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## Goat anti-Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody, Alexa Fluor 488

Catalog Number A-11034 Product data sheet

Outding Humber A 1100+		
Details		
Size	500µL	
Host/Isotope	Goat / IgG	
Class	Polyclonal	
Туре	Secondary Antibody	
Immunogen	Gamma Immunoglobins Heavy and Light chains	
Target Class	IgG	
Cross Adsorption	Against bovine IgG, goat IgG, mouse IgG, rat IgG and human IgG	
Antibody Form	Whole Antibody	
Conjugate	Alexa Fluor® 488	
Form	liquid	
Concentration	2 mg/ml	
Purification	purified	
Storage buffer	PBS, pH 7.5	
Contains	5mM sodium azide	
Storage Conditions	4° C, store in dark	

Species Reactivity	
Tested species reactivity	Rabbit
Published species reactivity	Not Applicable
Tested Applications	Dilution *
Flow Cytometry (Flow)	1-10 μg/mL
Immunocytochemistry (ICC)	1-10 µg/mL
Immunofluorescence (IF)	1-10 µg/mL
Published Applications	
Immunocytochemistry (ICC)	See 4 publications below
Immunohistochemistry (Frozen) ( IHC (F))	See 1 publications below
Miscellaneous PubMed (MISC)	See 348 publications below

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

### Product specific information

To minimize cross-reactivity, the goat anti-rabbit IgG whole antibodies have been highly cross-adsorbed against bovine IgG, goat IgG, mouse IgG, rat IgG, and human IgG. Cross-adsorption or pre-adsorption is a purification step to increase specificity of the antibody resulting in higher sensitivity and less background staining. The secondary antibody solution is passed through a column matrix containing immobilized serum proteins from potentially cross-reactive species. Only the nonspecific-binding secondary antibodies are captured in the column, and the highly specific secondaries flow through. Further passages through additional columns result in 'highly cross-adsorbed' preparations of secondary antibody. The benefits of these extra steps are apparent in multiplexing/multicolor-staining experiments where there is potential cross-reactivity with other primary antibodies or in tissue/cell fluorescent staining experiments where there may be the presence of endogenous immunoglobulins. Alexa Fluor dyes are among the most trusted fluorescent dyes available today. Invitrogen™ Alexa Fluor 488 dye is a bright, green-fluorescent dye with excitation ideally suited to the 488 nm laser line. For stable signal generation in imaging and flow cytometry, Alexa Fluor 488 dye is pH-insensitive over a wide molar range. Probes with high fluorescence quantum yield and high photostability allow detection of low-abundance biological structures with great sensitivity. Alexa Fluor 488 dye molecules can be attached to proteins at high molar ratios without significant self-quenching, enabling brighter conjugates and more sensitive detection. The degree of labeling for each conjugate is typically 2-8 fluorophore molecules per IgG molecule; the exact degree of labeling is indicated on the certificate of analysis for each product lot. Using conjugate solutions: Centrifuge the protein conjugate solution briefly in a microcentrifuge before use; add only the supernatant to the experiment. This step will help eliminate any protein aggregates that may have formed during storage, thereby reducing nonspecific background staining. Because staining protocols vary with application, the appropriate dilution of antibody should be determined empirically. For the fluorophore-labeled antibodies a final concentration of 1-10 µg/mL should be satisfactory for most immunohistochemistry and flow cytometry applications.

### **Background/Target Information**

We offer an extensive line of Invitrogen™ secondary antibody conjugates with well-characterized specificity and labeled with a wide selection of premium fluorescent dyes, including Invitrogen™ Alexa Fluor™ fluorescent dyes. Fluorescent secondary antibody conjugates are useful in the detection, sorting, or purification of its specified target and ideal for fluorescence microscopy and confocal laser scanning microscopy, flow cytometry, and fluorescent western

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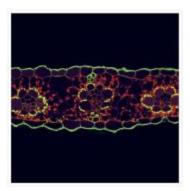
detection. The breadth of fluorescent markers we offer allows our reagents to be tailored to almost any fluorescent detection system. Secondary antibodies may be provided in three formats: whole IgG, divalent F(ab')2 fragments, and monovalent Fab fragments. Because of the high degree of conservation in the structure of many immunoglobulin domains, most class-specific secondary antibodies must be affinity-purified and cross-adsorbed to achieve minimal cross-reaction with other immunoglobulins. Our secondary antibody conjugates are most commonly prepared by immunizing the host animal with a pooled population of immunoglobulins from the target species and can be further purified and modified (e.g., immunoaffinity chromatography, antibody fragmentation , label conjugation, etc.) to generate highly specific reagents. In the first round of purification, whole immunoglobulins binding to the immunizing antibody are recovered and mainly consist of the ~150-kDa IgG class. Further purification, for example, with Protein A or G, removes all unwanted immunoglobulin classes except the affinity-purified antibodies that react with the target-specific immunoglobulin heavy and/or light chains.

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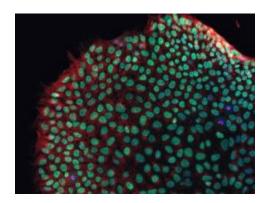


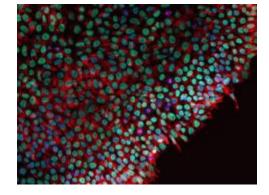
A 2.0 µm maize leaf section illustrating the immunolocalization of the enzyme ribulose bisphosphate carboxylase (rubisco) in the chloroplasts of the bundle sheath cells surrounding the vascular bundles. Maize is a C4 plant and, as a result, spatially segregates components of the photosynthetic process between the leaf mesophyll and the bundle sheath. Rubisco was localized using a rabbit anti-rubisco antibody and visualized using the highly cross-adsorbed Alexa Fluor® 488 goat anti-rabbit IgG antibody ( Cat. No. A11034). The remaining fluorescence is due to the autofluorescence of chlorophyll, which appears red and is localized to the mesophyll plastids; lignin, which appears dull green and is localized to the xylem of the vascular bundle; and cutin, which appears bright green and is localized to the cuticle outside the epidermis. Image contributed by Todd Jones, DuPont.

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Human iPSC Staining Human iPSCs were cultured on glass slides under feeder-free conditions in StemPro® hESC Medium (A1000701 ). Cells were fixed and permed with the Image-iT ® Fixation/ Permeabilization Kit ( R37602). Oct4 (green) expression was visualized using anti-Oct4 primary Ab and Alexa Fluor® 488 secondary Ab (A11034). Tubulin (red) expression was visualized using anti-tubulin primary Ab ( 322600) and Alexa Fluor® 594 secondary Ab ( A11005). Nuclei (blue) were labeled with NucBlue ™ Fixed Cell Stain ( R37606). Images were collected on the FLoid™ Cell Imaging Station ( 4471136).

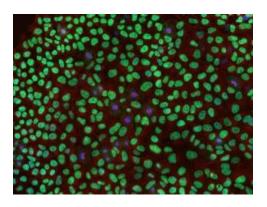
### Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody (A-11034)

Human iPSC Staining Human iPSCs were cultured on glass slides under feeder-free conditions in StemPro® hESC Medium (A1000701 ). Cells were fixed and permed with the Image-iT ® Fixation/ Permeabilization Kit ( R37602). Oct4 (green) expression was visualized using anti-Oct4 primary Ab and Alexa Fluor® 488 secondary Ab (A11034). Tubulin (red) expression was visualized using anti-tubulin primary Ab ( 322600) and Alexa Fluor® 594 secondary Ab ( A11005). Nuclei (blue) were labeled with NucBlue ™ Fixed Cell Stain ( R37606). Images were collected on the  $\mathsf{FLoid}^\mathsf{TM}$ Cell Imaging Station ( 4471136).

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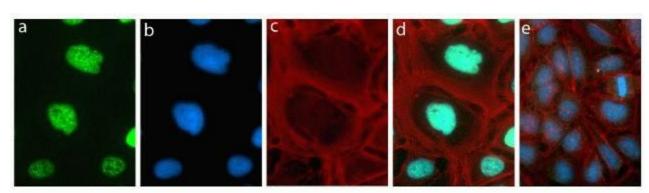




Human iPSCs were cultured on glass slides under feeder-free conditions in StemPro® hESC Medium (A1000701 ). Cells were fixed and permed with the Image-iT ® Fixation/ Permeabilization Kit ( R37602). Oct4 (green) expression was visualized using anti-Oct4 primary Ab and Alexa Fluor® 488 secondary Ab (A11034). Tubulin (red) expression was visualized using anti-tubulin primary Ab ( 322600) and Alexa Fluor® 594 secondary Ab ( A11005). Nuclei (blue) were labeled with NucBlue ™ Fixed Cell Stain ( R37606). Images were collected on the FLoid™ Cell Imaging Station ( 4471136).

### Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody (A-11034)

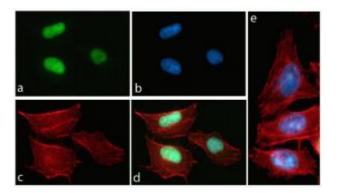
Immunocytochemistry analysis of U2OS cells stained with Rb [pSpT249/ 252] ABfinity™ Recombinant Rabbit Monoclonal Antibody, using (A) Alexa Fluor® 488 Goat Anti-Rabbit was used as secondary (green ). (B) DAPI was used to stain the nucleus (blue) and (C) Alexa Fluor® 594 phalloidin was used to stain actin (red). (D) Composite image of cells showing nuclear localization of phosphorylated Rb. (E) Composite image of cells showing competition with the phospho Rb [pSpT249 /252].



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### Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody (A-11034)

Immunocytochemistry analysis of HeLa cells stained with Tau [pT231] ABfinity™ Recombinant Rabbit Monoclonal Antibody, using (A) Alexa Fluor® 488 Goat Anti-Rabbit was used as secondary (green). (B) DAPI was used to stain the nucleus (blue) and (C) Alexa Fluor® 594 phalloidin was used to stain actin (red). (D) Composite image of cells showing nuclear localization of phosphorylated Tau. (E) Composite image of cells showing competition with the phospho Tau [pT231] peptide.

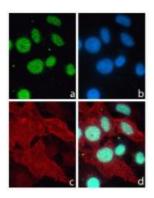
### Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody (A-11034)

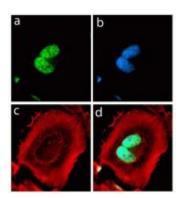
Immunocytochemistry analysis of HeLa cells serum starved for 36 hr, followed by serum release for 8 hr. The cells were then stained with Ki-67 ABfinity™ Recombinant Rabbit Oligoclonal Antibody, using (A) Alexa Fluor® 488 Goat Anti-Rabbit was used as secondary (green). (B) DAPI was used to stain the nucleus (blue) and (C) Alexa Fluor® 594 phalloidin was used to stain actin (red). (D) Composite image of cells showing nuclear localization of Ki-67.

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Immunocytochemistry analysis of U2OS cells stained with PiTX3, Rabbit mAb (5H10L5), ABfinity™ Recombinant Monoclonal Antibody, using a: Alexa Fluor®488 goat anti-rabbit was used as secondary ( green). b: DAPI was used to stain the nucleus (blue) and c: Alexa Fluor® 594 wheat germ agglutinin was used to stain glycoconjugates on cell membrane (red). d: Composite image of cells showing nuclear localization of PiTX3.

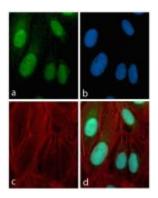
### Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody (A-11034)

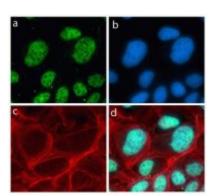
Immunocytochemistry analysis of HeLa cells stained with Alpha-Synuclein Rabbit Recombinant Oligoclonal Antibody using (A) Alexa Fluor® 488 Goat Anti-Rabbit was used as secondary (green). (B) DAPI was used to stain the nucleus (blue) and (C) Alexa Fluor® 594 phalloidin was used to stain actin (red). (D) Composite image of cells showing cytoplasmic and nuclear localization of Alpha-Synuclein. (Cat. No. 701085)

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Immunocytochemistry analysis of U2OS cells stained with Pax 3 (HUP2) , Rabbit mAb (16H22L10). ABfinity™ Recombinant Monoclonal Antibody (Cat. No. 701147), using (A) Alexa Fluor® 488 Goat Anti-Rabbit was used as secondary (green). (B) DAPI was used to stain the nucleus (blue) and (C) Alexa Fluor® 594 phalloidin was used to stain actin (red). (D) Composite image of cells showing nuclear localization of Pax 3.

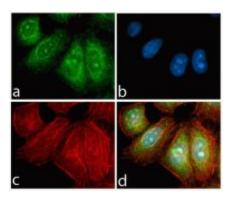
### Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody (A-11034)

Immunocytochemistry analysis of U2OS cells stained with p21 (CDKN1), Rabbit mAb (2H2L13) (Cat . No. 701151), ABfiniťy<sup>⊤</sup>™ Recombinant Monoclonal Antibody, using (A) Alexa Fluor® 488 Goat Anti-Rabbit was used as secondary (green). (B) DAPI was used to stain the nucleus (blue) and (C) Alexa Fluor® 594 phalloidin was used to stain actin (red). (D) Composite image of cells showing nuclear localization of p21.

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### Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody (A-11034)

Immunocytochemistry analysis of HeLa cells stained with PDGFR-alpha ABfinity™ Recombinant Rabbit Monoclonal Antibody using (A) Alexa Fluor® 488 Goat Anti-Rabbit was used as secondary (green). (B) DAPI was used to stain the nucleus (blue) and (C) Alexa Fluor® 594 phalloidin was used to stain actin (red). (D) Composite image of cells showing cytoplasmic and nuclear membrane localization of PDGFR-alpha.

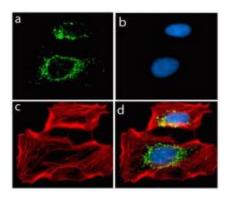
### Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody (A-11034)

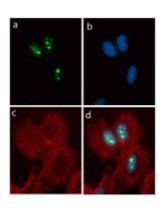
Immunocytochemistry analysis of U2OS cells stained with Nucleostemin (GNL3), Rabbit mAb ( 3H20L2), ABfinity™ Recombinant Monoclonal Antibody, using a: Alexa Fluor® 488 goat anti-rabbit was used as secondary (green). b: DAPI was used to stain the nucleus (blue) and c: Alexa Fluor® 594 phalloidin was used to stain actin (red). d: Composite image of cells showing nuclear localization of Nucleostemin.

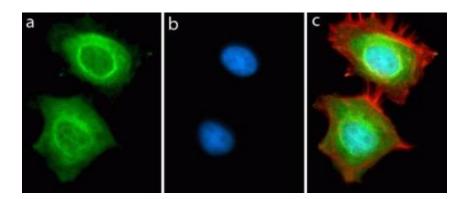
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Immunocytochemistry analysis of HeLa cells stained with Adiponectin ABfinity™ Recombinant Rabbit Monoclonal Antibody, using (A) Alexa Fluor® 488 Goat Anti-Rabbit was used as secondary (green). (B) DAPI was used to stain the nucleus (blue) and (C) Alexa Fluor® 594 phalloidin was used to stain actin (red). (D) Composite image of cells showing sub cellular localization in perinuclear region.

### Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody (A-11034)

Immunocytochemistry analysis of HeLa cells stained with Nucleostemin ABfinity™ Recombinant Rabbit Oligoclonal Antibody, using (A) Alexa Fluor® 488 Goat Anti-Rabbit was used as secondary (green). (B) DAPI was used to stain the nucleus (blue) and (C) Alexa Fluor® 594 phalloidin was used to stain actin (red). (D) Composite image of cells showing nuclear localization of Nucleostemin.

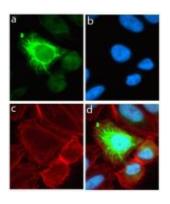
### Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody (A-11034)

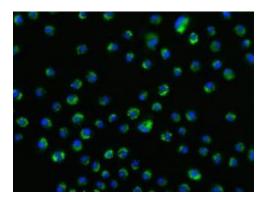
Immunocytochemistry analysis of HeLa cells stained with Visfatin ABfinity™ Recombinant Rabbit Monoclonal Antibody using (A) Alexa Fluor® 488 Goat Anti-Rabbit was used as secondary (green). (B) DAPI was used to stain the nucleus (blue). Alexa Fluor® 594 phalloidin was used to stain actin (red). ( D) Composite image of cells showing cytoplasmic and nuclear localization of Visfatin.

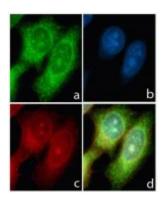
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Immunocytochemistry analysis of HeLa cells stained with Musashi-1 ABfinity™ Recombinant Rabbit Oligoclonal Antibody, using (A) Alexa Fluor® 488 Goat Anti-Rabbit was used as secondary (green). (B) DAPI was used to stain the nucleus (blue) and (C) Alexa Fluor® 594 phalloidin was used to stain actin (red). (D) Composite image of cells showing cytoplamic localization of Musashi-1.

### Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody (A-11034)

Gibco® Human Corneal Epithelial cells were treated with 30uM Chloroquine overnight. The following day the cells were fixed and permeabilized and labeled with 0.5ug/mL anti-LC3B with a goat anti mouse Alexa Fluor® 488 secondary (Green) Cells were counter stained with 1ug/mL Hoechst 33342 ( Blue) Images were acquired with a Molecular **Devices ImageXpress** High content imager

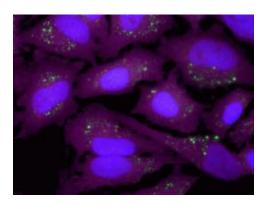
### Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody (A-11034)

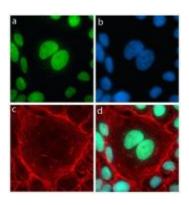
Immunocytochemistry analysis of HeLa cells stained with STAT6 ABfinity™ Recombinant Rabbit Monoclonal Antibody using (A) Alexa Fluor® 488 Goat Anti-Rabbit was used as secondary (green). (B) DAPI was used to stain the nucleus (blue) and (C) Alexa Fluor® 594 phalloidin was used to stain actin (red). (D) Composite image of cells showing cytoplasmic localization of STAT6.

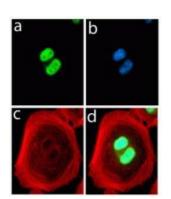
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HeLa cells were treated with 60um Chloroquine overnight. The following day cells were fixed and permeabilized and subsequently labeled with 1ug/mL anti-LC3B with a goat anti rabbit Alexa Fluor® 488 secondary ( green). Cells were counter stained with HCS Cellmask™ Deep Red ( purple) and 1ug/mL Hoechst 33342 (Blue). Cells were imaged on a Nikon D200 wide field microscope

### Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody (A-11034)

Immunocytochemistry analysis of U2OS cells stained with GATA2 ABfinity™ Recombinant Rabbit Oligoclonal Antibody, using (A) Alexa Fluor® 488 Goat Anti-Rabbit was used as secondary (green). (B) DAPI was used to stain the nucleus (blue) and (C) Alexa Fluor® 594 phalloidin was used to stain actin (red). (D) Composite image of cells showing nuclear localization of GATA2.

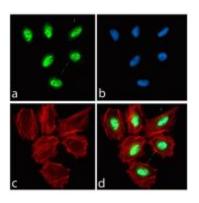
### Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody (A-11034)

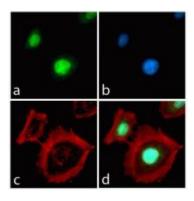
Immunocytochemistry analysis of HeLa cells stained with FABP4 ABfinity™ Recombinant Rabbit Monoclonal Antibody, using (A) Alexa Fluor® 488 Goat Anti-Rabbit was used as secondary (green). (B) DAPI was used to stain the nucleus (blue) and (C) Alexa Fluor® 594 phalloidin was used to stain actin (red). (D) Composite image of cells showing nuclear localization of FABP4.

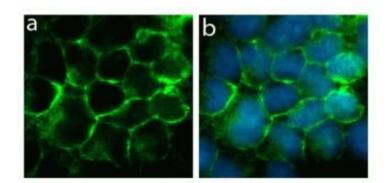
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Immunocytochemistry analysis of HeLa cells stained with Pax 3 ABfinity ™ Recombinant Rabbit Monoclonal Antibody, using (A) Alexa Fluor® 488 Goat Anti-Rabbit was used as secondary (green ). (B) DAPI was used to stain the nucleus (blue) and (C) Alexa Fluor® 594 phalloidin was used to stain actin (red). (D) Composite image of cells showing nuclear localization of Pax 3.

### Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody (A-11034)

Immunocytochemistry analysis of HeLa cells stained with Rex1 ABfinity ™ Recombinant Rabbit Monoclonal Antibody, using (A) Alexa Fluor® 488 Goat Anti-Rabbit was used as secondary (green ). (B) DAPI was used to stain the nucleus (blue) and (C) Alexa Fluor® 594 phalloidin was used to stain actin (red). (D) Composite image of cells showing nuclear localization of Rex1.

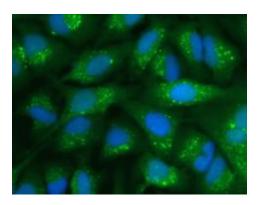
### Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody (A-11034)

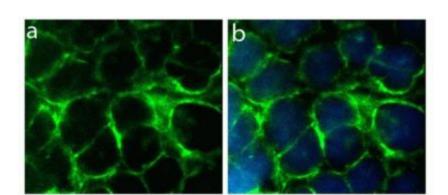
Immunocytochemistry analysis of U2OS cells stained with Connexin 40 (C-term) ABfinity™ Recombinant Rabbit Oligoclonal Antibody, using (A) Alexa Fluor® 488 Goat Anti-Rabbit was used as secondary (green ). DAPI was used to stain the nucleus (blue). (B) Composite image of cells showing cell junction localization of Connexin 40 (C-term).

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HeLa cells were treated with 60um Chloroquine overnight. The following day cells were fixed and permeabilized and subsequently labeled with 1ug/mL anti-LC3B with a goat anti rabbit Alexa Fluor® 488 secondary (green). Cells were counter stained with 1ug/mL Hoechst 33342 (Blue). Cells were imaged on a Nikon D200 wide field microscope

### Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody (A-11034)

Immunocytochemistry analysis of HeLa cells stained with Connexin 36 ABfinity™ Recombinant Rabbit Monoclonal Antibody, using (A) Alexa Fluor® 488 Goat Anti-Rabbit was used as secondary (green). DAPI was used to stain the nucleus (blue). (B) Composite image of cells showing cell membrane and junction localization of Connexin 36.

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LC3] and p62 to assess autophagy.   Analytical blochemistry (May 2015; 477: 13)   *Routine Western blot to check autophagic flux: cautions and recommendations."     Author(s):Gomez-Sanchez R. Pizarro-Estrella E. Yakhine-Diop SM. Rodriguez-Arribas M.Bravo-San Pedro JM. Fuentes JM Goraciaez-Polo R.     Author(s):Gomez-Sanchez R. Pizarro-Estrella E. Yakhine-Diop SM. Rodriguez-Arribas M.Bravo-San Pedro JM. Fuentes JM Goraciaez-Polo R.     Author(s):Gomez-Sanchez R. Pizarro-Estrella E. Yakhine-Diop SM. Rodriguez-Arribas M.Bravo-San Pedro JM. Fuentes JM Goraciaez-Polo R.     Author(s):Gomez-Sanchez R. Pizarro-Estrella E. Yakhine-Diop SM. Rodriguez-Arribas M.Bravo-San Pedro JM. Fuentes JM.     Author(s):Arribos W. Author(s):Arribos W. Author(s):Arribos W.     Author(s):Arribos W.	4 Immunocytochemistry Re	eferences
LC3) and p62 to assess autophagy.  Analytical biochemistry (May 2015; 477: 13)  "Acutine Western blot to check autophagic flux: cautions and recommendations."  Author(s):Comez-Sanchez R, Pizarro-Sariella E, Takhime-Diop SM, Rodriguez-Ambas M, Bravo-San Pedro JM, Fuentes JM, PubMed Article URL-they/dx.doi.org/10.1016/j.ab. 2015.02.020  A-11034 was used in immunocytochemistry to identify and characterize the Nrb1z-Beclin 1 interaction.  The Journal of biological chemistry (Sep 2014; 289: 28021)  **Not Applicable / Not Cited and Common Co	Species / Dilution	Summary
*Routine Western blot to check autophagic flux: cautions and recommendations." Author(s):Gömes-Sanchez R.P.Ezrac-Esrella E.Yakhine-Diop SM, Rodríguez-Arribas M, Bravo-San Pedro JM, Fuentes JM González-Polo RA PubMed Article URL:http://dx.doi.org/10.1016/j.ab.2015.02.020  A-11034 was used in immunocytochemistry to identify and characterize the Nrb12-Beclin 1 interaction.  The Journal of biological chemistry (Sep 2014; 289: 28021)  **Not Applicable / Not Cited  **PLOS pathogens (Jan 2014; 10; null) **Parvovirus-induced depletion of cyclin B1 prevents mitotic entry of infected cells." Author(s):Anglicable / Not Cited  **Not Applicable / Not Cited  **Not Applica	Not Applicable / 1:1000	A-11034 was used in immunocytochemistry to discuss the limitations of using microtubule-associated protein 1 light chain 3 (LC3) and p62 to assess autophagy.
The Journal of biological chemistry (Sep 2014; 289: 26021)  "Not Applicable / Not Cited  "Not Zprotein suppresses autophagy by modulating Atgr14L protein-containing Beclin 1-Vps34 complex architect and reducing intracellular phosphatidylinositoria-3 phosphate levels."  Author(s): Zhong Y, Morris DH, Jin L, Patel MB, Karunakaran SK, Fu YJ, Matuszak EA, Weiss HL, Chait BT, Wang QJ PubMed Article URL: http://dx.doi.org/10.1074/jbc.M114.561134  A-11034 was used in immunocytochemistry to elucidate how parvovirus minute virus of mice halts the cell cycle.  PLoS pathogens (Jan 2014; 10: null)  "Parvovirus-Induced depletion of cyclin B1 prevents mitotic entry of infected cells." Author(s):Adeyemi RO, Pintel DJ PutMed Article URL: http://dx.doi.org/10.1371/journal.ppat.1003891  A-11034 was used in immunocytochemistry to use ROCK1 and ROCK2 knock out mice to study doxorubicin-induced stre fiber disassembly and cell detachment.  Not Applicable / Not Cited  Old Cell cycle (Georgeotewn. Tex.) (May 2013; 12: 1492)  "Dissecting the roles of ROCK Isoforms in stress-induced cell detachment." Author(s):Shi.1, Surma M. Zhang I. Wei L PubMed Article URL: http://dx.doi.org/10.4161/cc.24699  Immunohistochemistry (Frozen) References  Species / Dilution  Summary  A-11034 was used in immunohistochemistry - frozen section to assess the effects of tissue-type plasminogen activator treatment after oral anticoagulation with invaroxaban or aphaban compared with warfarin or placebo  Not Applicable / 1:500  Not Applicable / 1:500  Not Applicable / 1:500  Summary  A-11034 was used in immunocytochemistry to characterize STAG3-deficient mice.  The EMBO journal (Jun 2014; 33: 12:56)  "Molecular induced depletion of the protein of place and pl		"Routine Western blot to check autophagic flux: cautions and recommendations."  Author(s):Gómez-Sánchez R,Pizarro-Estrella E,Yakhine-Diop SM,Rodríguez-Arribas M,Bravo-San Pedro JM,Fuentes JM, González-Polo RA
Not Applicable / Not Cited  **Not/2 protein suppresses autophagy by modulating Atg14L protein-containing Beclin 1-Vps34 complex architec and reducing intracellular phosphatidylinosito-3 phosphate levels."  Author(s):Zhong Y,Morris DH,Jin L,Patel MS,Karunakaran SK,Fu YJ,Matuszak EA,Weiss HL,Chait BT,Wang QJ PubMed Article URL:http://dx.doi.org/10.1074/jbc.M114.561134  A-11034 was used in immunocytochemistry to elucidate how parvovirus minute virus of mice halts the cell cycle.  PLoS pathogens (Jan 2014; 10: null)  "Parvovirus-induced depletion of cyclin B1 prevents mitotic entry of infected cells."  Author(s):Adeyem RO,Pintel D PubMed Article URL:http://dx.doi.org/10.1371/journal.ppat.1003891  A-11034 was used in immunocytochemistry to use ROCK1 and ROCK2 knock out mice to study doxorubicin-induced stre fiber disassembly and cell detachment.  **Cell cycle (Georgetown, Tex.) (May 2013; 12: 1492)  **Dissecting the roles of ROCK isoforms in stress-induced cell detachment."  Author(s):Shi J,Surma M,Zhang L,Wei L  PubMed Article URL:http://dx.doi.org/10.4161/cc.24699  1 Immunohistochemistry (Frozen) References  Species / Dilution  Summary  A-11034 was used in immunohistochemistry - frozen section to assess the effects of tissue-type plasminogen activator treatment after oral anticoagulation with rivanoxaban or apixaban compared with warfarin or placebo  Stroke: a journal of cerebral circulation (Aug 2014; 45: 2404)  "Rivaroxaban and apixaban reduce hemorrhagic transformation after thrombolysis by protection of neurovasculurit in rat.  Author(s):Kono S, Yamashita T, Deguchi K,Omoch Y,Yunoki T, Sato K,Kurata T, Hishikawa N,Abe K  PubMed Article URL:http://dx.doi.org/10.1161/STROKEAHA.114.005316  Author(s):Was used in immunocytochemistry to characterize STAG3-deficient mice.  The EMBO journal (Jun 2014; 33: 1256)  "Meiotic cohesin STAG3 is required for chromosome axis formation and sister chromatid cohesion."  Author(s):Karisson AB,Washingion J,Dimitrova V,Hooper C,Shekhtman A,Bakowska JC  PubMed Article URL:http://dx.d		A-11034 was used in immunocytochemistry to identify and characterize the Nrbf2-Beclin 1 interaction.
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