INSTRUCTIONS FOR AUTHORS

Please note important changes:
p. 473—new manuscript processing fee
p. 474—new style for References section

CANCER Research is the official organ of the American Association for Cancer Research, Inc., and is devoted to the publication of significant, original research in all the subfields of cancer research, including: biochemistry and physiology; chemical and physical carcinogenesis and mutagenesis; clinical investigations; endocrinology; epidemiology and biostatistics; immunology; molecular and cell biology; preclinical pharmacology and experimental therapeutics; radiobiology; and virology. Clinical investigations and epidemiological studies are published in a separate section from papers in the basic sciences.

Only those papers that report results of novel and timely studies and that meet high standards of scientific quality will be accepted. Papers are subjected to stringent review and are published within about three months of acceptance.

CATEGORIES OF PUBLICATION

The following types of manuscripts are considered for publication. (1) Papers containing results of original experimental, clinical, or statistical studies that are sufficiently well documented to be acceptable to the critical reader. (2) Concise reviews on subjects of importance to cancer researchers. Authors of unsolicited reviews should submit an outline of the proposed article for approval by the Editorial Board. If submission of the complete article is encouraged, the review will be given particularly stringent editorial evaluation before acceptance. (3) Perspectives in Cancer Research, which are invited articles analyzing either very active or undeveloped areas of research and presenting fresh insights and personal viewpoints on where research in that area may or should be heading. (4) Letters to the Editor which deal with issues of importance to cancer researchers. If experimental data are included, these should be kept to the minimum required for adequate understanding. Included under this category are commentaries on manuscripts published in the Journal. If accepted, replies by the authors are invited for simultaneous publication. (5) Brief reports of meetings, symposia, and conferences related to cancer research. (6) Proceedings of symposia, published as external supplements to the Journal (Cancer Research Supplements), the full expenses of which are assumed by the sponsoring agency. These proceedings are published at the discretion of the Editor and do not undergo the usual review process. (7) Brief announcements of scientific meetings of interest to readers, of courses in cancer-related biomedical science, and of the availability of fellowships and scholarships. These should be submitted at least 3 months prior to the expected month of issue. (8) Brief announcements of recent deaths among distinguished contributors to the field of cancer research.

EDITORIAL POLICY

When a manuscript is received for consideration, the Editors assume that no similar paper, other than an abstract or preliminary report, has been or will be submitted for publication elsewhere. The Editors assume that all authors listed on a manuscript have agreed to its submission. Submission of a manuscript implies acceptance of the strict policy of the journal that under no circumstances will the identities of the editors and reviewers be revealed.

Typically, the scientific review of a manuscript is handled by an Associate Editor who selects two investigators in the field as referees. The reviewers' critiques and the Associate Editor's recommended decision are then approved by the Editor. Editorial decisions, forwarded from the Editorial Office to authors, are rendered as promptly as possible, consistent with thoroughness of review; the review process usually takes an average of 8-10 weeks. If there is a marked discrepancy in the opinions of the reviewers, the Editor may send the paper to other reviewers for additional comments. In this case, more time may be needed to finalize the review process.

The Editorial Office cannot accept collect telephone calls from authors.

SUBMISSION OF MANUSCRIPTS

Contributions must be submitted in quadruplicate (with at least two sets of original illustrations) to Dr. Peter N. Magee, Editor, Cancer Research Editorial Office, Fels Research Institute, Temple University School of Medicine, Philadelphia, Pa. 19140 (Telephone: 215-221-4720). Papers should be submitted only by an author, preferably the senior author, who should request in the covering letter that the paper be considered for publication in Cancer Research. The exact address to which all related correspondence should be sent and a telephone number at which the author can be reached should also be given. Contributors should indicate in the covering letter if their papers would be appropriate for the Clinical and Epidemiological Investigations section; however, the final decision as to placement within the journal will be made at the discretion of the Editors. The original plus a photocopy of this letter are required.

If the manuscript contains any quoted information conveyed by either personal communication or release of unpublished experimental data, the covering letter should state specifically that authorization to use this material has been given.

A manuscript processing fee of $75 is assessed for each manuscript to cover the cost of editorial review. Payment of this processing fee may accompany the manuscript. An invoice will be mailed with the acknowledgment of receipt of the manuscript at the Editorial Office if payment has not already been made. As a courtesy to authors editorial review will not be delayed for receipt of payment. Authors are requested to fulfill their own institutional obligations with respect to purchase orders and call numbers so that payment of the manuscript handling charge can be expedited.

Revised manuscripts should be submitted in quadruplicate, with two sets of original illustrations. A covering letter in duplicate must accompany all revised manuscripts and indicate clearly what alterations have been made in response to the reviewers' comments. Satisfactory reasons should be given for noncompliance with any of the recommendations for revision. Revised manuscripts may undergo another review by an Associate Editor and/or referees, particularly if the original submission required extensive changes.

If a new author has been added or an author has been deleted since the original submission, it is the responsibility of the corresponding author to ensure that the authors involved are aware of and agree to the changes in authorship. Cancer Research accepts no responsibility for such changes.

FORMAT AND STYLE

Papers should conform strictly to Journal style. A recent issue of Cancer Research will provide authors with assistance in the proper arrangement of papers. Manuscripts are to be written in clear, grammatical English. Papers that are not in good idiomatic English will be returned to the author without review. Laboratory slang as well as terminology and abbreviations not consistent with internationally accepted guidelines should be avoided.


Data must be presented concisely. Large masses of data of peripheral significance to the main thesis of the investigation will not be published in Cancer Research but may be deposited with the National Auxiliary Publications Service, c/o Microfiche Publications, P.O. Box 3513, Grand Central Station, New York, N. Y. 10017. The manuscript should contain a footnote that indicates how this ancillary material can be obtained. Such data should be submitted for review along with the manuscript.

The manuscript should be typed on 21.6- x 28-cm (81/2- x 11-inch)
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paper with double spacing throughout, allowing for ample margins. Manuscripts with typing on both sides of the page will be returned to the authors. Consecutive numbering of all pages is requested, with the title page and page 1. The typescript should be arranged in the following order: (a) title, (b) author(s) and complete name(s) and location(s) of institution(s) or laboratory(ies), (c) running title, (d) footnotes, (e) text, (f) tables, (g) legends for all illustrations (charts and figures), (h) illustrations, and (i) other material. Numbered and lettered sections in the text should be avoided. The appropriate location for each table and illustration should be indicated by marginal notes. Simple chemical formulas or mathematical equations should be presented in a form that allows their reproduction in single horizontal lines of type; more complicated mathematical formulas or chemical structures difficult to set in type should be provided in the form of India ink drawings or glossy photographs for camera-ready reproduction.

Title. Titles should be brief but informative, and limited if possible to about 100 characters. It is important for literature retrieval to include in the title the key words necessary to identify the nature of the subject matter, including, if applicable, the species of animal on which the work is done. Use of expressions such as “Studies on...” or “Observations of...” should be avoided, since they are not sufficiently informative. Chemical formulas or abbreviations should not be used. Titles in the form of declarative or interrogative sentences are not acceptable. Also, do not use typographic numerals to indicate that the paper is one in a series (see section below on Footnotes).

Authors and Their Affiliations. Authors are urged to include their full names, complete with first and middle names or initials. Confusion often arises in the literature when authors are identified by surname and initials only. Authors’ academic degrees should not be included. The full names of institutions and subsidiary laboratories should be given, together with a useful address (including postal number). If several authors and institutions are listed on a paper, it should be clearly indicated with which department and institution each author is affiliated.

Running Title. A brief running title should be provided, not to exceed 50 characters. Running titles in the form of declarative or interrogative sentences are not acceptable.

Footnotes. Longish footnotes are discouraged since the same information can in most instances be presented more effectively in the text. Footnotes to the title page and text are to be designated consecutively with superscript Arabic numerals. A footnote to the title should contain information on financial support, including the source(s) and number(s) of the grant(s). If the paper is one of a series, a footnote to this effect may be included. Authors should also include a footnote designating to whom reprint requests should be addressed. An all-inclusive abbreviation footnote should contain a definition for every nonstandard abbreviation used in the paper.

For footnotes to tables, see section on Tables below.

Abstract. The abstract, to appear at the beginning of the paper, should be a concise summary of the results. As abstracts are often copied directly by the secondary services, they should recapitulate in abbreviated form the purpose of the study and the experimental technique, results, and interpretations of the data. Data such as the number of test subjects and controls, strains of animals or viruses, drug dosages and routes of administration, tumor yields and latent periods, length of observation period, and magnitude of activity should be included. Vague, general statements such as “The significance of the results is discussed,” or “Some physical properties were studied,” are uninformative and not acceptable. All important terms relevant to the content of the paper should be incorporated into the abstract to assist indexers in the derivation of key words. Abbreviations should be kept to an absolute minimum; however, if they are needed, they must be properly identified so as to make the abstract independent of the text. Authors may wish to keep in mind that “Medline,” a computerized monthly bibliography prepared by the National Library of Medicine, includes only those abstracts that contain fewer than 200 words; with very few exceptions, longer abstracts are not accessible through that service.

Introduction. It is not necessary to include all of the background in the introduction. Brief reference to the most pertinent papers generally suffices to acquaint the reader with the findings of others in the field and with the problem or question to which the author’s particular investigation is being addressed.

Materials and Methods. Explanation of the experimental methods should be brief but adequate for repetition by qualified investigators.

Procedures that have been published previously should not be described in detail but merely cited in appropriate references. Only new and significant modifications of previously published procedures need complete exposition. The sources of special chemicals or preparations used should be given along with their locations (city and state (country, if foreign)).

This Journal endorses the principles embodied in the Declaration of Helsinki and expects that all investigations involving humans will have been performed in accordance with these principles. For animal experimentation reported in this Journal, it is expected that the Guiding Principles in the Care and Use of Animals approved by the American Physiological Society will have been observed.

Results. This section should include a concise textual description of the data presented in tables, charts, and figures. Excessive elaboration of data already given in tables and illustrations should be avoided. The Results and Discussion sections may be combined if, by so doing, space is saved or the logical sequence of the material is improved.

Discussion. In this section, the data should be interpreted concisely without repeating material already presented in the Results section. Speculation is permissible, but it must be well founded.

References. Number references in the order of their first mention in the text; cite only the number assigned to the reference. [Please note that this is a change from our previous style of arranging the references section in alphabetical order.] References should be typed in double-spaced form to ensure accurate copy editing. The bibliography should be limited to only those citations essential to the author’s presentation. When comprehensive review articles are available, they are preferred to many separate references.

Before submission of the paper authors should verify the accuracy of all references and should check that all references have been cited in the text. Examples of the two most common types of Journal references are:


For tables, include the title, page number, inclusive pages, and year should be given. Serial compendia, such as Advances in Cancer Research and the Annual Review of Biochemistry, which appear annually in numbered sequence, should be cited as journals rather than books, thus omitting the names of publishers and editors. Biological Abstracts should be consulted for abbreviations of journals and serials.

Books and chapter citations. Citation of a specific chapter or article in a book should carry the author(s) of the chapter, its title, editor(s) of the book, title of the book, volume, inclusive pages of the chapter, location and name of the publisher, and year. For references to complete books, give all of the above information that is pertinent.

Papers in press and unpublished material. Papers in press may be listed among the references with the journal name and tentative year of publication. Papers in preparation or submitted for publication should be cited in a footnote, not in the References section. This rule also applies to unpublished data or personal communications. The names of all authors should be given, along with manuscript titles if possible.

Addenda. Data acquired after acceptance of the paper, by the authors themselves or by others, cannot be added to the text. An addendum may be added in proof upon approval by the Editor. Addenda should be kept extremely brief. The full expense of printing an addendum will be charged to the author.

Tables. Tables should be constructed so that when typeset, they will fit within a single Journal column (8.9 cm or 3½ inches). Tabular material should not duplicate data already presented in the charts. Unnecessary columns of data that can easily be derived from the rest of the results in the table should not be included. Large groups of individual values should be avoided; instead, these should be averaged and an appropriate designation of the dispersion such as standard deviation or standard error included.

Authors are obliged to indicate the significance of their observations by appropriate statistical analysis.

Every table must have a descriptive title and an explanatory paragraph.
that clearly gives the experimental details for understanding by the reader without reference to the text. Each column must carry an appropriate heading and, if numerical measurements are given, these units should be added to the column heading. Tables should be numbered with Arabic numerals and table footnotes should be indicated with superscript italic letters (a, b, c, etc.).

All units of measurement and concentration should be clearly designated. Exponential terminology is discouraged (the term m is preferable to $10^{-3}$). If exponents are absolutely unavoidable in column headings, the quantity expressed should be preceded, not followed, by the power of 10 by which its value has been multiplied, i.e., $10^{-3}$ x concentration (m). This will prevent confusion as to whether the quantity should be multiplied or divided to obtain the correct value.

Charts. Line-out illustrations (graphs and drawings) are to be designated charts. Flow diagrams and complex biochemical structures should be prepared professionally (not simply typewritten) and considered charts.

Graphs should be used sparingly and only when a salient point needs illustration. Straight-line functions such as relationships between concentration and absorbance, or Lineweaver-Burk plots when these are linear, should instead be described in a few lines in the text. To conserve space those curves that may appropriately appear together should be included in a single chart.

The use of exponents for labeling coordinates in charts is considered ambiguous and should, if at all possible, be avoided. If exponents must be used, the quantity expressed should be preceded by the power of 10 by which its value has been multiplied, i.e., $10^{-3}$ x concentration (m). The form "Concentration ($m x 10^{-3}$)" is not acceptable. If powers of 10 are used, the legend should designate how the quantity is to be calculated (whether multiplied or divided) to give the correct value.

Preparation of charts. Charts must be drawn with professional instruments and may be on Bristol board, tracing paper or cloth, or coordinate paper printed in light blue. Charts should not be mounted on heavy cardboard. Clear, glossy prints are acceptable in lieu of original drawings, provided that all parts of the chart are in focus. X-ray films or Polaroid photographs are not acceptable. If original drawings are submitted, they should not be larger than 21.6 x 28 cm ($8^{1/2} x 11$ inches).

Except for especially complicated drawings showing large amounts of data, all charts are published at one-column width (8.9 cm or 3½ inches) or less. It is recommended that charts be submitted in one-column size. If larger charts are submitted, it is the responsibility of the author to see that the abscissas, ordinates, lines, and especially the symbols are sufficiently large to permit reduction. When the charts are reduced to the size of a single column, the letters and numbers must be at least 1.5 mm high and the smallest part of the illustration must be discernible or the charts will be returned to the author for correction. In original charts, this can be insured by having the minimum height for lower-case letters 5 mm; numerals and upper-case letters 6 mm; and symbols within the drawings 5 mm. The thickness of ruled lines on charts is also vital for clear presentation of the data. Size recommendations for lines are as follows: #1 Leroy for graph grids, bonds, and arrows; #2 Leroy for graph borders or reference lines; and #5 Leroy for graph curves or emphasis lines.

Points of observations should be denoted with different symbols rather than with different types of lines; their significance can be explained directly in the body of the chart or in the legend. Only those common symbols for which the printer has type ($\times$, O, ●, ○, □, ■, ■, △, □, ○) should be used.

Charts should be ruled off close to the area occupied by the curve, and abscissas and ordinates should be clearly marked with appropriate units. Explanations of the coordinates should not extend beyond the respective lines. Do not box-in charts with top and right-hand frame lines unless these are essential for reference. If a chart contains a left- and a right-hand ordinate, explanation of the left ordinate should read in the upward direction and that of the right ordinate should read downward. Titles printed outside the confines of the charts waste space; all of this information can easily be included in the legend.

Each chart should be labeled in pencil on an adhesive label on the reverse side with Arabic numerals and the first author's name.

Chart legends. Legends are required for all charts. They should briefly describe the data shown; details in the text need not be repeated. Each legend should adequately identify all units, abbreviations, mathematical expressions, abscissas, ordinates, and symbols.

Figures. Because half-tone illustrations (photomicrographs and photographs) are expensive to reproduce, only those photographs that are absolutely essential to the clarity of the presentation can be accepted. It is often difficult for the printer to judge accurately the amount of reduction possible with electrophoretic patterns. Therefore, this material should be submitted in reduced form ready for photoreproduction.

Color photographs are discouraged and will be published only if the Editors deem them indispensable. The complete expense of reproducing such photographs will be charged to the author. The author is also responsible for submitting prints that are of sufficient quality to permit accurate reproduction, and for approving the final color proof. If mounted, color photographs must be on a flexible backing. Cancer Research assumes no responsibility for the quality of the photographs as it appears in the journal. Current estimates for color reproduction can be obtained from the Editorial Office.

Preparation of plates. Photographs should be mounted on "plates" of white cardboard; the overall dimensions of photographs on a given plate should not exceed 18.4 x 22.4 cm (7½ x 9 inches). All photographs should be correctly exposed, sharply focused, and submitted on glossy white paper.

Karyotypes should be presented in the form of cardboard plates onto which chromosome sections from an original photomicrograph are pasted. This "original" is needed for clear Journal reproduction. The back of the plate should indicate how much it can be reduced in size if published. For review purposes, two additional glossy photographs are requested.

Considerable space may be saved by cropping figures so that 4 to 6 photographs can be presented on one plate. Plates with single photographs are not acceptable and will be returned for cropping or reduction unless the authors can justify their necessity. The contrast among photographs on a plate should be consistent for better clarity of reproduction. Photographs should be buttressed together; tooling (thin white lines) between the photographs will be added by the printer.

Figure numbers, in Arabic numerals, should appear in India ink directly on the photographs and, if possible, should be in the lower right-hand corner of each photograph. These numbers should be no larger than ⅛ inch. Wax-banded lettering such as PRES-TYPE or LETRASET is discouraged because of its tendency to crumble and adhere to vinyl overlays. Photomicrographs should have internal scale markers. Symbols, arrows, or letters used in photomicrographs should contrast with the background.

Tissue overlays on plates are a necessary protection for figures. Indicate on overlays the important areas of the photographs that must be reproduced with greatest fidelity. The top of the figure and the first author's name should be given in pencil on an adhesive label on the reverse side of each plate.

Figure legends. An appropriate legend for each figure, including stains and magnifications where applicable, is required. Any abbreviations or reference points on a figure should be explained in the legend. All attempts will be made to place legends under the plates to which they refer. To facilitate proper layout, authors should keep their plates to 7¼ x 8½ inches.

ABBREVIATIONS

Abbreviations are in general a hindrance to readers in fields other than that of the author(s), to abstractors, and to scientists in foreign countries. Authors should limit their use to an absolute minimum. Single words should not be abbreviated, e.g., daunomycin, folate, vincristine. Abbreviations are not to be used in titles, but running titles may carry abbreviations for purposes of brevity. Abstracts may contain abbreviations for terms mentioned many times in that section but their identification is mandatory.

Authors should follow the recommendations of the IUPAC-IUB Commission on Biochemical Nomenclature (see section below on Terminology). All nonstandard abbreviations should be identified in an inclusive abbreviation footnote to the first such abbreviation after the Abstract. Abbreviations that form recognizable words, such as EAT and MOPS, are discouraged.

Standard Abbreviations. Authors may use, without definition, the abbreviations in the following lists.
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NAD+, NADH
nicotinamide adenine dinucleotide and its reduced form

NADP+, NADPH
nicotinamide adenine dinucleotide phosphate and its reduced form

(DPN*, TPN*, and their reduced forms are not acceptable.)

CoA, acyl-CoA
coenzyme A and its acyl derivatives (e.g., acetyl)

AMP, GMP, IMP, UMP
the 5'-phosphates of ribosyladenine, -guanine, -inosine, -uracil, -cytosine, and -thymine

ADP, etc.
the 5'(pyro)diphosphates of adenosine, etc.

ATP, etc.
the 5'(pyro)triphosphates of adenosine, etc.

dAMP, dGMP, dIMP
the 5'-phosphates of 2'-deoxyribosyladenine, etc.

RNA, DNA
ribonucleic acid, deoxyribonucleic acid

RNase, DNase
ribonuclease, deoxyribonuclease

mRNA
messenger RNA

nRNA
nuclear RNA

rRNA
ribosomal RNA

tRNA
transfer RNA (sRNA is not recommended for RNA preparations that accept amino acids.)

P, PP
orthophosphate, pyrophosphate

Tris
tris(hydroxymethyl)methylamine

EDTA
ethylenediaminetetraacetate

POPOP
1,4-bis(2-(5-phenyloxazolyl))benzene

DEAE, TEAE
diethylaminoethyl, triethylaminoethyl

UV, IR
ultraviolet, infrared

RBC, WBC
red blood cell(s), white blood cell(s)

Units of Concentration
molar (moles/liter)

millimolar (millimoles/liter)

micromolar (micromoles/"

nanomolar

picomolar

The expression mg % should be avoided; weight concentrations should be given as g per ml, g per 100 ml, g per liter, etc.

Units of Length, Area, Volume, Mass, Time

The abbreviations below are correct for both singular and plural forms of each term.

meter

centimeter

square centimeter

millimeter

micrometer (not micron)

nanometer (not millimicron)

picometer (not micromicron)

Angstrom (0.1 nm)

liter

milliliter

microliter

gram

milligram

microgram

kilogram

hour

minute

second

Physical and Chemical Units

Rt

acceleration of gravity

g

sedimentation coefficient

s

sedimentation coefficient in water at 20°

K

degree Celsius (Centigrade)

°C

degree Fahrenheit

°F

diffusion coefficient

D

equilibrium constant

K

inhibition constant

K_i

Michaelis constant

K_m

maximum velocity

V_{max}

Others

mole

mol

curie

Ci

equivalent

eq

counts per minute

cpm

disintegrations per minute

dpm

revolutions per minute

rpm

volt

V

Svedberg unit

S

absorbance

A (not O.D.)

probability

P

roentgen

R

standard deviation

SD

standard error of the mean

SE

logarithm (Briggsian)

log

logarithm (natural)

ln

entropy

S

molecular weight

M

In chemical compounds

ortho

o

meta

m

para

p

secondary

sec-

tertiary

tert-

Routes of administration

intramuscular

i.m.

intraperitoneal

i.p.

intravenous

i.v.

oral

p.o.

subcutaneous

s.c.

TERMINOLOGY

Approved terms and abbreviations for chemical substances have been collected in Biochemical Nomenclature and Related Documents, International Union of Biochemistry, Third Edition, 1978. This volume is available from: The Biochemical Society, 7 Warwick Court, London WC1R 5DP, United Kingdom. Included are all recommendations issued by the IUPAC-IUB Commission on Biochemical Nomenclature in the following areas: general abbreviations and symbols; abbreviations and symbols for chemical names of special interest in biological chemistry; stereochemistry; natural products and related compounds; isotopically labeled and modified compounds; biochemical equilibrium data; α-amino acids; symbols for amino-acid derivatives and peptides; synthetic modifications of natural peptides; synthetic polypeptides or polymerized amino acids; amino-acid sequences; conformation of polypeptide chains; peptide hormones; human immunoglobulins, multiple forms of enzymes; nucleic acids, polynucleotides, and their constituents; lipids; steroids,
quinones with isoprenoid side chains; carotenoids; tocopherols and related compounds; carbohydrates; cyclitols; phosphorus-containing compounds of importance in biochemistry; folic acids and related compounds; vitamins B-6 and related compounds; cornoids.

**Isotopically Labeled Compounds.** A radioactive nuclide is indicated by its mass number as a superscript to the left of the symbol (³²P); when written out, it should correspond to the spoken word (phosphorus-32).

In an isotopically labeled compound, the isotopic prefix should be placed in *square brackets and immediately precede the name (word) to which it refers*, as in [³¹C]thymidine, [α-³¹C]leucine, L-[(methyl-³¹C)me-thionine, [³⁷H]-3-hydroxykynurenine. When more than one position in a substance is labeled by means of the same isotope and the positions are not indicated, the number of labeled atoms is added as a subscript to the right of the element, as in [¹⁴C]glycine. The symbol U indicates uniform labeling and G, general labeling, e.g., [U-¹⁴C]glucose (where the [¹⁴C] is uniformly distributed among all six positions) and [G-¹⁴C]glucose (where the [¹⁴C] is distributed among all six positions, but not necessarily uniformly).

The isotopic prefix precedes that part of the name to which it refers, as in sodium [¹⁴C]formate, iodio[¹⁴C]acetic acid, 1-aminio[¹⁴C]methylcyclopentanol, α-naphthyl[¹⁴C]acetic acid, 2-acetamido-7-[(¹¹C)]iodofluorene, fructose 1,6-[³²P]bisphosphate, 17β-[³⁵S]estradiol. Terms such as [³¹H]-labeled albumin should not be contracted to [³¹H]albumin (since native albumin does not contain iodine), and [¹⁴C]-labeled amino acids should similarly not be written as [¹⁴C]amino acids (since there is no carbon in the amino group).

When isotopes of more than one element are introduced, their symbols should be arranged in alphabetical order, e.g., [³-¹⁴C; 2-³D; ¹⁵N]serine. Deuterium and tritium may be designated as [²H] and [³H] or as D and T, respectively.

When not sufficiently distinguished by the foregoing means, the positions of isotopic labeling are indicated by Arabic numerals, Greek letters, or prefixes in italics, as appropriate; these are to be placed within square brackets to appear before the symbol of the element concerned and are attached to it by a hyphen. Examples of this style are [¹-¹⁴C]alanine, L-[²-¹⁴C]leucine or L-[(methyl-¹⁴C)leucine, (carboxy-¹⁴C)leucine, [2,3-¹⁴C]maleic anhydride, [3,4-¹⁴C, ³⁵S]methionine, L-[(methyl-¹⁴C)leucine. The symbol indicating configuration always precedes the bracketed isotope, and a hyphen is used to separate it from the brackets, e.g., d-[¹⁴C]-glucose; L-[¹-¹⁴C]leucine.

The same rules apply when the labeled compound is designated by a standard abbreviation or symbol other than the atomic symbol, e.g., [α-³²P]ATP, [²⁵P]CMP, or [³¹H]dU. The square brackets are not to be used, however, with atomic symbols, or when the isotopic symbol is attached to a word that is not a specific chemical name, abbreviation, or symbol. Proper usage here is: [¹⁴C]O₂, [³²H]O₂, H₂[³⁵S]O₄, [³⁵P]H-labeled, [³⁷H]-lignans, [⁴⁻³⁷C]-steroids.

**Enzymes.** Authors should use the Recommended Name given by the Nomenclature Committee of the International Union of Biochemistry on the Nomenclature and Classification of Enzymes (Academic Press, Inc., New York, 1979) and Supplements 1 to 3: Corrections and additions. Eur. J. Biochem., 104: 1–4, 1980; 118: 423–435, 1981; 125: 1–13, 1982. In some cases the Systematic Name or the reaction catalyzed should also be included. It is strongly recommended that the Enzyme Commission number be stated at first mention.


**Histones.** Histone nomenclature should conform to the following system proposed at a Ciba conference held on April 4–5, 1974, in London: the six histone fractions are to be labeled H1, H1°, H2A, H2B, H3, and H4, rather than F1, F1°, F2a2, F2b, F3, and F2a1, respectively.

**Interferon Assays.** When reporting the calibration of interferon assays, authors should state the name, identifying number, and assigned potency of the international standard used to calibrate their assay, along with the observed geometric mean titer of the standard, the standard deviation of that value, the number of titrations performed to obtain that value, and the technical details of the assay.


**Outbred Animal Stocks.** Nomenclature for outbred laboratory animals should conform to that recommended by the Committee on Nomenclature, Institute of Laboratory Animal Resources: "A Nomenclature System for Outbred Animals," Lab. Animal Care, 20: 903–906, 1970.

**Drugs.** Generic names of drugs are preferred; a proprietary name may be used only after the first mention of the generic name and should be avoided in titles unless both names can easily be listed. If a foreign proprietary name is used, the name of the comparable U. S. product should not necessarily uniformly.

Authors should refer to the formally adopted generic names listed in *USAN and the USP Dictionary of Drug Names* (1984). In addition, the Council on Drugs reports in *The Journal (New Names)* drug names adopted by the USAN (United States Adopted Names) Council. These monographs include both the generic and proprietary names for the newest drugs, usually prior to their publication elsewhere.

**Tumors.** Tumors used in experimental investigations should be clearly described and identified in acceptable terminology. If these tumors are well known and have been identified in previous publications, extended descriptions and photomicrographs are unnecessary.

**General.** The composition of all solutions and buffers should be specified in sufficient detail so that the concentration of each component can be determined. The word "saline" should be replaced by "NaCl solution," along with the exact concentration. Inexact terms such as "physiological saline" or "phosphate-buffered saline" are not permitted; exact contents and concentrations should be given.

Decimals are preferred to factions; the form 0.01, not .01, is required in text, tables, and charts.

**Ionic charge** should be designated by a superscript immediately following the chemical symbol, e.g., Mg²⁺, S⁻.

**Advice on biochemical nomenclature** is readily available from Dr. Waldo E. Cohn, Director, Office of Biochemical Nomenclature, Biology Division, Oak Ridge National Laboratory, Box Y, Oak Ridge, Tennessee 37830; (615) 574-0808.

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