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After the manuscript has been checked by journal staff, the Corresponding Author will receive an e-mail acknowledging receipt of the manuscript. The e-mail contains links to the Submission Form and Color Charges Form (if applicable). The Corresponding Author must download, sign, and fax these forms to 301-634-7831 to complete the submission. The manuscript will not be sent for review until *The JI* Editorial Office receives these forms signed by the Corresponding Author. Please contact infoji@aai.org if you do not receive the acknowledgment e-mail. Please do NOT use the hard copy forms found in old copies of the printed journal.

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#### MANUSCRIPT PREPARATION

General Guidelines: A 12-point serif font, preferably Times New Roman, is required. Do not use compressed type format. Double-space entire manuscript. The average length of full-length articles is eight printed pages. Instructions for estimating the printed length of a manuscript are included below. Each of the following components should begin on a separate page:

- 1. The **Title Page** must include the full title; a running title (not to exceed 60 characters); each author's full name (first name, middle initial, last name); the affiliations of all authors and their institutions, departments or organizations (use the following symbols in this order: \*, †, ‡, §, ¶,  $\parallel$ , #, \*\*, ††, ‡‡, §§, ¶¶,  $\parallel$   $\parallel$ , ##); and three to five keywords, selected from the Keywords list (http://www.jimmunol.org/misc/authorfulllength.shtml#keyword) that describe the topic of the manuscript. (Keywords are used in editor and reviewer assignments and are not published with the manuscript. Please note that the list of keywords does not represent an exhaustive view of what *The JI* considers important topics, but it has been found useful for reviewer assignment purposes.)
- 2. The **Abstract** must be 250 words or less for full-length manuscripts; 150 words or less for Cutting Edge. Reference citations should not be included in the *Abstract*. The species of animals or species of origin of cells used in the manuscript must be clearly stated in the Abstract.
- 3. The **Introduction**, **Materials and Methods**, **Results**, and **Discussion** sections should begin on separate pages. Do not combine the *Results* and *Discussion* sections for full-length papers.
- 4. If the manuscript contains human or animal studies, the *Materials and Methods* section must state that the studies have been reviewed and approved by an appropriate institutional review committee.
- 5. **Acknowledgments** appear immediately after the *Discussion* and before *References*.
- 6. **Grant support** must not be included in the *Acknowledgments* but should be cited as a footnote to the title.
- 7. **Disclosures** contain conflict of interest disclaimers. For more information, see the Editorial Policies Regarding Manuscript Submission.
- 8. **References** must be numbered as they appear in the text. All authors must be listed for each reference. If citations are included in tables or in figure legends, they must be numbered according to the position of citation of the table or figure in the text. Only published papers and papers "in press" may be included in the *References*. "In press" articles, i.e., not yet published, must be submitted as online attachments in PDF format at the time of article submission. NOTE: Do NOT submit as an attachment papers that are already published, e.g., manuscripts published ahead of print. Such papers must be incorporated into the *References* and cited with the DOI numbers and the publication dates. Citations of "manuscripts in preparation," "unpublished observations," and "personal communications" must appear parenthetically in the

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Format for references:

Periodicals: Wells, A. D., M. C. Walsh, D. Sankaran, and L. A. Turka. 2000. T cell effector function and anergy avoidance are quantitatively linked to cell division. *J. Immunol.* 165: 2432–2443. Books: McIntyre, T. M., and W. Strober. 1999. Gut-associated lymphoid tissue: regulation of IgA B-cell development. In *Mucosal Immunology*, 2nd ed. P. L. Ogra, J. Mestecky, E. Lamm, W. Strober, J. Bienenstock, and J. R. McGhee, eds. Academic Press, San Diego, CA. 319–356.

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- 10. **Abbreviations** that may be used without definition are provided in the Standard Abbreviations list. Spell out nonstandard abbreviations used less than three times. Nonstandard abbreviations used three or more times must be defined in a footnote. Abbreviations and their definitions must be consistent throughout the text.
- 11. **Figure legends** must be numbered with Arabic numerals in order of appearance in the text and should include a short title after the figure number. Where possible, symbols and patterns used to distinguish data should be defined in a key placed within the graphic rather than in the figure legend.
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- 13. **Figures** At initial submission, low-resolution figures may be submitted. At submission of a revised manuscript, high-resolution figures that meet the following specifications must be submitted.
  - Sizes: Figures should be submitted in final size (printed 1:1).
     Figures may be printed in one of two formats: single-column (width from 3.37 to 8.23 cm or 20 picas) or double-column (width from 12.65 to 17.1 cm or 42 picas). The single-column format is preferred. Unless the file is too large, multi-piece figures should be submitted as a single file.
  - Text and Lines: Text in figures should be 6–8 points in size, except for single letter markers, which may be 12 points. Helvetica should be used for all figure text (except for the use of symbols). If Helvetica is not available, Times Roman may be used. Line widths must be greater than one point thick or they will not appear on the PDF version of the article.
  - Numbering: Figures must be numbered as they appear in the text.
  - File Format: Figures should be in TIFF (better for halftone art, e.g., blots, photographs) or EPS (better for line art or monochrome art, i.e., anything that involves sharply delineated lines) format. While images from PowerPoint files are not usually suitable quality, as their resolution is too low for print, they may sometimes be acceptable but will result in delayed processing of the manuscript.
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- 14. **Cover art** changes with each issue of *The JI*. Authors are encouraged to submit color figures with their manuscripts for possible use as cover illustrations.
- 15. **Estimation for printed pages:** One printed page in *The JI* contains approximately 8,000 characters, including spaces. Thus, an 8-page, full-length article would contain approximately 64,000 characters. Each line in a table occupies about 60 characters for a single-column table, and 120 characters for a double-column table. Figures occupy about 180 characters per centimeter height for single-column figures, and 360 characters for double-column figures. Determine the total character count for the text of your manuscript and add the character-equivalents for the tables and figures. This will provide a reasonable estimate for the printed length of a manuscript.

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- 1. List the phone number, fax number, and e-mail address of the corresponding author on the title page.
  - 2. The *Abstract* is limited to 150 words.
- 3. The *Materials and Methods* section may be sharply limited but should be sufficient to allow the evaluation of results and conclusions.
  - 4. Authors may combine the Results and Discussion sections.
- 5. Cutting Edge articles, including figures and references, *must fit within four journal pages*. Authors should estimate the size of figures and tables and limit the text accordingly. One printed page in *The JI* contains approximately 8,000 characters, including spaces. Thus, a 4-page Cutting Edge article would contain approximately 32,000 characters. The formula for estimating the number of pages is provided in Manuscript Preparation.

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Individual manuscript, figure, and table files, and a point-bypoint reply to all referee comments, must be uploaded to the system. The revised manuscript text must be marked to show changes, using highlighting or a colored font (Microsoft Word files preferred). High-resolution figure files should be submitted. Figures The Journal of Immunology 723

must be in **TIFF** or **EPS** format; PowerPoint may be acceptable but will result in a delay in processing the submission. Instructions for figures are available at the Cadmus Digital Art website (http://cjs.cadmus.com/da/index.jsp). Authors should retain for themselves copies of all the files in their original formats.

After the manuscript has been checked by journal staff, the Corresponding Author will receive an e-mail acknowledging receipt of the revised manuscript. The e-mail contains links to the Submission Form and Color Charges Form (if applicable). The Corresponding Author must download, sign, and fax these forms to 301-634-7831 to complete the submission of the revised manuscript. Your manuscript will not be sent for review until *The JI* Editorial Office receives these forms signed by the Corresponding Author. Please contact infoji@aai.org if you do not receive the acknowledgment e-mail. Please do NOT use the hard copy forms found in old copies of the printed journal.

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Supplemental Data: All supplemental material accompanying an article must be submitted with the original paper for peer review. When submitting online, upload the file when requested. Supplemental material should be limited to short videos (must be no longer than 30 seconds and under 10 MB, with no sound or voice-over) or large tables, large sequence alignments, or large data sets such as those obtained with microarray hybridization experiments. Such supplemental data must be larger than two printed journal pages; smaller pieces of data should be included in the manuscript.

Videos must be 320 480 pixels or smaller for best viewing within a browser. Submit videos in MPG or QuickTime. Change QuickTime file extensions to ".mov" so that Web browsers will recognize the file type and play the movie. Compress videos as much as possible to help control file size. Name videos by order of citation appearance (e.g., video1.mov). Authors will be notified if problems exist with videos as submitted and will be asked to take responsibility for modifications. No editing will be done to videos at the Editorial Office. Legends or short explanations of the material must accompany all supplemental material. Links to the material will appear in two places in the online journal, in the Table of Contents and in the information box associated with the first page of the full-text article. There will not be any links in the body of the article. In the printed paper, supplemental material will be footnoted the first time mentioned: "The online version of this article contains supplemental material."

#### Distribution and depositing of materials:

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High-resolution structural data: Any paper submitted to The JI that contains new high-resolution structural data requires an accession number from the Protein Data Bank (http://www.rcsb.org/pdb/Welcome.do) and assurance that unrestricted release will occur at or before the time of publication. The accession number should be accompanied by the website address of the databank.

Nucleotide sequences: Sequences of nucleotides or amino acids longer than 50 bases/residues should not be presented in the text or in table form, but rather should be submitted as a publication-quality figure. Original nucleotide sequences, and determined nucleotide sequences encoding reported amino acid sequences, de-

scribed in the manuscript must be submitted to Gen-Bank or EMBL DataLibrary at the time of manuscript submission; an accession number and sequence availability are required at the time of publication. The accession number should be accompanied by the website address of the databank. Instructions on submission of data may be obtained directly from GenBank (Mail Stop K710, Los Alamos National Laboratory, Los Alamos, NM 87545) or from the European Molecular Biology Library, Nucleotide Sequence Library (Postfach 10.2209, Meyerhofstrasse 1, 6900 Heidelberg, Germany) or see NCBI's GenBank site (http://www.ncbi.nlm. nih.gov/Genbank/index.html).

Microarray data: The JI will not publish descriptive manuscripts that report microarray data, unless such information can be considered of unusual immunological significance and/or include functional experiments that provide novel insight into mechanism. As with other scientific approaches, current experimental, quantitation, verification, and statistical analyses are expected. Microarray experiments should be Minimum Information About a Microarray Experiment (MIAME) compliant (for guidelines see www.mged.org). Whereas limited online space may be available for supplemental tables associated with the manuscript, complete microarray data must be deposited in the appropriate public database; e.g., GEO (http://www.ncbi.nlm.nih.gov/geo/), ArrayExpress (http://www.ebi.ac.uk/ arrayexpress/), or CIBEX (http:// cibex.nig.ac.jp/index.jsp), and must be accessible without restriction from the date of publication. An entry name or accession number must be included in the paper before publication. The accession number should be accompanied by the website address of the databank.

#### STYLE GUIDE

General style conventions: In general, The JI follows Scientific Style and Format, The CSE Style Manual for Authors, Editors, and Publishers, seventh edition, published by the Council of Science Editors, Inc., in instances where journal style issues are not directly addressed.

Abbreviations for references: BIOSIS is the primary source for journal name abbreviations; *Index Medicus* is the secondary source.

**Nomenclature**: The most current links for nomenclature guidelines are posted online (http://www.jimmunol.org/misc/authorfulllength.shtml#style).

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Chemical names: The JI uses The Merck Index (http://library.dialog.com/bluesheets/html/bl0304.html) and the IUPAC-IUB Commission on Biochemical Nomenclature-Chemical Abstracts (http://www.chem.qmul.ac.uk/iupac/bibliog/white.html) as the primary references for proper spelling and style of chemical names.

**Chemokine/chemokine receptor nomenclature:** The systematic name for chemokines and chemokine receptors should be used. The original name may be given in parenthesis if desired. See *Cytokine* 21:48–49, 2003.

**Enzyme Nomenclature:** (http://www.chem.qmul.ac.uk/iubmb/enzyme/), is *The JI* source for style and spelling of enzyme names.

**Gene nomenclature:** The HUGO guidelines for gene nomenclature (http://www.gene.ucl.ac.uk/nomenclature/) may be used for naming human genes. Mouse Genome Informatics (http://www.informatics.jax.org/searches/marker\_form.shtml) is a reference source for naming mouse genes.

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**HLA nomenclature:** HLA nomenclature is updated periodically by the WHO Nomenclature Committee for Factors of the HLA System. A recent reference is *Hum. Immunol.* 64: 919–920, 2003. Annual comprehensive revisions are published in *Human Immunology*, usually in the spring.

#### STANDARD ABBREVIATIONS

The abbreviations listed here are used without definition in articles published in *The JI*. The form may be used for both singular and plural, or made plural with "s" at the author's option. The list of standard abbreviations is published in the first issue of each volume.

#### STANDARD ABBREVIATIONS

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A, angstrom

aa, amino acid (only with numbers)

Ab, antibody

ABTS, 2,2'-azinobis(3-ethylbenzthiazoline-6-sulfonic acid)

Ag, antigen

AIDS, acquired immunodeficiency syndrome

ANOVA, analysis of variance

AP-1, activator protein 1

APC, Ag-presenting cell

ATP, adenosine triphosphate (also ADP, AMP, CMP, CTP, GDP, GMP, GTP, ITP, NTP, TMP, UDP, and UTP)

AZT, 3'-azido-3-deoxythymidine

BALT, bronchus-associated lymphoid tissue

BAPTA-AM, 1,2-bis(2-aminophenoxy)ethane-*N*,*N*,*N'*,*N'* tetraacetic acid acetoxymethyl ester

BCR, B cell receptor

bp, base pair (only with numbers)

BrdU, 5-bromo-2'-deoxyuridine

BSA, bovine serum albumin

C, complement

C region, constant region of Ig

cAMP, cyclic AMP

CCL, CC chemokine ligand

CCR, CC chemokine receptor

CD40L, CD40 ligand

cDNA, complementary DNA

CDR, complementarity determining region

C/EBP, CCAAT/enhancer-binding protein

CFA, complete Freund's adjuvant

CFSE, 5-(and 6-)-carboxyfluorescein diacetate succinimidyl ester

CFU, colony-forming unit

cGMP, guanosine 3',5'-cyclic monophosphate

 $CHAPS,\ 3\hbox{-}[(3\hbox{-cholamidopropyl})dimethylammonio]\hbox{-}1\hbox{-propane-sulfonate}$ 

Ci, curie

CIITA, class II transactivator

CLIP, class II-associated invariant-chain peptide

CMV, cytomegalovirus

CNS, central nervous system

CoA, coenzyme A

Con A, concanavalin A

CpG, cytosine guanine dinucleotide

cpm, counts per minute

CREB, cAMP response binding protein

cRNA, complementary RNA

CSF, colony-stimulating factor

CTL, cytotoxic T lymphocyte

CTLA, cytolytic T lymphocyte-associated Ag

CXCL, CXC chemokine ligand

CXCR, CXC chemokine receptor

d, deoxy; distilled (as in dH<sub>2</sub>O)

D region, diversity region of Ig or T cell receptor for Ag

Da, dalton (only with numbers)

dATP, 2'-deoxyadenosine triphosphate

DEAE, diethylaminoethyl

df, degrees of freedom

DMEM, Dulbecco's modified Eagle's medium

DMSO, dimethylsulfoxide

DNA, deoxyribonucleic acid

DNase, deoxyribonuclease

DNP, dinitrophenyl

dNTP, 2'-deoxynucleoside 5'-triphosphate

dpm, disintegrations per minute

ds, double-stranded (as dsDNA)

DTT, dithiothreitol

E, erythrocyte

EBV, Epstein-Barr virus

EC<sub>50</sub>, 50% effective concentration

ECL, enhanced chemiluminescence

ED<sub>50</sub>, 50% effective dose

EDTA, ethylenediaminetetraacetic acid

EGTA, ethylene glycol-bis( $\beta$ -aminoethyl ester)-N,N,N',N'-tetraacetic acid

ELISA, enzyme-linked immunosorbent assay

ELISPOT, enzyme-linked immunospot

EMSA, electrophoretic mobility shift assay

ERK, extracellular signal-regulated kinase

E:T ratio, effector to target ratio

Fab, Ag-binding fragment

F-actin, filamentous actin

FACS, fluorescence-activated cell sorter

FAM, 6-carboxyfluorescein

FBS, fetal bovine serum

FcR, Fc receptors (e.g., Fc $\gamma$ RI)

FCS, fetal calf serum

FITC, fluorescein isothiocyanate

FLICE, Fas-associated death domain-like IL-1 $\beta$ -converting enzyme

FLIP, FLICE inhibitory protein

fMLP or FMLP, formyl-methionyl-leucyl-phenylalanine

Fura 2-AM, fura 2-acetoxymethyl ester

g, gram (only with numbers)

GALT, gut-associated lymphoid tissue

GAPDH or G3PDH, glyceraldehyde-3-phosphate

dehydrogenase

G-CSF, granulocyte CSF

GFP, green fluorescent protein

GM-CSF, granulocyte-macrophage CSF

gp, glycoprotein (e.g., gp100)

GPI, glycosylphosphatidylinositol

GST, glutathione *S*-transferase

h, hour (only with numbers)

H chain, heavy chain

H&E, hematoxylin and eosin

HBSS, Hanks' balanced salt solution

HEPES, N-2-hydroxyethylpiperazine-N'-2-ethanesulfonic acid

HIV, human immunodeficiency virus

HLA, human histocompatibility leukocyte Ag

HPLC, high performance liquid chromatography

HRP, horseradish peroxidase

HSV, herpes simplex virus

HUVEC, human umbilical vein endothelial cells

IC<sub>50</sub>, 50% inhibition/inhibitory concentration

ICAM, intercellular adhesion molecule

ICOS, inducible costimulator

Id, idiotype; idiotypic determinant

ID<sub>50</sub>, 50% infective dose or 50% inhibiting dose

IDO, indoleamine 2,3-dioxygenase

IFA, incomplete Freund's adjuvant

IFN, interferon (e.g., IFN-γ)

Ig, immunoglobulin

IgH, Ig heavy chain

IκB, inhibitory NF-κB IL, interleukin (e.g., IL-2)

i.m., intramuscular

IMDM, Iscove's modified Dulbecco's medium

IMEM, Iscove's minimal essential medium

i.p., intraperitoneal

ITAM, immunoreceptor tyrosine-based activation motif

ITIM, immunoreceptor tyrosine-based inhibitory motif

IU, international unit

i.v., intravenous
J region, joining region of Ig or T cell receptor for Ag

JAK or Jak, Janus kinase

JNK, c-Jun N-terminal kinase

kb, kilobase (only with numbers)

kbp, kilobase pair (only with numbers)

 $K_{\rm a}$ , association constant

 $K_{\rm d}$ , distribution coefficient; dissociation constant

 $K_{\rm D}$ , affinity constant

kDa, kilodalton (only with numbers)

L chain, light chain

LD<sub>50</sub>, 50% lethal dose

LFA, leukocyte (lymphocyte) function-associated Ag

LIF, leukemia inhibitory factor

LPS, lipopolysaccharide

LU, lytic unit

mAb, monoclonal Ab

MACS, magnetic-activated cell sorting

MALDI, matrix-assisted laser desorption ionization

MALDI-TOF, matrix-assisted laser desorption ionization-time

of flight

MALT, mucosa-associated lymphoid tissue

MAPK, mitogen-activated protein kinase

MCP-1, monocyte chemoattractant protein-1

M-CSF, macrophage CSF

2-ME, 2-mercaptoethanol

MEK, mitogen-activated protein kinase kinase

MEM, minimum essential medium

MES, 2-(N-morpholino)ethanesulfonic acid

mg, milligram (only with numbers)

MHC, major histocompatibility complex

min, minute (only with numbers)

MIP, macrophage-inflammatory protein

ml, milliliter (only with numbers)

MLC, mixed lymphocyte culture

MLR, mixed leukocyte reaction

mo, month(s) (only with numbers)

MOPS, 4-morpholinepropanesulfonic acid

 $M_{\rm r}$ , relative molecular mass mRNA, messenger RNA

MTT, 3-(4,5-dimethylthiazol-2-yl)-2,5-dimethyltetrazolium

bromide

 $\mu$ g, microgram (only with numbers)

 $\mu$ l, microliter (only with numbers)

m.w., molecular weight

MyD88, myeloid differentiating factor 88

n, number in study or group

NAD, nicotinamide adenine dinucleotide

NADH, reduced NAD

NaDodSO<sub>4</sub>, sodium dodecyl sulfate

NADP, NAD phosphate

NADPH, reduced NAD phosphate

NBT. nitroblue tetrazolium

ND, not determined

NDP, nucleoside 5'-diphosphate

NF, nuclear factor

NFAT or NF-AT, nuclear factor of activated T cells

NF-κB, nuclear factor κB

Ni-NTA, nickel-nitrilotriacetic acid

NK cell, natural killer cell

NMP, nucleoside 5'-monophosphate

NO, nitric oxide

NOD, nonobese diabetic

NS, not significant

nt, nucleotide (only with numbers)

OCT, octamer-binding factor

OD, optical density

OVA, ovalbumin

p, probability

PAGE, polyacrylamide gel electrophoresis

PBL, peripheral blood lymphocyte

PBMC, peripheral blood mononuclear cell

PBS, phosphate-buffered saline PCR, polymerase chain reaction

PE, phycoerythrin

PECAM-1, platelet endothelial cell adhesion molecule-1

PerCP, peridinin chlorophyll protein

PFU, plaque-forming unit

PG, prostaglandin

PHA, phytohemagglutinin

PI3K, phosphatidylinositol 3-kinase

PIPES, piperazine-N,N'-bis(2-ethane sulfonic acid)

PMA, phorbol myristate acetate

PMSF, phenylmethylsulfonyl fluoride

PWM, pokeweed mitogen

r, recombinant (e.g., rIFN- $\gamma$ )

R, receptor (e.g., IL-2R)

RACE, rapid amplification of cDNA end

RAG, recombination-activating gene

RANTES, regulated upon activation, normal T cell expressed

and secreted

RBC, red blood cell

RFLP, restriction fragment length polymorphism

RIA, radioimmunoassay

RNA, ribonucleic acid

RNase, ribonuclease

rpm, revolutions per minute rRNA, ribosomal RNA

RT-PCR, reverse transcriptase polymerase chain reaction

s, second (use only with numbers)

s.c., subcutaneous

SCID, severe combined immunodeficiency

SD, standard deviation SDS, sodium dodecyl sulfate

SE, standard error

SEM, standard error of the mean

SHIP, src homology 2-containing inositol 5'-phosphatase

SIV, simian immunodeficiency virus

sp. act., specific activity

SRBC, sheep red blood cells

ss, single-stranded (e.g., ssDNA)

SSC, standard saline citrate

STAT, signal transducer and activator of transcription

SV40, simian virus 40  $t_{1/2}$ , half-life, half-time

TAMRA, 5-(and 6)-carboxytetramethylrhodamine

TAP, transporter associated with Ag processing

Tat, terminal deoxynucleotidyltransferase

TBS, Tris-buffered saline

TBST, TBS with Tween 20

TCA, trichloroacetic acid

TCR, T cell receptor for Ag

TdR, thymidine deoxyribose (also UdR, AdR)

TdT, terminal deoxynucleotidyltransferase

TGF, transforming growth factor

Th cell, T helper cell

TLC, thin layer chromatography

TLR, Toll-like receptor

TNF, tumor necrosis factor

TNP, trinitrophenyl

TRAIL, TNF-related apoptosis-inducing ligand

Tris, tris(hydroxymethyl)aminomethane

tRNA, transfer RNA

TUNEL, Tdt-mediated dUTP nick end labeling

U, unit (only with numbers)

UV, ultraviolet

v/v, volume to volume ratio (%)

V region, variable region of Ig

VCAM, vascular cell adhesion molecule

V(D)J, variable diversity joining

VLA, very late activation Ag

W, watt (only with numbers)

wk, week (only with numbers)

xid, X-linked immunodeficiency

Zap70, ζ-associated protein 70 (or ζ-chain-associated protein

#### **Keywords**

Animals Human Rodent Other Animals

Cells B Cells Dendritic Cells Endothelial Cells Eosinophils Mast Cells/Basophils

Monocytes/Macrophages Natural Killer Cells

Neutrophils Stem Cells Stromal Cells

T Cells, Cytotoxic Th1/Th2 Cells

Diseases

Autoimmunity Diabetes EAE/MS

Endotoxin Shock Graft Versus Host Disease Immunodeficiency Diseases Rheumatoid Arthritis Systemic Lupus Erythematosus

Infections AIDS Bacterial

Fungal Parasitic-Helminth Parasitic-Protozoan

Molecules AcutePhase Reactants Adhesion Molecules

Antibodies Antigens/Peptides/Epitopes

Autoantibodies
Cell Surface Molecules

Chemokines Complement Cytokine Receptors

Cytokines Fc Receptors Lipid Mediators Lipopolysaccharide MHC

Nitric Oxide Protein Kinases/Phosphatases

Superantigens T Cell Receptors Transcription Factors

Processes

Allergy Antigen Presentation/Processing

Apoptosis Cell Activation Cell Differentiation Cell Proliferation Cell Trafficking

Chemotaxis Comparative Immunology/Evolution

Costimulation Cytotoxicity Gene Rearrangement

Neuroimmunology Phagocytosis Repertoire Development Reproductive Immunology Signal Transduction Tolerance/Suppression/Anergy Transplantation Tumor Immunity Vaccination

**Techniques** 

Gene Regulation

Hematopoiesis Inflammation

Memory

Gene Therapy Molecular Biology Transgenic/Knockout Mice

Tissues Lung Mucosa

Skin Spleen & Lymph Nodes Thymus

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The American Association of Immunologists (AAI) is a professional organization whose members have a strong interest in, and have made substantial contributions to, the science of immunology. AAI is a member of the Federation of American Societies for Experimental Biology (FASEB) and is responsible for the publication of *The Journal of Immunology*. To be eligible for election to membership in the AAI, a candidate must meet **one** of the following criteria:

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