than DNA polymerase. The incorporated nucleotides form a random heteropolymer in a ratio that is optimized to promote anti-digoxigenin antibody binding or to minimize fluorescein self-quenching. The anti-digoxigenin antibody fragment carries either a conjugated reporter enzyme (peroxidase, see Figure 17) or one of two fluorescent molecules, fluorescein or sulforhodamine, to the reaction site. The localized peroxidase enzyme catalytically generates an intense signal from chromogenic substrates that can be observed using light microscopy. Alternatively, the bound fluorescein or sulforhodamine may be observed by flow cytometry or fluorescence microscopy. (Analysis of sulforhodamine by flow cytometry requires the use of special filters in the flow cytometer.)

In the direct labeling strategy (used in Cat. No. S7160), the antibody detection steps are eliminated, and nucleotides labeled with fluorescein are directly incorporated into the DNA. These specimens may be analyzed by flow cytometry or fluorescent microscopy.

#### ApopTag Advantages

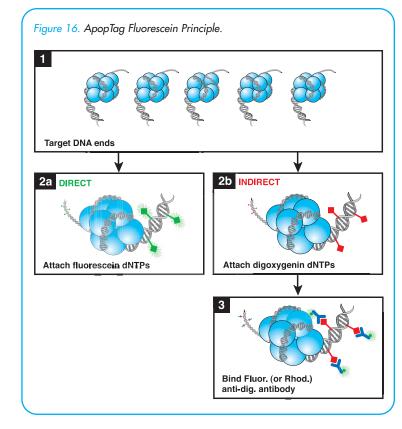
**Flexible:** Choice of detection methods (flow cytometry or microscopy)

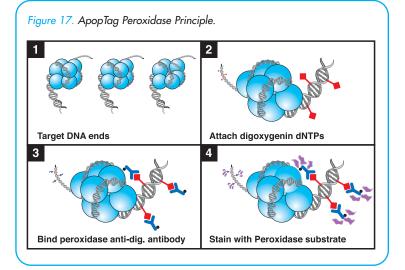
**Versatile:** Peroxidase, fluorescein or rhodamine kits available

Accuracy: Control slides available to monitor and confirm accuracy

Reliable: Hundreds of citations in peer-reviewed journals

Description	Quantity	Cat. No.	Price
ApopTag Peroxidase In Situ Apoptosis Detection Kit	40 assays	S7100	\$380
ApopTag Plus Peroxidase In Situ Apoptosis Detection Kit	40 assays	S7101	\$430
ApopTag Fluorescein In Situ Apoptosis Detection Kit	40 assays	S7110	\$405
ApopTag Plus Fluorescein In Situ Apoptosis Detection Kit	40 assays	S7111	\$415
ApopTag Fluorescein Direct In Situ Apoptosis Detection Kit	40 assays	S7160	\$385
ApopTag Red In Situ Apoptosis Detection Kit	40 assays	S7165	\$390



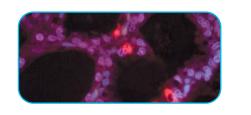


#### Table 3: ApopTag TUNEL Kit Components

Description	Cat. No.	Buffers* & Coverslips	Blocking Solution	Anti- Digoxigenin- Peroxidase Antibody	Anti- Digoxigenin- Fluorescein Antibody	Anti- Digoxigenin- Rhodamine Antibody	Positive Control Slides	DAB Substrate
ApopTag Peroxidase	S7100	•		•				
ApopTag PLUS Peroxidase	S7101	•		•			•	•
ApopTag Fluorescein	\$7110	•	•		•			
ApopTag PLUS Fluorescein	\$7111	•	٠		٠		•	
ApopTag Fluorescein Direct	S7160	٠						
ApopTag Fluorescein	\$7165	•				•		

\*Equilibration Buffer, Reaction Buffer, TdT Enzyme, Stop/Wash Buffer

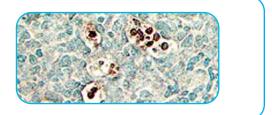
Figure 18. Detection of apoptotic cells with ApopTag Red in human lymph node tissue, fixed with 10% neutral-buffered formalin and paraffin-embedded.



# Individual Kit Components

Description	Quantity	Cat. No.	Price
ApopTag Equilibration Buffer	15 mL	S7105	\$115
ApopTag Plastic Coverslips	1000 ea	S7117	\$255
ApopTag Positive Control Slides	5 slides	S7115	\$65
ApopTag Reaction Buffer	1 mL	S7105	\$115
ApopTag Stop/Wash Buffer	20 mL	S7108	\$49
ApopTag TdT Enzyme	300 µL	S7107	\$115

Figure 19. Detection of apoptotic cells in human lymph node with ApopTag Peroxidase. The tissue was fixed in 10% neutral-buffered formalin and paraffin-embedded.



## **Ancillary Products**

/			
DAPI/Antifade Solution	1 mL	S7113	\$60
Propidium lodide Solution	1 mL	S7109	\$60
Propidium Iodide/Antifade Solution	1 mL	S7112	\$60

# CleavaLite Caspase-3 Activity Assay Kits

Chemicon's CleavaLite Caspase-3 Activity Assay Kit from Millipore provides a highly sensitive assay employing CleavaLite, a novel bioluminescent substrate specific for Caspase-3. CleavaLite is a Renilla luciferase mutant containing the Caspase-3 cleavage site, DEVD. Upon cleavage, it exhibits significantly decreased bioluminescence relative to the amount of Caspase-3. As shown in Figure 20, CleavaLite is highly specific for Caspase-3.

## **CleavaLite Caspase-3 Advantages**

Sensitive: 50 to 100 times more sensitive than standard colorimetric assays Fast: Obtain results in about 2 hours Convenient: 96-well or 384-well format

Highly Specific: Only detects Caspase-3

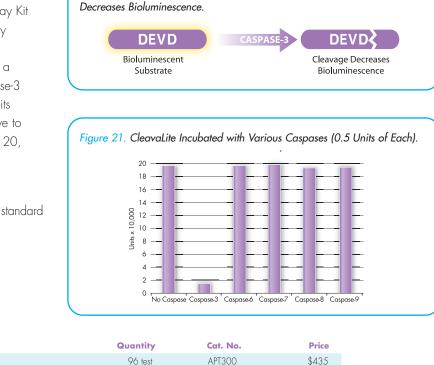


Figure 20. Cleavage of Bioluminescent Substrate by Caspase-3

Cleavalite Caspase 3 Activity Kit Cleavalite Caspase 3 Activity Kit

Description

# Other Caspase Activity Assay Kits

Caspase Colorimetric and Fluorometric Kits provide a simple means for assaying caspase activity. The colorimetric assay is based on spectrophotometric detection of a chromophore following cleavage from a labeled substrate. The fluorometric assay is based on detection of cleavage of a fluorescently labeled substrate. Comparison of the colorimetric or fluorescent signal from an apoptotic sample with an uninduced control allows determination of the fold increase in caspase activity.

APT301

\$550

384 test

Caspase 1 Colorimetric	100 assays	APT161	\$475
Caspase 1 Fluorometric	100 assays	APT160	\$500
Caspase 2 Colorimetric	100 assays	APT163	\$475
Caspase 2 Fluorometric	100 assays	APT162	\$500
Caspase 3 Colorimetric	25 assays	APT131	\$205
Caspase 3 Colorimetric	100 assays	APT165	\$390
Caspase 3 Fluorometric	130 assays	17-198	\$349
Caspase 5 Colorimetric	100 assays	APT167	\$475
Caspase 5 Fluorometric	100 assays	APT166	\$500
Caspase 6 Colorimetric	100 assays	APT169	\$475
Caspase 6 Fluorometric	100 assays	APT168	\$500
Caspase 8 Colorimetric	25 assays	APT129	\$205
Caspase 8 Colorimetric	100 assays	APT171	\$390
Caspase 8 Fluorometric	100 assays	APT170	\$500
Caspase 9 Colorimetric	100 assays	APT173	\$390
Caspase 9 Fluorometric	100 assays	APT172	\$500
Caspase 10 Colorimetric	100 assays	APT176	\$475
Caspase 10 Fluorometric	100 assays	APT174	\$500

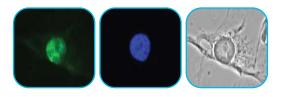
# Apoptosis and Histones H2A.X and H2B

# The Role of Histone Modifications in Apoptosis and DNA Damage

Many key scientific discoveries have been made that link specific histone modifications with important biological phenomena. In particular, several histone modifications have been discovered that are useful biomarkers of apoptosis and DNA damage. Reagents and kits specific for these modifications are of great value to any researcher studying these processes.

Apoptosis is defined as a growth-limiting regulatory mechanism by which cells can trigger their own demise due to advanced age in response to extracellular signals or as a result of irreparable cellular or DNA damage.

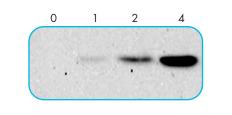
Figure 22: H2A.X Phosphorylation During Apoptosis. Immunofluorescence of HeLa cells using Anti-phospho-Histone H2A.X (Ser139), clone JBW301 (Cat. No. 05-636). Cells were treated with 1 µg/ml Staurosporine for two hours to induce DNA damage and apoptosis. Left panel, Anti-phospho-H2A.X (Ser139), FITC conjugate (Cat. No.16-202A). Middle panel, DNA stained with DAPI. Right panel, phase contrast image.



Staurosporine

Many key regulators of apoptosis are critical to cancer development, as the function of one or more apoptotic inducers (e.g., p16, p53, BRCA1) is lost in most types of cancer. The loss of these key growth checkpoint proteins allow cells to become immortalized by escaping important regulatory growth restrictions. Additionally, some genes that normally function to block premature apoptosis are hyperactivated in specific cancers (Bcl2), thus preventing cells from terminating their growth. Apoptosis is also an important developmental mechanism, such as preventing the overgrowth of particular neuronal cell lineages in the developing brain.

Figure 23: H2A.X Phosphorylation Western Blot. Western blot using Anti-phospho-Histone H2A.X (Ser139), clone JBW301 (Cat. No. 05-636) antibody at 1:2000 dilution to detect accumulation of g-H2A.X in Jurkat cells treated with 1 µg/ml staurosporine to induce apoptosis. Cells were harvested after the number of hours indicated above each lane.



# Apoptosis and DNA Damage Related Products

Antibodies			
Description	Quantity	Cat. No.	Price
Anti-phospho-Histone H2A.X (Ser139), clone JBW301	200 µg	05-636	\$349
Anti-phospho-H2A.X (Ser139)	200 µg	07-164	\$319
Anti-phospho-Histone H2B (Ser14)	200 µL	07-191	\$319
Anti-phospho-Histone H2B (Ser14), clone MC603	100 hã	05-751	\$329
Anti-phospho-Histone H2A.X (Ser139), clone JBVV301, biotin conjugate	100 ha	16-193	\$359
Active Caspase Detection Kits			
TUNEL Apoptosis Detection Kit	1 kit	17-141	\$475
H2A.X Phosphorylation Assay Kit (Chemiluminescence Detection)	1 kit	17-327	\$509
H2A.X Phosphorylation Assay Kit (Flow Cytometry)	1 kit	17-344	\$429
Antibodies to Apoptosis Specific Cleavage Products			
Fas Ligand, Membrane Bound	1 vial	01-210	\$309

200 µg

19-123

\$179

# H2A.X Phosphorylation as a Biomarker for DNA Damage and Apoptosis

Many agents, such as ionizing radiation and DNAbinding drugs, cause damage to nuclear DNA - the most severe type being double-strand breakage (DSB). DSBs must be repaired quickly and with high fidelity, as they can lead to rearrangements in the genome through recombination. Specific protein complexes exist within the nucleus for just this purpose. The process of apoptosis occurs in successive stages, each with characteristic changes in cell morphology. Along with these changes, specific post-translational modifications occur on the histones. The histone variant H2A.X is rapidly phosphorylated at serine 139 by the ATM kinase in response to even a few double strand DNA breaks. The accumulation of high levels of H2A.X phosphorylation resulting from severe DNA damage is a very early and accurate indicator of apoptosis. Interestingly, H2A.X phosphorylation also occurs when chromosomal DNA is cleaved intentionally, such as during meiotic recombination between sister chromatids and immunoglobulin gene VDJ recombination. The detection of H2A.X phosphorylation is illustrated in Figure 25.

#### References

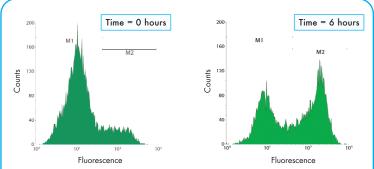
- 1. Rogakou, E. P. et al., J. Biol. Chem., 273: 5858-5868, 1998.
- 2. Rogakou, E. P. et al., J. Cell. Biol., 146: 905-916, 1999.
- 3. Talasz, H. et al., Cell Death Differ., **9**: 27-39, 2002.
- 4. Rogakou, E. P. et al., J. Biol. Chem., 275: 9390-9395, 2000.
- 5. Paull, T. T. et al., Curr. Biol., 10: 886-895, 2000.

# H2B Phosphorylation as a Biomarker of Irreversible Commitment to Apoptosis

Another modification, phosphorylation of serine 14 on histone H2B, occurs at the stage in apoptosis when the condensed nuclear DNA is cleaved by caspaseassociated DNase enzymes. It is a hallmark of a cell irrevocably committed to the apoptotic pathway. H2A.X phosphorylation is also observed during this stage as a result of the DNase-induced DNA breaks. A third phosphorylation event linked to apoptosis, at serine 32 of histone H2B, has also been reported. Upstate antibodies from Millipore that recognize these modifications, by themselves or as part of kits, are extremely useful in monitoring DNA damage and apoptosis.

#### References





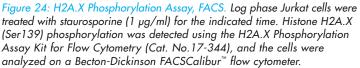


Figure 25. Detection of Histone H2B phosphorylated at serine 14 with Antiphospho-Histone H2B (Ser14) (Cat. No. 07-191) by immunofluorescence in HL-60 cells stimulated to undergo apoptosis by treatment with VP-16. Top panel, Anti-phospho-Histone H2B (Ser14) (Cat. No. 07-191) staining. Bottom panel, DAPI staining. Note the distinctive morphology of the nuclear DNA as it breaks down, concomitant with the detection of H2B serine 14 phosphorylation (arrows). Courtesy David Allis, The Rockefeller University.

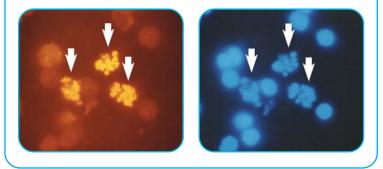
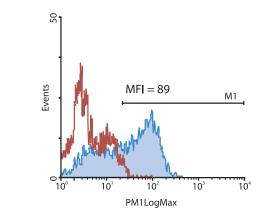


Figure 26. HL-60 cells were cultured with 20 µg/ml of etoposide for 18 hours. Cells were aliquoted at 3 X 10<sup>5</sup> per well into a 96-well round-bottom microtiter plate, permeabilized with a buffer containing Tween<sup>®</sup>-20 and stained with 20 µg/ml of Anti-phospho-Histone H2B (Ser14), clone MC603 (Cat. No. 05-751) (filled histogram) or 20 µg/ml of Normal Rabbit IgG (Cat. No. 12-370) (open histogram) followed by a goat anti-rabbit IgG (H+L) phycoerythrin conjugate. Data was acquired and analyzed using a Guava<sup>®</sup> PCA-96 System.



# Biotools Spotlight

# Visualizer™

## A Flexible Family of Western Blot Detection Kits – Discover More and Publish Faster™

Every assay is different. Whether you need a quality reagent for routine detection or require a high sensitivity kit for low protein concentrations, Millipore offers the perfect choice. With the right reagent, you can increase your signal and decrease your background so that you can discover more and publish faster.

#### Low cost, everday use reagents

Upstate Visualizer EC Western Blot Detection Kits from Millipore are designed with a high glow reaction chemiluminescent substrate with outstanding sensitivity, long-lasting signal and low cost. Visualizer EC offers the highest signal-to-noise ratios of any existing luminol peroxidase formulations on the market, resulting in high sensitivity and low backgrounds at an economical price (see Figure 27 below).

Visualizer Spray & Glow<sup>™</sup> is a simple, but novel, enhanced chemiluminescent (ECL) detection reagent with the "at-home" convenience of a spray bottle! Spray the Western membrane and detect your target protein using standard detection methods, without the mixing of substrate and enhancer solutions. The new Upstate Visualizer Spray & Glow ECL Western Blotting

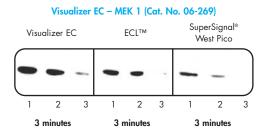


Detection System from Millipore will revolutionize Western blotting detection by putting everything you need for a high glow reaction chemiluminescent substrate together in one bottle.

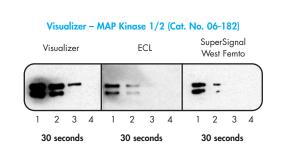
#### High sensitivity reagents for difficult to detect proteins

Upstate Visualizer Western Blot Detection Kits from Millipore, mouse and rabbit, are designed to make Western detection of proteins easier by providing the highest level of sensitivity with the least amount of background. This means less effort optimizing exposure times, and for hard to detect proteins needing long exposures, significantly lower background (see Figure 27 below). Visualizer Western Blot Detection Kits utilize a superior version of the chemiluminescent HRP substrate luminol that results in the fastest and most sensitive detection, a long-lasting signal and the highest signal-tonoise ratio. Kits include chemiluminescent substrates paired with an HRP-conjugated secondary antibody to either rabbit or mouse IgG. Combined with Millipore's extensive line of Upstate and Chemicon primary antibodies, Visualizer Western Blot Detection Kits provide you with the quality results you require for your research. For more information, please visit www.millipore.com/upstate.

#### Figure 27: Western Blot Detection Kit Product Comparison.



Visualizer EC Western Blot: A 4-12% Bis-Tris SDS-PAGE gel in MOPS buffer was loaded and electrophoresed with HeLa Cell Lysate (Cat. No. 12-501) in a dilution series; 10 µg, 5 µg and 2 µg (lanes 1-3 respectively), and transferred to PVDF membrane. The membrane was cut, and antibody incubations were performed according to each reagent manufacturers' guidelines. Each blot was incubated in the detection reagent as listed and laid side-by-side for 3 minutes exposure to film.



Visualizer Western Blot: A 4-12% Bis-Tris SDS-PAGE gel in MOPS buffer was loaded and electrophoresed with Jurkat Cell Lysate (Cat. No. 12-303) in a dilution series; 500 ng, 250 ng, 100 ng and 50 ng (lanes 1-4 respectively), transferred to PVDF membrane. The membrane was cut, and antibody incubations were performed according to each reagent manufacturers' guidelines. Each blot was incubated in the detection reagent as listed and laid side-by-side for 30 seconds exposure to film.

# Visualizer Family of Western Blot Detection Kits

Visualizer Spray & Glow Products	Quantity	Cat. No.	Price
New Visualizer Spray & Glow ECL Western Blotting Detection System	40 mL Spray Bottle (20 mini-gel-sized blots)	1 <i>7-</i> 373SP	\$59
New Visualizer Spray & Glow ECL Western Blotting Detection System	100 mL Spray Bottle (50 mini-gel-sized blots)	17-373	\$149
New Visualizer Spray & Glow ECL Western Blotting Detection System	3 X 100 mL Spray Bottles (150 mini-gel-sized blots)	1 <i>7-</i> 373BP	\$359

### **Visualizer EC Products**

New Visualizer EC Western Blot Detection Kit	5 mini-blots	64-301SP	\$49
New Visualizer EC Western Blot Detection Kit	20 mini-blots	64-301	\$99
New Visualizer EC Western Blot Detection Kit	50 mini-blots	64-301BP	\$149

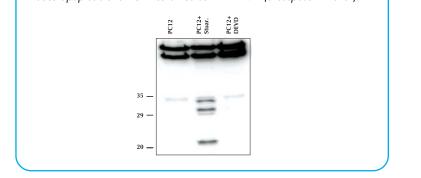
#### **Visualizer Products**

Visualizer Western Blot Detection Kit, mouse	5 mini-blots	64-201SP	\$95
Visualizer Western Blot Detection Kit, mouse	20 mini-blots	64-201	\$249
New Visualizer Western Blot Detection Kit, mouse	50 mini-blots	64-201BP	\$359
Visualizer Western Blot Detection Kit, rabbit	5 mini-blots	64-202SP	\$95
Visualizer Western Blot Detection Kit, rabbit	20 mini-blots	64-202	\$249
New Visualizer Western Blot Detection Kit, rabbit	50 mini-blots	64-202BP	\$359

# Apoptosis and Neuroscience

A prominent feature of Alzheimer's Disease is the formation of senile plaques, composed mainly of Amyloid- $\beta$ , in selected regions of the brain. Amyloid- $\beta$  is neurotoxic at high concentrations, and its precursor, Amyloid- $\beta$  precursor (APP), is directly and efficiently cleaved by caspases 3, 6, 8 and 9 during apoptosis (Barnes et al., 1998; Gervais et al., 1999; LeBlanc et al., 1999; Pelligrini et al., 1999; Weidemann et al., 1999; Lu et al., 2000). In vitro, Amyloid- $\beta$ -induced apoptosis is mediated by the activation of caspase-3 (Marin et al., 2000). The proteolytic cleavage of APP by caspase-3 generates a novel epitope at the carboxy-terminal part of the APP protein. In dying neurons of Alzheimer's Disease brains, the levels of caspase-3 are elevated and there is co-localization of the caspase-3-generated APP cleavage product and Ab in senile plaques (Gervais et al., 1999).

Figure 28. Rabbit anti-Caspase-Cleaved Amyloid Precursor Protein (Cat. No. AB5942) Western blot of PC12 cells, PC12 cells treated with Staurosporine to induce apoptosis and PC12 cells treated with DEVD (a caspase inhibitor).



Caspase-cleaved Amyloid B Precursor Protein	100 hð	AB5942	\$320
Amyloid Precursor Protein, C-terminus	100 µL	AB5352	\$240
Amyloid Precursor Protein, Universal	100 µL	AB5300	\$240

# Apoptosis Products

Note: Additional species and applications may apply. Call Tech Support at 800 437 7500 for more information.

#### **Tested Applications**

Abbr.	Description	Abbr.	Description	Abbr.	Description
ABA	Affinity Binding Assay	HDAC	Histone Deacetylase Assay	NEUT	Neutralizing
ACT	Activity Assay	HI	Hemagglutination Inhibition	NT	Nitration
ADH	Stimulates ECM Adhesion	HMT	Histone Methyltransferase Assay	NUEX	Nuclear Extraction
Al	Agonist or Inhibitor	IAP	Immunoaffinity Purification	PA	Phosphatase Assay
AMP	DNA Amplification	IC	Immunocytochemistry (Cells)	PC	Positive Control
APA	Affinity Precipitation Assay	ID	Immunodiffusion	PCU	Protein Clean-up
BD	Beadlyte® Assay	IEP	Immunoelectrophoresis	PD	Protein Determination
CA	Caspase Assay	IF	Immunofluorescence	PIA	Peptide Inhibition Assay
CC	Culture Confirmation	IFIX	Immunofixation	RIA	Radioimmunoassay
ChIP	Chromatin Immunoprecipitation	IH	Immunohistochemistry (Tissue)	RNAi	RNAi/siRNA/Gene Knockdown
CULT	Cell Culture	IH(P)	Immunohistochemistry (Paraffin)	RPA	Ribonuclease Protection Assay
DB	Dot Blot	IND	Induces Function	RT-PCR	Reverse Transcriptase Polymerase Chain Reaction
EA	Enzyme Assay	INHIB	Inhibits Activity/Function	SW	Software Needed
ELISA	Enzyme Immunoassay (ELISA)	IP	Immunoprecipitation	TFX	Transfection
EM	Electron Microscopy	IPK	IP-Kinase Assay	UC	Uncharacterized Antiserum
EMSA	Electrophoretic Mobility Shift Assay	IPX	Immunoperoxidase Staining	WB	Immunoblotting (Western)
FC	Flow Cytometry (FACS)	IRMA	Immuno Radio-Metric Assay	Web*	Important additional product reactivity
FUNC	Affects Function	IT	Immunotoxin		information available on datasheet
GPA	G-Protein Assay	KA	Kinase Assay		
HA	Hemagluttination	LFA	Lateral Flow Assay		
HAT	Histone Acetyltransferase Assay	NB	Northern Blot		

#### **Tested Species Reactivity**

Abbr.	Description	Abbr.	Description	Abbr.	Description
A	All Species	Gr	Gerbil	R	Rat
Am	Amphibian	Gs	Ground Squirrel	Rb	Rabbit
As	Aspergillus	Gt	Goat	Rc	Raccoon
Av	Avian	Н	Human	rH	Recombinant Human Protein
В	Bovine	H-sp	Human Only	Rp	Reptilian
Bab	Baboon	Ht	Hamster	Sal	Salamander
Bact	Bacterial	In	Insect	Seal	Seal
Bat	Bat	lnv	Invertebrates	Sh	Sheep
Ca	Canine (Dog)	Kn	Kangaroo	Shk	Shark
Ch	Chicken	lg	ligia	SHm	Syrian Hamster
Chp	Chimpanzee	Lz	Lizard	Shp	Shrimp
Crb	Crab	M	Mouse	Sj	Schistosoma japonicum
Crf	Crawfish	Ma	Mammals	Sn	Snail
Di	Dictyostelium	Md	Mule Deer	Snk	Snake
Dr	Drosophila	Mi	Mink	Spd	Spider
Ec	E. coli Bacteria	Mk	Monkey	Sqd	Squid
Ech	Echinoderms	M	Mollusk	Su	Sea Urchin
Ecl	Enterobacter cloacae	Nem	Nematode	Т	Tetrahymena
Elk	Elk	Nr	Neurospora crassa	Vo	Vole
Eq	Equine (Horse)	Op	Opposum	Vrt	Vertebrates
Eu	Eukaryote	Ox	Ox	VVR	Most common vertebrate species tested
F	Fish	Pl	Green Plants	Xn	Xenopus
Fe	Feline (Cat)	Pm	Primate	Y	Yeast (S. cerevisiae)
Fg	Frog	Pn	Penicillium	Zf	Zebra Fish
Ft	Ferret	Po	Porcine (Pig)		
Gp	Guinea Pig	Qu	Quail		

<b>Description</b> t/PKB (see page 8 for a product spotlight on Akt/PKB)	Species	Applications	Host	Quantity	Cat. No.	Price
Anti-Akt/PKB, PH Domain, clone SKB1	HMR	WB IP IPK FC	M IgG	100 µg	05-591	\$329
Anti-Akt/PKB, PH Domain, agarose conjugate	HMR	IP IPK		100 µg	16-185	\$329
kt1/PKBα				. o o pg	10 100	φ0 <i>2</i> ,
Anti-Akt1/PKBα, PH domain, polyclonal	НM	WB	Rb IgG	100 µL	06-885	\$319
Anti-Akt1/PKBα	H M R	WB IPK	Rb IgG	100 µg	07-416	\$329
Anti-Akt1/PKBα, clone AVV24	HMR	WB IP	Rb IgG	100 pg 100 µL	05-796	\$319
Anti-phospho-Akt1/PKBα (Thr308)	M	WB IH	Rb IgG	100 pL	06-678	\$319
Anti-phospho-Akt1/PKB $\alpha$ (Ser473)	HM	WB	Rb IgG	200 µg	07-310	\$309
Anti-Akt1, phospho-specific (Ser473)	HMR	WB ELISA	Rb	200 рд 50 µд	AB3132	\$280
Anti-phospho-Akt1/PKB $\alpha$ (Thr308), clone NL50	HM	WB	Rb IgG	100 µL	05-802	\$319
Anti-phospho-Akt1/PKB $\alpha$ (Ser473), clone 11E6	НМ	WB	M IgG <sub>1x</sub>	100 µg	05-669	\$329
Anti-phospho-Akt1/PKB $\alpha$ (Ser473), clone SK703	НМ	WB	Rb IgG	100 µg	05-736	\$349
Akt1/PKB $\alpha$ Immunoprecipitation-Kinase Assay Kit		IP KA		1 kit	17-188	\$419
Akt1/PKBa SMARTpool <sup>®</sup> siRNA reagent	Н	RNAi		5 nmol	M-003000	\$375
siRNA plasmid, pKD™-Akt1/PKBα-v2	Н	RNAi		5 µg	62-041	\$349
Akt1/PKB $\alpha$ cDNA (dominant negative) Expression Kit		TFX		1 kit	17-252	\$699
Akt1/PKB $\alpha$ cDNA (activated) Expression Kit		TFX		1 kit	17-253	\$699
Akt1/PKB $\alpha$ cDNA Allelic Pack		TFX		1 kit	17-254	\$1,369
Akt1/PKB $\alpha$ cDNA (activated) in pUSEamp		TFX		5 µg	21-151	\$599
Akt1/PKB $\alpha$ cDNA (dominant negative) in pUSEamp		TFX		5 µg	21-152	\$599
Akt1/PKB $\alpha$ cDNA (wt) in pUSEamp		TFX		5 µg	21-153	\$599
New Anti-phospho-Akt1 (Thr34)	Н	WB ELISA	Rb IgG	100 µL	07-789	\$309
			0			
kt1/PKBβ	HR		Rb IgG	100 µL	05 771	¢ つ 1 (
Anti-Akt2/PKBβ, clone AW114	H M R	VVB IP VVB IP	U U	1	05-771 07-372	\$319 \$319
Anti-Akt2/PKBβ Akt2/PKBβ <i>SMART</i> pool siRNA reagent	H	RNAi	Rb IgG	200 µg 5 nmol	M-003001	\$375
siRNA plasmid, pKD-Akt2/PKBβ-v1	Н	RNAi		5 µg	62-103	\$349
				o pg	02 100	ψ04 γ
kt3/PKBγ						
Anti-Akt3/PKBg	Н	VVB IPK	Rb IgG	100 µL	07-383	\$329
Anti-Akt3, N-terminus	H M R	WB ELISA	Rb	50 µg	AB3136	\$260
Anti-Akt3/PKBg, clone GMA104	Н	VVB IP	$M \log_{2a}$	100 hð	05-780	\$319
Akt3/PKBg SMARTpool siRNA reagent	Н	RNAi		5 nmol	M-003002	\$375
siRNA plasmid, pKD-Akt3/PKBg-v2	Н	RNAi		5 µg	62-138	\$349
RAS40 (Proline-Rich Akt Substrate)						
New Anti-PRAS40	Н	WB IP	M lgG <sub>1ĸ</sub>	200 µL	05-988	\$309
New Anti-phospho-PRAS40 (Thr246)	НM	WB	Rb IgG	100 µL	07-888	\$309
			0	·		
nnexin				100		¢00
Anti-Annexin I, clone 6E4/3	Н	WB ELISA IF	M IgG1	100 µg	MAB3773	\$233
Anti-Annexin II, clone 3D5/4	Н	IP ELISA IH	Μ lgG <sub>2aκ</sub>	100 µg	MAB3774	\$235
Apopnexin FITC Apoptosis Kit	Ma			100 assays	APT750	\$380
Apopnexin Biotin Apoptosis Kit	Ma			100 assays	APT700	\$380
paf-1						
Anti-Apaf-1, C-terminus	HMR	WB	Rb	100 µg	AB16503	\$220
Anti-Apaf-1, N-terminus	НМ	VVB	Rb	100 µg	AB16941	\$220
Anti-Apaf-1, clone 2E12	Н	WB IP ELISA FC IH	R IgG <sub>2ak</sub>	100 hð	MAB3503	\$220
Anti-Apaf-1, clone 13F11	Μ	WB IP ELISA FC	R IgG <sub>2ak</sub>	100 hð	MAB3504	\$263
Anti-Apaf-1, clone 18H2	ΗM	WB IP ELISA FC	R IgG <sub>2ak</sub>	100 µg	MAB3505	\$263
F (Apoptosis Inducing Factor)						
Anti-AIF	HMR	WB	Rb IgG	200 µg	07-208	\$299
Anti-AIF, internal domain	H M R	WB IH(P) Web*	Rb	100 µg	AB16501	\$220
		WB	Rb	100 µg	AB16502	\$220
Anti-AIF, N-terminus	Н	VVD	ND .	roo pq		
Anti-AIF, N-terminus siRNA plasmid, pKD-AIF-v3	H	RNAi	KD.	5 µg	62-128	\$349

APRIL	Description	Species	Applications	Host	Quantity	Cat. No.	Price
7 di Idiz	Anti-APRIL	Н	WB	Rb	100 hð	AB3635	\$205
ARC							
	Anti-ARC, N-terminus	H M R	WB	Rb	100 µg	AB16504	\$220
	Anti-ARC, C-terminus	Н	WB	Rb	100 µg	AB17005	\$220
	Anti-ARC	HMR	WB	Rb	100 µL	AB4517	\$195
	ARC, control peptide for AB16504		PIA			AG679	\$145
Arto							
Arts	Anti-Arts	Н	WB	Rb	50 µL	AB4512	\$195
4.01.0							• · · -
AS160	J Anti-AS160 (Rab GAP)	H M R	WB		200	07-741	\$269
-		H M R	WB	Rb IgG	200 µL		\$209 \$309
New	Anti-phospho-AS160 (Thr642)		VVD	Rb IgG	100 µL	07-802	\$20A
ASC							
	Anti-ASC	H WB		Rb	100 µg	AB3607	\$220
ASK1							
	Anti-ASK1	H WB		Rb IgG	200 µg	07-302	\$309
	Anti-ASK-1, C-terminus	H WB		Rb	100 µg	AB16505	\$220
Aven							
Aven	Anti-Aven	H,M	WB	Rb IgG	200 µg	07-640	\$309
Bad					100 1	120005	+005
	Anti-Bad	Н	WB IP IC	Rb	100 µL	AB2925	\$205
	Anti-Bad, clone BYC001	НМ	WB IP	M IgG	200 µg	05-605	\$329
	Anti-phospho-Bad (Ser 112)	H		Rb	100 µL	AB3569	\$315
-	Anti-phospho-Bad (Ser128)	H		Rb	100 µL	AB3567	\$315
New	Anti-phospho-Bad (Ser 136)	Н		Rb	100 µL	AB3571	\$315
	Anti-phospho-Bad (Ser155), clone JBW101	H M R	WB	$M \ IgG_{2b}$	100 µL	05-628	\$319
	Anti-Bad, phospho-specific (Ser155), clone 27AT381	НМ	WB	M lgG1	100 hð	MAB3730	\$295
	Bad, agarose		KA		100 hB	14-281	\$299
	Bad, soluble		KA		100 hð	14-357	\$319
	siRNA plasmid, pKD-Bad-v3		RNAi		5 µg	62-243	\$349
Baff							
	Anti-BAFF, C-terminus	HMR	WB	Rb	100 µg	AB16530	\$220
	BAFF, recombinant human				20 µg	GF119	\$220
Bag							
bug	Anti-BAG-1, clone 3.10G3E2	Н	WB IH	M IgG1	50 µg	MAB4611	\$210
							<b>4</b> = 1 =
Bak						01 501	****
	Anti-Bak, NT	НМ	WB IP IH	Rb IgG	200 µg	06-536	\$329
	APOPTOPAK, (Anti-Bcl2, Bak, Bax) Miniature Set		WB IP IC IH		1 kit	17-178	\$275
BAR							
	Anti-BAR	Н	WB IP IH	Rb	100 µL	AB3530	\$205
	Anti-BarH2	НМ	IH	Rb	100 µg	AB5720	\$245
	Anti-Barx 1	НМ	WB IC IH	Rb	100 hð	AB5825	\$245
Bax							
	Anti-Bax, NT	НМ	WB IP IC IH	Rb IgG	200 µg	06-499	\$319
	Anti-Bax	Μ	VVB IH(P)	Rb	100 µL	AB2915	\$255
	Anti-Bax, a.a. 43-61 hBax	Н		Rb	100 µL	AB2930	\$205
	Anti-Bax, N-terminus, aa3-16 hBax protein, clone 2D2	Н	VVB IF IH(P)	M IgG1	100 µg	MAB4601	\$290
	Anti-Bax, azide free, clone 2D2	Н	WB IP ELISA IF IH(I	-	100 µg	MAB4601Z	\$290
	APOPTOPAK, (Anti-Bcl2, Bak, Bax) Miniature Set		WB IP IC IH		1 kit	17-178	\$275
	siRNA plasmid, pKD-Bax-v2	Н	RNAi		5 µg	62-042	\$349
	siRNA plasmid, pKD-Bax-v3	H M R	RNAi		5 µg	62-132	\$349
	Bax cDNA (wt) in pUSEamp		TFX		5 µg	21-159	\$599
					10		

Bcl2	Description	Species	Applications	Host	Quantity	Cat. No.	Price
DCIZ	Anti-Bcl-2			Rb	50 µL	AB1720	\$210
	Anti-Bcl-2, a.a. 68-86 of mBCL2	M R		Rb	100 µL	AB1722	\$205
	Anti-A1	Μ	WB IP	Rb	100 µL	AB3155	\$205
	Anti-Bcl2, clone 100	H M Ca	WB FC IH	M lgG	100 µg	05-729	\$319
	Anti-Bcl2, clone AW604	Н	WB IP	M IgG <sub>1a</sub>	200 µg	05-826	\$319
	Anti-Bcl-2, clone 4D7	Н	IP DB	M IgG1	100 µg	MAB2900	\$280
	Anti-phospho-Bcl2 (Ser70), clone CT7	Н	WB IP	Rb IgG	100 µL	05-843	\$339
	IHC Select® Anti-Bcl-2, prediluted, clone 124	Н	IH(P)	M IgG <sub>1x</sub>	6 mL	IHC2033-6	\$210
	IHC Select Anti-Bcl-2, prediluted, clone 124	Н	IH(P)	M IgG <sub>1x</sub>	6 mL	IHCR2033-6	\$210
	APOPTOPAK, (Anti-Bcl2, Bak, Bax) Miniature Set		WB IP IC IH	0 11	1 kit	17-178	\$275
	Bcl-2, Sandwich ELISA	Н	ELISA		1 kit	APT230	\$525
	Bcl2 SMARTpool siRNA reagent	Н	RNAi		5 nmol	M-003307	\$375
	Bcl2 siRNA/siAb™ Assay Kit	Н	WB RNAi		1 kit	60-004	\$649
	Bcl2 cDNA (wt) in pUSE amp		TFX		50 µg	21-158	\$599
Bcl3					10		
DCIO	Anti-Bcl3	Н	WB IP IC	Rb IgG	200 µg	06-415	\$329
	Anti-Bcl-3, clone HAM150-3.5	НМ	WB ELISA	AHmst IgG <sub>11</sub>	100 µL	MAB2350	\$240
	siRNA plasmid, pKD-Bcl3-v1	Н	RNAi	3-11	5 µg	62-133	\$349
	siRNA plasmid, pKD-Bcl3-v4		RNAi		5 µg	62-233	\$349
	en en le construction de la constru				10		
Bcl6	Anti-Bcl-6, clone BL6.02 (PG-B6p)	H R Sh Rb B Po	IH(P)	M lgG <sub>1k</sub>	250 µL	MAB4618	\$210
D 110				MilgO <sub>K</sub>	200 pt	1110-010	ΨΖΙΟ
Bcl10	Anti-Bcl-10, N-terminus	Н	WB	Rb	100 µg	AB16506	\$195
	Anti-Bel10, clone 151	Н	WB FC IH	CD .	100 µL	05-732	\$319
	Anii-bci TO, cione 151	П	VVD FC IN		100 pt	03-732	Φ21A
BclW				DI	50	401700	¢010
	Anti-Bcl-W, a.a. 16-29	H M R B Mk	WB IH(P) Web*	Rb	50 µg	AB1723	\$210
	Anti-Bcl-W, clone 16H12	НМ	WB IP FC	R IgG <sub>2a</sub>	100 hâ	MAB17002	\$350
BclXL/	/XS						
New	Anti-phospho-Bcl-XL (SerS62)	Н	WB	Rb	100 µL	AB3573	\$315
	Anti-Bcl-XL, Phospho-specific	Н	WB	Rb	100 µL	AB3116	\$290
	Anti-Bcl-XL, N-terminus, clone 7D9	Μ	IF FC	M lgG <sub>2a</sub>	50 µg	MAB4619	\$210
	Anti-Bcl-XL, clone BXLO3	Н	IF	M IgG1	50 µg	MAB4620	\$210
	Anti-Bcl-X, clone 2H12	H M R Po	WB IP IF FC IH	$M \log G_{2a}$	50 µg	MAB4625	\$210
	Anti-Bcl-XL and BclXs, clone 7B2.5	Н	WB IP FC IH	M lgG <sub>3</sub>	50 µg	MAB3121	\$210
	Anti-Bcl-XL, clone 7B2.5	Н	FC IH	Μ	50 µg	MAB3121B	\$210
	Anti-Bcl-XL, clone 7B2.5	Н	FC IH	Μ	50 µg	MAB3121F	\$235
	Anti-Bcl-XL, clone 7B2.5	Н	FC IH	Μ	50 µg	MAB3121P	\$250
BID							
	Anti-BID	H M R	WB IP IH	Rb	100 µg	AB1730	\$330
	Anti-BID	H M R Rb	WB IP IH	Rb	100 µL	AB1735	\$205
New	Anti-Mouse BID Cleavage site specific	Μ	WB	Rb	100 µL	AB10002	\$315
Bim							
New	Anti-Bim, internal epitope, pan-Bim isoforms	H M R	WB	Rb	100 µg	AB17003	\$220
	Anti-phospho-Bim EL (Ser65)	H M R	WB IP	Rb IgG	200 µg	36-004	\$319
	Anti-phospho-Bim EL (Ser55)	Н	WB	Rb IgG	200 µg	36-005	\$319
	Anti-Bim, clone 14A8	НМ	WB IP FC IH	R IgG <sub>2a</sub>	250 µg	MAB17001-250UG	\$750
	Anti-Bim, clone 14A8	НМ	WB IP FC IH	R IgG <sub>2a</sub>	50 µg	MAB17001-50UG	
	Anti-BimL, clone 5E.5	НМ	WB IP ELISA FC IH	R lgG <sub>2b</sub>	50 µg	MAB3129	\$210
	Anti-BimL, clone 5E.5	НМ	WB IP ELISA FC IH	R IgG <sub>2a</sub>	250 µg	MAB3129-250UG	
	Bim-GST		WB KA		100 µg	12-483	\$275
	Bim-GST (3A)		WB KA		100 µg	12-484	\$275
New	Anti-phospho-Bim EL (Ser69)	HR	WB	Rb	100 µL	AB3579	\$315
Bin 1							
	Anti-Bin 1, clone 99D	HMR	WB IP IH	Μ lgG <sub>1κ</sub>	200 µg	05-449	\$319

<b>Description</b> lys	Species	Applications	Host	Quantity	Cat. No.	Price
Anti-Blys/TALL-1/BAFF/THANK	Н	WB	Rb IgG	100 ha	07-167	\$319
siRNA plasmid, pKD-Blys-v2	Н	RNAi		5 µg	62-129	\$349
siRNA plasmid, pKD-Blys-v3	Н	RNAi		5 µg	62-130	\$349
nip3L						
Anti-Bnip3L, internal	НМ	WB	Rb	100 hð	AB16507	\$220
AD						
Anti-CAD, C-terminus	Μ	WB	Rb	100 µg	AB16925	\$210
Calpain						
Anti-Calpain, small subunit of µ- or m-Calpains	HRB	WB ELISA IC IH	M lgG1	100 µL	MAB3083	\$190
(Calpain I or II), clone P1	THE B		111901	100 pt		ψ170
Anti-Calpain I, large subunit (Calpain I), clone P-6	Н	WB ELISA IC IH	M lgG1	100 µL	MAB3082	\$190
Anti-Calpain I, large subunit, clone P9	H M R Ht Ca	WB IP ELISA IC	M IgG1	100 µL	MAB3104	\$190
Anti-Calpain II, large (catalytic) subunit	Н	WB ELISA IC IH	Rb	100 µL	AB1625	\$170
Anti-Calpain II, large subunit	R	WB IP ELISA IH	Rb	100 µg	AB81013	\$300
Anti-Calpain LP85 and LP82, large subunit	R	WB IP ELISA IH	Rb	100 µg	AB81011	\$265
Anti-Calpain-94, large subunit, domain IV	Н	WB IP ELISA IH	Rb	100 µg	AB81018	\$300
Anti-Calpain-94, large subunit, domain II	Н	WB IP ELISA IH	Rb	100 µg	AB81019	\$300
Anir-Calpanir94, large subunit, domain in	11	VVD IF LUJA II I	ND	roo µg	ADOIUI9	φ300
aspase						
an-Caspase		4.CT		0.5	A DT 400	¢ 1 C C
CaspaTag™ Pan-Caspase <i>In Situ</i> Assay Kit, Fluorescein	Ma	ACT		25 assays	APT420	\$155
CaspaTag Pan-Caspase In Situ Assay Kit, Fluorescein	Ma	ACT		100 assays	APT400	\$435
CaspaTag Pan-Caspase In Situ Assay Kit, Sulforhodamine	Ma	ACT		25 assays	APT520	\$185
CaspaTag Pan-Caspase In Situ Assay Kit, Sulforhodamine	Ma	ACT		100 assays	APT500	\$470
CaspSCREEN, non-adherent cells only	Ma	ACT		25 assays	APT105	\$395
Caspase Family Inhibitor Set	Ma	ACT		1 kit	APT135	\$265
Caspase 1						
Anti-Caspase 1	H M R	WB IP IC	Rb IgG	200 µg	06-503	\$329
Anti-Caspase 1	HMR	WB IP IF	Rb	50 µg	AB1871	\$220
Caspase 1, recombinant human active		ACT		25 units	CC126	\$170
siRNA plasmid, pKD-Caspase 1-v3	Н	RNAi		5 µg	62-131	\$349
siRNA plasmid, pKD-Caspase 1-v2	Н	RNAi		5 µg	62-159	\$349
Caspase 1 Fluorometric Assay Kit, YVAD	Ma	ACT		100 assays	APT160	\$500
Caspase 1 Colorimetric Assay Kit, YVAD	Ma	ACT		100 assays	APT161	\$475
aspase 2						
Anti-Caspase 2, clone 10C6	H M Mk Ca	WB IP ELISA FC	R lgG <sub>2a</sub>	100 µg	MAB3501	\$290
Anti-Caspase 2, clone 11B4	H M Mk Ca	WB IP ELISA FC	R IgG <sub>2a</sub>	100 µg	MAB3507	\$290
Anti-Caspase 2, clone 4-1-1	Н	WB	M IgG	200 µL	05-711	\$299
Caspase 2 Fluorometric Assay Kit, VDVAD	Ma	ACT		100 assays	APT162	\$500
Caspase 2 Colorimetric Assay Kit, VDVAD	Ma	ACT		100 assays	APT163	\$475
Caspase 2, recombinant human active	1110	ACT		25 units	CC127	\$185
aspase 3 Anti-Caspase 3	H M R	WB	Rb IgG	200 µg	06-735	\$319
Anti-Caspase 3 Anti-Caspase 3, a.a. 104-117, proform & Irg subunit	HM	WB	Rb igG	200 µg 100 µL	AB3640	\$255
				1		
Anti-Caspase 3, large subunit & proform	HMR		Rb	50 µg	AB1899	\$220
Anti-Caspase 3, active (cleaved) form	HMR	WB IH IH(P) Web*	Rb	50 µg	AB3623	\$240
Anti-Caspase 3, clone 4-1-18	Н	WB	M lgG <sub>2a</sub>	100 µg	05-654	\$329
Anti-Caspase 3, large subunit & proform, clone 3CSP03	Н	VVB IP IH(P)	M lgG <sub>2a</sub>	50 µg	MAB4603	\$210
Anti-Caspase 3, large subunit & proform, clone 4-1-18	Н	WB IH(P)	M lgG <sub>2ακ</sub>	100 hð	MAB4703	\$220
Caspase 3, active		CA		20 µg	14-264	\$329
Human Caspase 3, recombinant human active		ACT		25 units	CC119	\$115
siRNA plasmid, pKD-Caspase 3-v2		RNAi		5 µg	62-245	\$349
siRNA plasmid, pKD-Caspase 3-v3		RNAi		5 µg	62-246	\$349
CleavaLite Caspase 3 Activity Assay Kit	Ma	ACT		96 assays	APT300	\$435
CleavaLite Caspase 3 Activity Assay Kit	Ma	ACT		384 assays	APT301	\$550

<b>Descript</b> CaspaTag	<b>ion</b> g Caspase 3,7 <i>In Situ</i> Assay Kit, Fluorescein	<b>Species</b> Ma	Applications ACT	Host	<b>Quantity</b> 25 assays	<b>Cat. No.</b> APT423	<b>Price</b> \$155
CaspaTag	g Caspase 3,7 In Situ Assay Kit, Fluorescein	Ma	ACT		100 assays	APT403	\$435
CaspaTag	g Caspase 3,7 In Situ Assay Kit, Sulforhodamine	Ma	ACT		25 assays	APT523	\$185
CaspaTag	g Caspase 3,7 In Situ Assay Kit, Sulforhodamine	Ma	ACT		100 assays	APT503	\$490
Caspase	3 Colorimetric Activity Assay Kit, DEVD	Ma	ACT		25 assays	APT131	\$205
Caspase	3 Colorimetric Activity Assay Kit, DEVD	Ma	ACT		100 assays	APT165	\$390
Caspase	3 Activity Detection Kit		СА		1 kit	17-198	\$349
Caspase	3/7 Assay Kit (Ac-DEVD-AMC Substrate)		CA		1 kit	17-367	\$420
Caspase 5			ACT		100	A DT 1 4 4	¢EOO
	5 Fluorometric Activity Assay Kit, WEHD	Ma	ACT		100 assays	APT166	\$500
Caspase	5 Colorimetric Activity Assay Kit, WEHD	Ma	ACT		100 assays	APT167	\$475
Caspase 6							
Anti-Casp	base 6	Н	WB IP	Rb IgG	200 µg	06-691	\$289
Anti-Casp	pase 6, proform & sm. subunit	H M R Ht Sh Rb B Po Mk Ca	WB	Rb	50 µg	AB1851	\$210
Caspase	6 Fluorometric Activity Assay Kit, VEID	Ma	ACT		100 assays	APT168	\$500
Caspase	6 Colorimetric Activity Assay Kit, VEID	Ma	ACT		100 assays	APT169	\$475
Caspase 7							
	pase 7, proform & lg. subunit	Н	WB	Rb	100 µL	AB1999	\$220
	pase 7, active (cleaved) form	HR	WB	Rb	50 µg	AB1999 AB3627	\$220
	pase 7, proform & lg. subunit	Н	WB	Ch	100 µL	AB3027	\$240
	pase 7, clone 7-1-11	H	WB IP	M IgG	100 µg	05-578	\$319
	pase 7, clone 10-1-62	Н	WB	Migo	100 µg	MAB4707	\$210
	7, recombinant human active	H	Act	111	25 units	CC125	\$170
	7, active	11	CA		25 units	14-398	\$289
	asmid, pKD-Caspase 7-v1	Н	RNAi		23 υniis 5 μg	62-004	\$349
	asmid, pKD-Caspase 7-v3	Н	RNAi		20 µg	62-004	\$349
	g Caspase 3,7 In Situ Assay Kit, Fluorescein	Ma	ACT		25 assays	APT423	\$155
	g Caspase 3,7 In Situ Assay Kit, Fluorescein	Ma	ACT		100 assays	APT403	\$435
	g Caspase 3,7 In Situ Assay Kit, Sulforhodamine	Ma	ACT		25 assays	APT523	\$185
	g Caspase 3,7 In Situ Assay Kit, Sulforhodamine	Ma	ACT		100 assays	APT503	\$490
	3/7 Assay Kit (Ac-DEVD-AMC Substrate)	1110	CA		1 kit	17-367	\$420
Caspase 8							
Anti-Casp		Н	WB	Rb IgG	200 µg	06-775	\$319
Anti-Casp		H M R B Mk	WB	Rb	50 µg	AB1879	\$220
	base 8, a.a. 410-424	Н	WB	Rb	100 µL	AB3641	\$255
	pase 8, clone 1-1-37	Н	WB IP	M lgG	100 hð	05-573	\$309
	pase 8, clone C51S	Н	WB IP	M IgG <sub>2a</sub>	50 µg	MAB3508	\$200
	pase 8, clone 8CSP01	Н	WB	Μ lgG <sub>1κ</sub>	50 µg	MAB4608	\$200
	pase 8, clone 1-1-37	Н	WB	M lg	100 µg	MAB4708	\$215
	8, recombinant human active	Н	ACT		25 units	CC123	\$170
	asmid, pKD-Caspase 8-v1	Н	RNAi		5 µg	62-145	\$349
	asmid, pKD-Caspase 8-v3		RNAi		5 µg	62-190	\$349
	g Caspase 8 <i>In Situ</i> Assay Kit, Fluorescein		ACT		25 assays	APT428	\$155
	g Caspase 8 <i>In Situ</i> Assay Kit, Fluorescein		ACT		100 assays	APT408	\$440
Caspaso					1)5	APT129	\$205
	8 Colorimetric Activity Assay Kit, IETD	Ma	ACT		25 assays		
Caspase	8 Fluorometric Activity Assay Kit, IETD	Ma	ACT		100 assays	APT170	\$500
Caspase							
Caspase	8 Fluorometric Activity Assay Kit, IETD	Ma	ACT		100 assays	APT170	\$500
Caspase Caspase	8 Fluorometric Activity Assay Kit, IETD	Ma	ACT	Rb	100 assays	APT170	\$500
Caspase Caspase Caspase 9 Anti-Casp	8 Fluorometric Activity Assay Kit, IETD 8 Colorimetric Activity Assay Kit, IETD	Ma Ma	ACT ACT	Rb Rb	100 assays 100 assays	APT170 APT171	\$500 \$390
Caspase Caspase Caspase 9 Anti-Casp Anti-Casp	8 Fluorometric Activity Assay Kit, IETD 8 Colorimetric Activity Assay Kit, IETD pase 9, proform & Ig. subunit	Ma Ma H M R	ACT ACT WB		100 assays 100 assays 100 µg	APT170 APT171 AB16969	\$500 \$390 \$220
Caspase Caspase 9 Anti-Casp Anti-Casp Anti-Casp	8 Fluorometric Activity Assay Kit, IETD 8 Colorimetric Activity Assay Kit, IETD pase 9, proform & lg. subunit pase 9, proform & sm. subunit	Ma Ma H M R H M	ACT ACT WB WB	Rb	100 assays 100 assays 100 µg 100 µg	APT170 APT171 AB16969 AB16970	\$500 \$390 \$220 \$220
Caspase Caspase 9 Anti-Casp Anti-Casp Anti-Casp More Anti-Casp	8 Fluorometric Activity Assay Kit, IETD 8 Colorimetric Activity Assay Kit, IETD base 9, proform & lg. subunit base 9, proform & sm. subunit base 9, active (cleaved) form	Ma Ma H M R H M H	ACT ACT WB WB WB IP IC	Rb Rb	100 assays 100 assays 100 µg 100 µg 50 µg	APT170 APT171 AB16969 AB16970 AB3629	\$500 \$390 \$220 \$220 \$240
Caspase Caspase 9 Anti-Casp Anti-Casp Anti-Casp Anti-Casp Anti-Casp Anti-Casp	8 Fluorometric Activity Assay Kit, IETD 8 Colorimetric Activity Assay Kit, IETD base 9, proform & Ig. subunit base 9, proform & sm. subunit base 9, active (cleaved) form base 9 cleavage site specific (315/316)	Ma Ma H M R H M H H	ACT ACT WB WB WB IP IC WB IC	Rb Rb Rb	100 assays 100 assays 100 µg 100 µg 50 µg 100 µL	APT170 APT171 AB16969 AB16970 AB3629 AB3577	\$500 \$390 \$220 \$220 \$240 \$315

<b>Description</b> Caspase 9, recombinant human active	Species H	Applications ACT	<b>Host</b> E. Coli	Quantity 25 units	<b>Cat. No.</b> CC120	<b>Price</b> \$155
CaspaTag Caspase 9 In Situ Assay Kit, Fluorescein		ACT		25 assays	APT429	\$155
CaspaTag Caspase 9 In Situ Assay Kit, Fluorescein		ACT		100 assays	APT409	\$440
Caspase 9 Fluorometric Activity Assay Kit, LEHD	Ma	ACT		100 assays	APT172	\$500
Caspase 9 Colorimetric Activity Assay Kit, LEHD	Ma	ACT		25 assays	APT139	\$195
Caspase 9 Colorimetric Activity Assay Kit, LEHD	Ma	ACT		100 assays	APT173	\$390
				100 4004/0	,	<i>4070</i>
Caspase 10	11.0		DI	100	451010	¢005
Anti-Caspase 10	H Ca	WB IP IH	Rb	100 µL	AB1010	\$205
Anti-Caspase 10, C-terminus, proform & sm. subunit	Н	WB	Rb	100 µg	AB16960	\$220
Caspase 10, recombinant human active	H	ACT		25 units	CC128	\$185
Caspase 10 Fluorometric Assay Kit, AEVD	Ma	ACT		25 assays	APT148	\$210
Caspase 10 Fluorometric Assay Kit, AEVD	Ma	ACT		100 assays	APT174	\$500
Caspase 10 Colorimetric Assay Kit, AEVD	Ma	ACT		100 assays	APT176	\$475
Caspase 12						
Anti-Caspase 12, prodomain, aa100-116 mcaspase 12.	H M R	WB	Rb	100 hð	AB3612	\$220
Anti-Caspase 12, N-terminus	HMR	VVB	Rb	100 hB	AB3613	\$220
Caspase 13						
Anti-Caspase 13	HMR	WB	Rb	100 µg	AB16531	\$220
				10		
Caspase 14				100	05/07	¢010
Anti-Caspase 14, clone 8-1-71	Н	WB	M lgG <sub>1κ</sub>	100 hð	05-687	\$319
Cathepsin						
Anti-Cathepsin B	M R	WB IH	Rb IgG	400 µg	06-480	\$319
Anti-Cathepsin B	Н	WB	Rb	100 hð	AB4064	\$240
Anti-Cathepsin D	НМ	WB IC	Rb IgG	200 µg	06-467	\$319
Anti-Cathepsin D, clone C5	Н	IH(P)	$M \ IgG_{2b}$	500 µL	MAB422	\$300
IHC Select Anti-Cathepsin D, prediluted, clone C5	Н	IH(P)	$M \ IgG_{2b}$	6 mL	IHC2105-6	\$235
IHC Select Anti-Cathepsin D, prediluted, clone C5	Н	IH(P)	$M \ IgG_{2b}$	6 mL	IHCR2105-6	\$235
Anti-Cathepsin G, clone AHN-11	Н	RIA IC	M IgG <sub>2a</sub>	100 µL	MAB1054	\$190
Anti-Cathepsin K, clone 182-12G5	Н	WB Web*	$M \; \text{IgG}_{1\kappa}$	100 hð	MAB3324	\$265
CIDE						
Anti-CIDE-A, C-terminus	Н	WB	Rb	100 µg	AB16508	\$195
Anti-CIDE-A, C-terminus	Μ	WB	Rb	100 µg	AB16922	\$210
Anti-CIDE-B, C-terminus	Μ	WB	Rb	100 µg	AB16923	\$195
				10		
Cyclophilin A				000 1	07.010	¢000
Anti-Cyclophilin A	H M R	WB	Rb IgG	200 µL	07-313	\$299
Cyclophilin A (human) SMARTpool siRNA reagent	H	RNAi		5 nmol	M-004979	\$375
Cyclophilin A (mouse) SMARTpool siRNA reagent	M	RNAi		5 nmol	M-040767	\$375
siRNA plasmid, pKD-Cyclophilin A-v1	Н	RNAi		5 µg	62-079	\$349
siRNA plasmid, pKD-Cyclophilin A-v2	Н	RNAi		5 µg	62-080	\$349
Cyclophilin A siRNA/siAb Starter Kit (human)	H	WB RNAi		1 kit	61-003	\$649
Cyclophilin A siRNA/siAb Starter Kit (mouse)	Μ	WB RNAi		1 kit	61-004	\$649
Cystatin						
Anti-Cystatin A	НМ	WB	Rb	200 µg	AB4065	\$240
Anti-Cystatin C	HMR	VVB IH	Rb IgG	250 µg	06-458	\$319
siRNA plasmid, pKD-Cystatin C-v3	Н	RNAi		20 µg	62-010	\$349
Cytochrome C						
Anti-Cytochrome C	Н	ELISA	Ch	100 µg	AB3425	\$325
Anti-Cytochrome C	H R Rb Ca	WB IC	Sh	100 µg	AB3547	\$235
Anti-Cytochrome C, clone C-7	H Eq	WB IH	M IgG	200 µL	05-479	\$329
Anti-Cytochrome C, clone 7H8.2C12	H M R	WB	M IgG <sub>2b</sub>	100 µg	MAB1800	\$330
Anti-Cytochrome C, clone CTC04	H B Eq Ca	FC	M IgG <sub>1</sub>	, со ру 50 µg	MAB4612	\$210
Cytochrome C ELISA Kit	H	ELISA		1 plate	APT200	\$525
Cytochrome C 6H2.B4 Monoclonal	HMR	IP IC EM	Μ	100 µL	MAB3914	\$315
Cytochrome C 7H8.2C12 Monoclonal	HMR	WB	M	100 pL	MAB3916	\$315
Systements Strift. Zerz monocional				. 00 pc		4010

Description	Species	Applications	Host	Quantity	Cat. No.	Price
DAPK				000	0/ 050	¢000
Anti-DAPK	H	WB	Rb IgG	200 µg	06-859	\$329
Anti-DAP Kinase 2	H M R	WB	Rb	100 hð	AB3606	\$220
Daxx						
Anti-Daxx	H M R Ht	WB IP IC	Rb IgG	200 µL	07-471	\$299
Anti-Daxx, C-terminus	Н	WB	Rb	100 hð	AB16959	\$210
Death Receptor			D	100	401/050	¢000
Anti-DR3, C-terminus extracellular domain	H M H M	WB WB	Rb Rb	100 µg	AB16952 AB16953	\$220 \$205
Anti-DR3, N-terminus Anti-DR3, clone B65	Н	IP IF FC	M IgG <sub>2a</sub>	100 µg 50 µg	MAB4614	\$203 \$210
Anti-DR4, C-terminus	Н	WB	Rb	100 µg	AB16955	\$220
Anti-DR4/TRAIL-R1	H M R	WB	Rb IgG	200 µg	06-744	\$299
Anti-DR5, C-terminus	Н	WB	Rb	100 µg	AB16942	\$220
				10		
			D	100	401/500	¢000
Anti-DcR1, extracellular	Н	WB	Rb	100 µg	AB16509	\$220
Anti-DcR2, intracellular domain	H	WB	Rb Rb	100 µg	AB16943	\$220
Anti-DcR3, N-terminus	H M R	WB	ND	100 hð	AB16510	\$195
DEDAF						
Anti-DEDAF	H M R	WB	Rb	100 µg	AB3637	\$220
DFF						
Anti-ICAD/DFF, N-terminus (a.a. 2–21 of mouse K	CAD) M	WB	Rb	100 µg	AB16965	\$220
Anti-DFF40	H M R	VVB	Rb	100 µg	AB16926	\$210
Anti-DFF45/35, N-terminus	Н	VVB IP	Rb	100 hð	AB16961	\$205
Anti-DFF45, C-terminus	Н	VVB IP	Rb	100 hð	AB16962	\$205
Single Stranded DNA						
Anti-DNA, single stranded specific, clone F7-	26 A	FC IC IH IH(P)	M IgM	50 µg	MAB3299	\$210
Anti-DNA, single stranded	A	ELISA FC IC IH(P)	$M \lg G_{2a}$	100 µg	MAB3868	\$265
Anti-DNA, single stranded, clone 16-19	А	ELISA	$M  lgG_{2a}$	500 µL	MAB3034	\$190
ssDNA Apoptosis ELISA Kit	Ma	ELISA	0 20	1 plate	APT225	\$325
DOK						
DOK Anti-DOK, C-terminus	Н	WB	Rb	100 µg	AB16948	\$205
Anibok, Cleminus	11	VVD	ND	100 pg	AD10940	Ψ200
DRAK						
Anti-DRAK1, N-terminus	H M R	WB	Rb	100 hð	AB16514	\$205
Anti-DRAK2, C-terminus	Н	WB	Rb	100 hð	AB16515	\$205
Endonuclease G						
Anti-Endonuclease G	HMR	WB	Rb	100 ha	AB3639	\$195
FADD						
Anti-FADD	Н	WB IP	Rb IgG	200 µg	06-711	\$289
Anti-FADD	H M R Mk	VVB IP	Rb	200 рд 50 µg	AB3102	\$210
Anti-FADD, clone 1F7	НM	WB	M lgG1	100 µg	05-486	\$319
			0	10		
FAS and FAS Ligand						
FAS Anti-Fas (human, activating), clone CH11	Н	WB FC IC	AA 1~AA	50 µg	05-201	\$339
Anti-Fas (human, neutralizing), clone ZB4	H	NEUT	M lgM M lgG1	100 µg	05-201	\$329
Anti-Fas, clone 7C10	MB	WB IP	$R  lgG_{2a}$	200 µg	05-351	\$299
Anti-Fas, clone B-G27	H R	FC IH	M $IgG_{2a}$	100 assays	CBL527B	\$240
Anti-Fas, clone B-G27	HR	FC IH	$M \log G_{2a}$	100 assays	CBL527F	\$240
Anti-Fas, clone B-D29	Н	IP FC	M IgG <sub>1</sub>	100 µg	CBL537	\$130
Anti-Fas, clone SM1/1	Н	C IND	$M \log G_{2a}$	100 µg	MAB3061	\$210
Anti-Fas, clone SM1/23	Н	FC INHIB	$M \ lgG_{2a}$	100 µg	MAB3065	\$220
Anti-Fas, clone 95C02	Н	IF	M lgG	150 µg	MAB4621	\$200
Anti-Fas, clone 95C03	Н	IH	$M \ lgG_1$	250 µL	MAB4622	\$210
Soluble Fas ELISA Kit		ELISA		1 plate	APT 1 <i>77</i>	\$575

<b>Description</b> S Ligand	Species	Applications	Host	Quantity	Cat. No.	Pric
Anti-Fas Ligand	H M R	WB ELISA	Rb	100 µL	AB1665	\$39
Anti-Fas Ligand	HMR	WB ELISA	Rb	50 µg	AB16982	\$29
Anti-Fas Ligand, clone ALF1.2	Н	ELISA FC	Μ	100 µL	MAB3912	\$31
Anti-CD95L, clone MFL3	M	IF FC	AHt IgG	500 µg	CBL1369	\$41
Anti-CD95L, clone MFL3	M	IF FC	AHt IgG	500 µg	CBL1369B	\$43
Anti-CD95L, clone MFL3	M	FC	AHt IgG	100 µg	CBL1369P	\$29
Anti-CD178, clone B-R17	Н	FC INHIB NEUT	M IgG <sub>1</sub>	100 µg	CBL587	\$13
Anti-Fas Ligand, clone FSL01	H	WB	M IgM	250 µL	MAB4623	\$21
Fas Ligand, membrane bound		CC	i i i givi	500 ng	01-210	\$30
ASH		66		500 lig	01210	ψυυ
Asti-FLASH	Н	WB	Rb	100 hð	AB3605	\$20
P						
Anti-FLIP, NT	НМ	WB	Rb IgG	200 µg	06-697	\$31
Anti-FLIP, CT	H M R	WB	Rb IgG	200 µg	06-864	\$31
Anti-FLIP, N-terminus	HMR	WB	Rb	100 µg	AB16516	\$22
Anti-FLIP, C-terminus	Н	WB	Rb	100 µg	AB16963	\$22
Anti-FLIP $\gamma/\delta$ , C-terminus	НМ	WB	Rb	100 µg	AB16998	\$20
Anti-mFLIP, C-terminus	M	WB IP	Rb	100 µg	AB16964	\$19
actin				· · · · F3		<b>.</b>
Anti-Fractin, C-terminus	HMR	WB IC IH(P)	Rb	100 µL	AB3150	\$26
utathione						
Glutathione Detection Kit	Ma	ACT		100 assays	APT250	\$32
anzyme B						
Anti-Granzyme B, clone GrB-7	Н	VVB IH(P)	$M \ lgG_{2a}$	250 µL	MAB3070	\$19
Anti-Granzyme B, clone 2C5/F5	HR	WB ELISA FC IC	$\rm M~lgG_{2a}$	100 hB	MAB3170	\$29
5K-3						
Anti-GSK-3	H M R Mk Ch	WB IHC	Rb	500 µL	AB9258	\$30
Anti-phospho-GSK-3β (Ser9)	HR	WB	Rb IgG	100 µL	07-835	\$30
Anti-GSK-3β	HMRB	WB IP ICC	Rb	50 µg	AB8687	\$18
Anti-GSK-3α	Н	ELISA Web*	Chk	100 µg	AB3441	\$33
Anti-GSK-3β	Н	ELISA Web*	Chk	100 µg	AB3443	\$30
Anti-GSK-3α	HR	WB IPK	Rb IgG	200 µg	07-389	\$30
Anti-phospho-GSK3β (Ser9), clone 2D3	HMR	WB EMSA	M lgG <sub>1ĸ</sub>	100 µg	05-643	\$33
Anti-phospho-GSK3α (Ser21), clone BK202	H R	WB	M IgG <sub>1ĸ</sub>	100 µg	05-676	\$3
Anti-GSK3 $\alpha$ , clone AW103	HMR	WB IPK	Rb IgG	100 µg	05-737	\$3
Anti-phospho-GSK3 $\alpha$ (Ser21)	Н	WB	Rb IgG	100 pg 100 µL	07-393	\$3
Anti-phospho-GSK3α (Ser21) Anti-phospho-GSK3α (Ser21)	HR	WB	Sh IgG	200 µg	07-393	\$3
	1 I K	VVD	Shirigo	200 µg	07-420	φJ
stone H2A.X Anti-phospho-H2A.X (Ser139)	WR	WB IP	Rb IgG	200 µg	07-164	\$3
			-	10		
Anti-phospho-Histone H2A.X (Ser139), clone JBW301	VVR	WB IF IC	M lgG1	200 µg	05-636	\$34
Anti-phospho-Histone H2A.X (Ser139), clone JBW301, biotin conjugate	WR	WB IC	M lgG1	100 hâ	16-193	\$3:
Anti-phospho-Histone H2A.X (Ser139), FITC conjugate	VVR	FC	M lgG1	100 µg	16-202A	\$3
H2A.X Phosphorylation Assay Kit		ELISA		1 kit	17-327	\$50
(Chemiluminescence Detection) H2A.X Phosphorylation Assay Kit (Flow Cytometry)		FC		1 kit	17-344	\$4
tone H2B						
Anti-phospho-Histone H2B (Ser14), clone MC603	Н	BD FC	Rb IgG	100 µg	05-751	\$32
Anti-phospho-Histone H2B (Ser14)	Н	WB ELISA IC IH	Rb IgG	200 µL	07-191	\$3
rA2/Omi						
Anti-HtrA2/Omi, clone 18-1-83	Н	WB	M lgG	200 µL	05-721	\$29
lew Anti-phospho-IκBα (Ser32/Ser36)	Н	WB	Rb IgG	100 µL	07-836	\$30

IKK	Description	Species	Applications	Host	Quantity	Cat. No.	Price
	Anti-IKK2	HMRB	WB	Ch	100 µg	AB3451	\$330
	Anti-phospho-IKK $lpha$ (Ser176/Ser180)	Н	WB	Rb IgG	100 µL	07-837	\$309
	Anti-IKKα, clone 14A231	H M Mk	WB	M lgG1	100 µg	05-536	\$329
	IKK $\alpha$ SMARTpool siRNA reagent	Н	RNAi	0.	5 nmol	M-003473	\$375
	siRNA plasmid, pKD-IKKα-v3	НМ	RNAi		5 µg	62-040	\$349
	siRNA plasmid, pKD-IKKα-v4	H M R	RNAi		5 µg	62-049	\$349
	Anti-ΙΚKβ	Н	WB	Rb	100 µg	AB16522	\$205
	Anti-IKKβ, clone 10AG2	H M R	WB	M lgG1	100 µg	05-535	\$329
	siRNA plasmid, pKD-IKKβ-v3	HR	RNAi	0,	5 µg	62-039	\$349
	siRNA plasmid, pKD-IKKβ-v2	Н	RNAi		5 µg	62-050	\$349
	IKK- $\beta$ SMARTpool siRNA reagent	Н	RNAi		5 nmol	M-003503	\$375
	Anti-ΙΚΚγ, N-terminus	Н	WB ELISA	Rb	100 µg	AB3458	\$220
	Anti-IKKy	НМ	WB	M lgG	100 hð	05-631	\$329
	siRNA plasmid, pKD-IKKy-v2	Н	RNAi	0	5 µg	62-053	\$349
	IKKy SMARTpool siRNA reagent	Н	RNAi		5 nmol	M-003767	\$375
	Anti-IKKε	Н	WB	Rb IgG	200 µg	07-580	\$319
	siRNA plasmid, pKD-IKKε-v2	Н	RNAi		5 µg	62-031	\$349
	siRNA plasmid, pKD-IKKε-v3	Н	RNAi		5 µg	62-051	\$349
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ILP-2			14/0	D	E0 1	AD 4 5 1 4	¢105
	Anti-ILP-2	H M R	WB	Rb	50 µL	AB4514	\$195
IAP (Ir	nhibitor of Apoptosis)						
	Anti-cIAP-1	HMR	WB	Rb	50 µg	AB3614	\$240
	Anti-HIAP-2 (cIAP-1)	H M R	WB	Rb IgG	200 µg	07-759	\$289
	Anti-cIAP-2	HMR	WB	Rb	50 µg	AB3615	\$240
	Anti-NAIP	Н	WB	Rb	50 µg	AB3617	\$240
	Anti-XIAP	Н	WB	Rb	50 µg	AB3616	\$240
	Anti-RIAP-3 (XIAP)	HR	WB	Rb IgG	200 µg	07-753	\$289
JNK							
	Anti-JNK/SAPK1	HMR	WB IP	Rb IgG	200 µg	06-748	\$309
	Anti-INK1	MR	IP KA	Sh	250 µg	AB4081	\$240
	Anti-phospho-JNK (Thr183/Tyr185, Thr221/Tyr223)	HMR	WB IP	Rb IgG	200 µg	07-175	\$319
	Anti-INK2	HMR	WB IP ELISA	Rb	50 µL	AB8910	\$189
	siRNA plasmid, pKD-JNK2α2/SAPK1a-v1	Н	RNAi		5 µg	62-097	\$349
	siRNA plasmid, pKD-JNK2α2/SAPK1a-v5	НM	RNAi		5 µg	62-098	\$349
	JNK2 SMARTpool siRNA reagent	Н	RNAi		5 nmol	M-003515	\$375
	JNK2/SAPK1a siRNA/siAb Assay Kit	Н	WB RNAi		1 kit	60-099	\$649
	Anti-JNK3/SAPK1b, clone C05T	HR	WB	Rb IgG	100 µL	05-893	\$319
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Livin							
	Anti-Livin	Н	WB	Rb	100 hâ	AB3632	\$280
Mcl-1							
	Anti-Mcl-1	Н	WB IP IH(P)	Rb	100 µL	AB2910	\$240
	Anti-Mcl-1, clone RC13	Н	WB IP IH(P)	M lgG1	100 µg	MAB4602	\$280
	Anti-Mcl-1, clone RC31	Н	WB IP	M IgG <sub>1</sub>	50 µg	MAB4624	\$185
				Ŭ ,	10		
MKK	Apti-AAKKZ /SKKA	Ц	WB	Phace	100	07-399	\$210
	Anti-MKK7/SKK4 Anti-phospho-MKK7/SKK4 (Thr275/Ser277)	H	VVB VVB	Rb IgG Rb IgG	100 µg 100 µL	36-010	\$319 \$319
	Anti-phospho-MKK7/SKK4 (Thr275)	Н	WB WB	Rb IgG	200 µg	36-010	\$319
			V V D	nu iyo	200 hð	30/013	Ψ017
mTOR							
	Anti-mTOR (mTAb1)	HMR	WB IPK KA	Rb	200 µL	07-231	\$299
	Anti-mTOR	HR	WB IP	Rb	100 hð	AB3882	\$235
p35							
	Anti-p35 (cdk5 regulator)	НВ	IP	M lgG	100 µg	05-380	\$329
				0	10		

	<b>Description</b> APK/SAPK	Species	Applications	Host	Quantity	Cat. No.	Price
New A	Anti-p38α	HMR	WB IP IPK ELISA	Rb IgG	100 µg	09-273	\$309
-	Anti-p38 MAPK, phospho-specific (Thr180/Tyr182)	HMR	WB IP IPK IF	Rb IgG	100 µL	09-272	\$319
-	GST-p38 $\alpha$ , recombinant human inactive			0	25 µg	SGT221	\$290
	o38α MAPK, recombinant human active				10 µg	SGT222	\$330
	Non-radioactive p38 Kinase Assay Kit				40 assays	SGT455	\$345
	Anti-p38/SAPK2, clone 2F11	HMR	WB IP	M lgG	200 µg	05-454	\$329
	p38α/SAPK2a Assay Kit, 50 Assays	TT / VI K	KA	migo	200 pg 1 kit	17-169	\$560
	538/SAPK2 SMARTpool siRNA reagent		RNAi		5 nmol	M-003512	\$375
		Н					
	38/SAPK2 siRNA/siAb™ Assay Kit		WB RNAi		1 kit	60-032	\$649
	Anti-p38γ/SAPK3	M R	WB	Rb IgG	200 µg	07-139	\$309
	Anti-p38γ/SAPK3	Н	WB	Rb IgG	200 µg	07-474	\$319
	Anti-p38γ/SAPK3	Н	WB	Rb IgG	200 µL	07-508	\$309
A	Anti-p388/SAPK4	HR	WB	Rb IgG	200 µg	07-603	\$309
<b>RP</b>							
	Anti-PARP Poly ADP-ribose Polymerase-1	H M R B	WB ELISA	Rb	100 µL	AB16661	\$330
	Anti-PARP Cleaved Form	Н	WB	Rb	50 µg	AB3620	\$240
	Anti-Poly ADP-ribose, clone 10H	A	WB IP IC	M IgG <sub>3a</sub>	50 pg	MAB3192	\$265
	Anti-PARP, clone A6.4.12	H M R Ht Xn Dr	WB IP ELISA IH IH(P)	M IgG <sub>1</sub>	2 mL	MAB3217	\$380
	Anti-PARP, clone C-2-10	H M R Ht Av	WB ELISA IC		50 µg	MAB3290	\$30
	Anti-VPARP, clone p193-4	Н	WB IH	M IgG <sub>1</sub>	100 µg	MAB4142	\$260
	Anti-phospho-PARP	НМ	WB IC	Rb	100 µL	AB3565	\$31
s DK	iRNA plasmid, pKD-PARP-v2	Н	RNAi		5 hð	62-156	\$34
	Anti-PDK 1	HR	WB IPK	Rb IgG	100 µL	07-707	\$30
	Anti-PDK 1	HR	WB IPK	Rb IgG	100 µL	07-707	\$30
	Anti-Pyruvate Dehydrogenase Kinase isoform 1	Н	WB	Rb	100 µg	AB4236	\$23
	Anti-Pyruvate Dehydrogenase Kinase isoform 2	Н	WB	Rb	100 µg	AB4238	\$23
	2DK1 Immunoprecipitation Kinase Assay Kit	11	IPK	ND	1 kit	17-279	\$45
	2DK1 Immunoprecipitation Kinase Assay Kit		IPK		1 kit	17-279	\$45
	,		KA				
	PDK1 Kinase Assay Kit				1 kit	17-280	\$45
	PDK1 Kinase Assay Kit		KA		1 kit	17-280	\$45
erforin							
A	Anti-Perforin, clone dG9	НМ	IP IF FC IH	$M \; \text{IgG}_{2b\kappa}$	50 µg	MAB4616	\$21
HAP							
A	Anti-PHAP, N-terminus	HMR	WB	Rb	50 µL	AB4515	\$19
	atidylserine						
	7	\ A /D		M I=C	200	05 710	\$ 2.2
	Anti-Phosphatidylserine, clone 1H6	WR NA/P	FC IH	M IgG	200 µg	05-719	\$33
P	Anti-Phosphatidylserine, clone 1H6, Alexa Fluor 488 conjugate	VVR	FC	M lgG	100 hB	16-256	\$33
3-Kind							
	Anti-PI 3-Kinase, p85	H M R Mk	WB IP	Rb IgG	125 µL	06-195	\$32
	Anti-PI 3-Kinase, p85, N-SH2 domain	H M R Mk	WB IP	Rb IgG	250 µg	06-496	\$32
	Anti-PI 3-Kinase, p85	H M R Mk	WB IP	Rb IgG	250 µg	06-490	\$31
	Anti-PI 3-Kinase, p85, N-SH3, clone AB6	H M	WB IP IC	-	230 µg 100 µg	05-212	\$33
		H M R	WB IP	M lgG <sub>1κ</sub>		05-212	\$33 \$32
	Anti-PI 3-Kinase, p85, N-SH2, clone UB93-3			M IgG	200 µL		
	Anti-PI 3-Kinase p $85\alpha$ , clone 8-2D-4D	HMR	IP IF FC	M lgG1	100 µg	MAB1143	\$26
	Pl 3-Kinase p85 $\alpha$ SMARTpool siRNA reagent	Н	RNAi		5 nmol	M-003020	\$37
	iRNA plasmid, pKD−Pl3 Kinase, p85-v3		RNAi		5 µg	62-222	\$34
	Anti-PI3 Kinase, p101	HR	WB	Rb IgG <sub>1</sub>	200 µg	07-281	\$30
	Anti-PI3 Kinase, p110α	H VVR	IP	Rb IgG	200 µg	07-658	\$30
A	Anti-P13 Kinase, p110β	HMRB	IP	Rb IgG	200 µg	06-568	\$32
A	Anti-PI3 Kinase, p1108, clone AW103	Н	WB	$M \; IgG_{1\kappa}$	200 µg	05-703	\$32

PKA	Description	Species	Applications	Host	Quantity	Cat. No.	Price
	Anti-PKA, NT	H M R Ht B Po	WB	Rb IgG	200 µg	06-903	\$319
	Anti-PKA, RII subunits	HR	WB IP IC	Gt IgG	500 µg	06-411	\$329
	Anti-phospho-PKA, RII (Ser96)	MR	WB	Rb IgG	100 µg	06-704	\$309
	Anti-phospho-PKA, Regulatory subunit IIβ (Ser114)	M	WB	Rb IgG	100 µL	07-869	\$309
	Anti-phospho-PKA catalytic subunits $\alpha/\beta$ (Thr197)	M	WB	Rb IgG	100 µL	07-867	\$309
	Anti-phospho-PKA Catalytic β subunit (Ser338)	M	WB	Rb IgG	100 µL	07-868	\$309
	PKA Assay Kit		KA	no igo	1 kit	17-134	\$229
	PKA Inhibitor Cocktail		KA		1 mL	20-114	\$94
	PKA/PKC Inhibitor Cocktail		KA		1 mL	20-129	\$94
	PKA/CaMK Inhibitor Cocktail		KA		1 mL	20-132	\$94
	siRNA plasmid, pKD-PKAa-v2	Н	RNAi		5 µg	62-069	\$349
	PKA siRNA/siAb Assay Kit	Н	WB RNAi		1 kit	60-115	\$649
	Anti-cAMP-Dependent Protein Kinase, Regulatory	H M Mk	WB IH	Rb	100 µg	AB1612	\$235
	Subunit I-B, internal				10		
	Anti-cAMP-Dependent Protein Kinase, Regulatory Subunit II-α	H M Mk	WB IP ELISA FC IH	Rb	100 hð	AB1613	\$235
	Anti-cAMP-Dependent Protein Kinase, Regulatory Subunit Il-β, internal	H M Mk	WB IH	Rb	100 hð	AB1614	\$235
D							
ΡΚϹ ΡΚϹα							
	Anti-PKC $\alpha$ , $\beta$ , $\gamma$	H M R Rb B	WB	Rb IgG	200 µg	06-870	\$309
	Anti-PKC $\alpha$ , clone M4	H M R Rb B	WB IP NEUT	M lgG1	100 µg	05-154	\$329
	Anti-Protein Kinase C $\alpha$ , clone 1F3.2	Н	WB ELISA	M lgM	100 µL	MAB3074	\$180
	Anti-phospho-PKC $lpha$ (Thr638)	НМ	WB	Rb IgG	100 µL	07-871	\$309
	Anti-phospho-PKC $lpha$ (Ser657)	H M R Rb B	WB	Rb IgG	200 µg	06-822	\$319
	PKC $\alpha$ SMARTpool siRNA reagent		RNAi		5 nmol	M-003523	\$375
	siRNA plasmid, pKD-PKCα-v4	Н	RNAi		5 µg	62-104	\$349
	siRNA plasmid, pKD·PKCα-ν6	НМ	RNAi		5 µg	62-105	\$349
ΡΚϹβ				-			****
	Anti-phospho-PKCβ I&II (Thr500)	Н	WB	Rb IgG	100 µL	07-870	\$309
	Anti-phospho-PKCβ I (Thr642)	Н	WB	Rb IgG	100 µL	07-872	\$309
	Anti-phospho-PKC $\beta$ II (Thr641)	Н	WB	Rb IgG	100 µL	07-873	\$309
ΡΚϹδ							
	Anti-PKC <b>o</b>	HMR	WB	Rb IgG	200 µg	06-990	\$309
	Anti-PKC $\delta$	НМ	WB IP IH	Sh	100 µg	AB1685	\$180
	Anti-phospho-PKCδ (Ser645)	Н	WB	Rb IgG	100 µL	07-874	\$309
	Anti-phospho-PKCδ (Ser664)	Н	WB	Rb IgG	100 µL	07-875	\$309
	siRNA plasmid, pKD-PKC <b>&amp;</b> -v3	Н	RNAi		5 µg	62-071	\$349
	siRNA plasmid, pKD-PKC <b>&amp;</b> -v6	HMR	RNAi		5 µg	62-072	\$349
ΡΚϹε							
I NCE	Anti-PKC <sub>e</sub>	H M R	WB IP IH	Rb IgG	200 µg	06-991	\$309
	( ) ( ) ( ) ( ) ( ) ( ) ( ) ( ) ( ) ( )	I I IVI K		-			
		HAAPPL	\A/R	Ph laC	200~	()6.201	4.3.70
	Anti-phospho-PKCe (Ser729)	h m r rb	WB PNIA;	Rb IgG	200 µg	06-821	\$329 \$340
	Anti-phospho-PKCe (Ser729) siRNA plasmid, pKD-PKCe-v1	Н	RNAi	Rb IgG	5 µg	62-121	\$349
	Anti-phospho-PKCe (Ser729) siRNA plasmid, pKD-PKCe-v1 siRNA plasmid, pKD-PKCe-v5		RNAi RNAi	Rb IgG	5 µg 5 µg	62-121 62-122	\$349 \$349
	Anti-phospho-PKCe (Ser729) siRNA plasmid, pKD-PKCe-v1 siRNA plasmid, pKD-PKCe-v5 PKCe SMARTpool siRNA reagent	Н	RNAi	Rb IgG	5 µg	62-121	\$349
РКСө	Anti-phospho-PKCe (Ser729) siRNA plasmid, pKD-PKCe-v1 siRNA plasmid, pKD-PKCe-v5 PKCe SMARTpool siRNA reagent	H	RNAi RNAi RNAi		5 μg 5 μg 5 nmol	62-121 62-122 M-004653	\$349 \$349 \$375
РКСө	Anti-phospho-PKCe (Ser729) siRNA plasmid, pKD-PKCe-v1 siRNA plasmid, pKD-PKCe-v5 PKCe SMARTpool siRNA reagent Anti-phospho-PKC <b>e</b> (Ser676)	H H H	RNAi RNAi RNAi WB	Rb IgG	5 μg 5 μg 5 nmol	62-121 62-122 M-004653 07-883	\$349 \$349 \$375 \$309
РКСө	Anti-phospho-PKCe (Ser729) siRNA plasmid, pKD-PKCe-v1 siRNA plasmid, pKD-PKCe-v5 PKCe SMARTpool siRNA reagent Anti-phospho-PKCO (Ser676) Anti-phospho-PKCO (Ser695)	H H H H	RNAi RNAi RNAi WB WB	Rb IgG Rb IgG	5 μg 5 μg 5 nmol 100 μL 100 μL	62-121 62-122 M-004653 07-883 07-884	\$349 \$349 \$375 \$309 \$309
РКСө	Anti-phospho-PKCe (Ser729) siRNA plasmid, pKD-PKCe-v1 siRNA plasmid, pKD-PKCe-v5 PKCe SMARTpool siRNA reagent Anti-phospho-PKC <b>e</b> (Ser676)	H H H	RNAi RNAi RNAi WB	Rb IgG	5 μg 5 μg 5 nmol	62-121 62-122 M-004653 07-883	\$349 \$349 \$375 \$309
	Anti-phospho-PKCe (Ser729) siRNA plasmid, pKD-PKCe-v1 siRNA plasmid, pKD-PKCe-v5 PKCe SMARTpool siRNA reagent Anti-phospho-PKC0 (Ser676) Anti-phospho-PKC0 (Ser695) Anti-phospho-PKC0 (Thr538)	H H H H	RNAi RNAi RNAi WB WB	Rb IgG Rb IgG	5 μg 5 μg 5 nmol 100 μL 100 μL	62-121 62-122 M-004653 07-883 07-884	\$349 \$349 \$375 \$309 \$309
ΡΚCθ ΡΚCη	Anti-phospho-PKCe (Ser729) siRNA plasmid, pKD-PKCe-v1 siRNA plasmid, pKD-PKCe-v5 PKCe SMARTpool siRNA reagent Anti-phospho-PKCO (Ser676) Anti-phospho-PKCO (Ser695) Anti-phospho-PKCO (Thr538)	H H H H	RNAi RNAi RNAi WB WB	Rb IgG Rb IgG Rb IgG	5 μg 5 μg 5 nmol 100 μL 100 μL 100 μL	62-121 62-122 M-004653 07-883 07-884 07-885	\$349 \$349 \$375 \$309 \$309 \$309
	Anti-phospho-PKCe (Ser729) siRNA plasmid, pKD-PKCe-v1 siRNA plasmid, pKD-PKCe-v5 PKCe SMARTpool siRNA reagent Anti-phospho-PKC0 (Ser676) Anti-phospho-PKC0 (Ser695) Anti-phospho-PKC0 (Thr538)	H H H H	RNAi RNAi RNAi WB WB WB	Rb IgG Rb IgG	5 μg 5 μg 5 nmol 100 μL 100 μL	62-121 62-122 M-004653 07-883 07-884	\$349 \$349 \$375 \$309 \$309

Description	Species	Applications	Host	Quantity	Cat. No.	Price
Anti-PKC <sup>c</sup>	H M R	WB	Rb IgG	200 µL	07-264	\$309
siRNA plasmid, pKD-PKCζ-v1	Н	RNAi		5 µg	62-073	\$349
siRNA plasmid, pKD-PKC5-v3	Н	RNAi		5 µg	62-074	\$349
PKCς SMARTpool siRNA reagent		RNAi		5 nmol	M-003526	\$375
C General Reagents						
PKC Assay Kit		KA		1 kit	17-139	\$229
PKC Lipid Activator		KA		1 ml	20-133	\$74
PKC/CaMK Inhibitor Cocktail		KA		1 ml	20-119	\$94
PKC Substrate Cocktail		KA		1 ml	20-131	\$135
22A						
Anti-Protein Phosphatase 2 A/C	H M R B Ch	VVB	Rb	100 µL	AB1621	\$265
P2A Immunoprecipitation Phosphatase Assay Kit		PA		25 assays	17-313	\$379
Anti-PP2A, A subunit	H M R Po Xn	WB IP	Rb IgG	100 ha	07-250	\$309
Anti-PTPA, clone 5G3	НМ	WB	Μ	100 µL	05-941	\$279
Anti-PP2A, A subunit, clone 4G7	H M R Xn	WB	M lgG1	200 µg	05-657	\$329
PP2A C $lpha$ subunit SMARTpool siRNA reagent	Н	RNAi		5 nmol	M-003598	\$375
Anti-PP2A, B' subunit	H M Rb	WB	Rb IgG	200 µg	07-334	\$309
Anti-PP2A, B' subunit, clone 2G9	h m r rb b	VVB IP IH Po Xn	M lgG	200 µL	05-592	\$319
Anti-PP2A, C subunit	H M R Rb	WB IP	Rb IgG	200 µg	06-222	\$309
Anti-PP2A, C subunit	HMRB	WB	Rb IgG	100 µL	07-324	\$309
Anti-Protein Phosphatase 2 C $\alpha/\beta$	HMRB	WB	Rb	100 µg	AB4090	\$240
Anti-PP2A, C subunit, clone 1D6	H M R Rb B Xn Y	WB IP IC	M IgG <sub>2bĸ</sub>	200 µg	05-421	\$329
Anti-PP2A, C subunit, clone 7A6	НМҮ	WB IP	M IgG	200 pg 200 µL	05-545	\$329
Anti-methyl-PP2A, C subunit, clone 2A10	H M R Rb Po	WB	M IgG	200 pt 2 mL	05-546	\$329
	Ch Dr Y	VVD	in igo	ZIIIL	05-540	ψυΖ.
Anti-phospho-PP2A, C subunit (Tyr307), clone 4B10	VVR	ELISA	$M \log G_1$	100 µL	05-547	\$329
Anti-PP2A, C subunit, demethylated, clone 4B7	VVR	WB IP IC	M lgG	200 µL	05-577	\$329
2A-methylesterase						
Anti-PP2A-methylesterase/PME-1	HMR	VVB IP	Rb IgG	200 µg	07-095	\$300
Anti-PP2A-methyltransferase/PPMT1, clone 4A4	M R	WB	M lgG1	100 µg	05-849	\$319
EN						
Anti-PTEN, C-terminus, clone A2b1	HMR	WB IP IC IH IH(P)	M lgG1	100 µg	MAB4037	\$245
Anti-phospho-PTEN (Ser370)	Μ	WB	Rb IgG	100 µL	07-889	\$309
Anti-phospho-PTEN (Ser385)	Н	VVB	Rb IgG	100 µL	07-890	\$309
Anti-phospho-PTEN (Ser380/Thr382/ Thr383/Ser385)	НМ	WB	Rb IgG	100 µL	07-891	\$309
PTEN Malachite Green Assay Kit		PA		1 kit	17-351	\$429
PTEN Enzyme Assay Buffer, 5X		PA		1 ml	20-165	\$54
PTEN SMARTpool siRNA reagent	Н	RNAi		5 nmol	M-003023	\$375
MA						φ0/ t
Anti-PUMA	Н	WB	Rb IgG	100 µL	07-669	\$309
Anti-PUMA, C-terminus	НМ	WB	Rb	50 µL	AB4510	\$19
f						
Anti-Raf-1	Н	WB IPK	Rb IgG	200 µg	07-396	\$319
Anti-B-Raf	HMR	WB IPK	Rb IgG	200 µg	07-453	\$319
Anti-B-Raf, NT	HMR	WB KA	Rb IgG	200 pg 200 µL	07-583	\$319
Anti-C-raf-1, a.a. 38-53	Н	WB ELISA	Sh	100 µg	CBL445	\$23
Anti-C-rai-1, d.a. 36-33 Anti-Raf-1, clone AM223	11	WB IP				\$319
			Rb IgG	100 µg	05-739	
Anti-phospho-Raf-1 (Ser338)	M	WB	R IgG1	100 µg	05-534	\$32
Anti-phospho-Raf-1 (Ser338)	M	WB	M lgG	200 µg	05-538	\$329
Anti-phospho-Raf1 (Ser259)	Н	WB	Rb IgG	100 µL	07-811	\$309
Anti-phospho-Raf1 (Ser43)	Н	WB	Rb IgG	100 µL	07-812	\$309
Anti-phospho-Raf-1 (Ser621)	Н	WB	Rb IgG	100 µL	07-813	\$309
Raf-1-RBD GST Protein, Ras binding domain				300 µg	SGT223	\$325

<b>Description</b> Raf-1 Kinase Cascade Assay Kit	Species	<b>Applications</b> IPK KA	Host	<b>Quantity</b> 1 kit	<b>Cat. No.</b> 17-357	<b>Price</b> \$474
B-Raf Kinase Cascade Assay Kit		IPK KA		1 kit	17-358	\$474
B-Raf Kinase Assay Kit, Chemiluminescence Detection		WB KA		1 kit	17-359	\$474
Raf-1 Kinase Assay Kit, Chemiluminescence Detection		KA		1 kit	17-360	\$474
Raf-1 SMARTpool siRNA reagent		RNAi		5 nmol	M-003601	\$375
siRNA plasmid, pKD-Raf-1-v4	HMR	RNAi		5 µg	62-085	\$349
siRNA plasmid, pKD-Raf-1-v6	Н	RNAi		5 µg	62-086	\$349
Raf-1 siRNA/siAb Assay Kit	Н	WB RNAi		1 kit	60-109	\$649
Raf-1 cDNA (wt) in pUSEamp		TFX		5 µg	21-111	\$599
AIDD						
Anti-RAIDD, C-terminus	Н	WB	Rb	100 µg	AB16958	\$205
siRNA plasmid, pKD-RAIDD/CRADD-v1	H	RNAi	NB	5 µg	62-157	\$349
siRNA plasmid, pKD-RAIDD/CRADD-v2	H	RNAi		5 µg	62-158	\$349
		IN W II		9 P9	02100	Ψ0 <del>Π</del> 7
ICK						
Anti-RICK, N-terminus	Н	WB	Rb	100 hð	AB17004	\$220
OCK						
Anti-ROCK-1	НМ	WB	Rb	100 µg	AB3885	\$240
siRNA plasmid, pKD-ROKβ/ROCK-I-v4		RNAi		5 µg	62-228	\$349
Anti-ROKa/ROCK-II	R	WB IP	Rb IgG	200 µg	07-443	\$319
Anti-ROK $\alpha$ /ROCK-II, clone A9W4	HR	WB IP	Rb IgG	100 µL	05-841	\$309
siRNA plasmid, pKD-ROKa/ROCK-II-v4	HMR	RNAi		5 µg	62-075	\$349
siRNA plasmid, pKD-ROKa/ROCK-II-v6	H M R	RNAi		5 µg	62-076	\$349
70.54 Vienee						
70 S6 Kinase Anti-S6 Kinase	H M R B Po	WB IP ELISA	Rb	100 µg	AB3191	\$195
	HMR	WB IF LLISA WB			06-926	\$309
Anti-p70 S6 Kinase			Rb IgG	200 µg		
Anti-p70 S6 Kinase	HMR	WB IP IPK	Rb IgG	200 µg	07-402	\$329
Anti-S6 Kinase II	H Mk	WB	Sh IgG	100 µg	07-173	\$299
Anti-p70 S6 Kinase, clone SB20	HMR	WB	Rb IgG	100 µL	05-781	\$319
Anti-phospho-Ribosomal Protein S6 (Ser235), clone MC27	H M R	WB	Rb IgG	100 µL	05-795	\$329
Anti-phospho-p70 S6 Kinase (Thr412)	НМ	WB	Rb IgG	100 hð	07-018	\$309
Anti-phospho-Ribosomal Protein S6 (Ser235)	M R Ch Xn	WB	Rb IgG	200 µg	07-433	\$329
p70 S6 Kinase Assay Kit		KA		1 kit	17-136	\$229
Kemptide, 1mM		KA		500 µL	20-199	\$159
siRNA plasmid, pKD-p70 S6 Kinase II-v2	Н	RNAi		5 µg	62-101	\$349
siRNA plasmid, pKD-p70 S6 Kinase II-v4	Н	RNAi		5 µg	62-102	\$349
S6 Kinase Substrate Cocktail		KA		1 mL	20-122	\$135
mac/Diablo						
Anti-Smac/Diablo	HMR	WB	Rb	100 µg	AB3609	\$220
Anti-Smac/Diablo, clone 17-1-87	Н	IP	M IgG1 <sub>K</sub>	100 µg	05-674	\$319
Anti-Smac/Diablo, clone 78-1-118	H	WB	M IgG <sub>1</sub>	100 µg	MAB4626	\$265
Anti-Smac/Diablo, clone 78-1-118	Н	WB	M IgG	200 µg	05-681	\$319
		,,,,,		200 99	00001	<i><b>Q</b></i> <b>OT</b>
ODD			DI	100	401/510	¢000
Anti-SODD, N-terminus	H M R	WB	Rb	100 hð	AB16518	\$220
RAD						
Anti-TRADD, clone 3E11	Н	WB IP	M lgG $_{2a\kappa}$	100 µg	05-473	\$309
RAIL						
Anti-TRAIL, C-terminus	Н	WB	Rb	100 µg	AB16957	\$220
TRAIL, recombinant human				50 µg	GF092	\$220
ansglutaminase				250	06 471	\$210
Anti-Transglutaminase type II Anti-Tissue Transglutaminase, FN binding domain,	M Gp H IC	WB IP M	Gt antiserum	250 µL 100 µg	06-471 MAB3839	\$319 \$295
Anti-Tissue Transgiutaminase, FIN binding domain, clone 4G3		/ • 1		roo ha	111403034	ΦΖΑϽ

<b>Description</b> Other Apoptosis Reagents	Species	Applications	Host	Quantity	Cat. No.	Price
Apoptosis Inducer Set				1 kit	APT800	\$230
Apoptosis Sampler Kit	VVR		Rb	1 kit	APT880	\$435
Propidium Iodide Solution				1 mL	S7109	\$60
Propidium Iodide/Antifade Solution				1 mL	S7112	\$60
DAPI/Antifade Solution, Ready-to-Use				1 mL	S7113	\$60
Antifade Solution				1 mL	S7114	\$60
Staurosporine		KA		200 µg	19-123	\$179

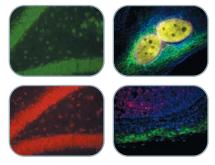


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