

CLINICAL SCIENCE

Guidance for Authors 1998

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I. POLICY OF THE JOURNAL

I.1. Scope

Clinical Science publishes papers in the field of clinical investigation, provided they are of a suitable standard and contribute to the advancement of knowledge in this field. The term 'clinical investigation' is used in its broadest sense to include studies in animals and the whole range of biochemical, physiological, immunological and other approaches that may have relevance to disease in man. Studies which are confined to normal subjects, or animals, or are purely methodological in nature may be acceptable. The material presented should permit conclusions to be drawn and should not be only of a preliminary nature. The journal publishes five types of manuscript, namely invited Editorial Reviews, Full Papers, Rapid Communications, Correspondence and invited Comments. In addition, *Clinical Science* publishes abstracts of the proceedings of the Medical Research Society (as Supplements) and also the Bayer Lecture.

I.2. Availability on the World Wide Web (WWW)

Abstracts of all articles are available on the journal's home page on the WWW (<http://cs.portlandpress.co.uk>). The journal will become fully online during 1998.

I.3. The editorial process

Membership of the Editorial Board covers as wide a range of interests as possible.

A submitted paper is considered by an appropriate editor together with (usually) two Referees from outside the membership of the Board. The Editor returns it with a recommendation to the Editor in Chief or Regional Editor, who then writes formally to the authors. The ultimate responsibility of acceptance for publication lies with the Editor in Chief.

Authors may suggest potential referees for their papers in the submission letter. The journal is under no obligation to follow such suggestions, but, if it does so, only one of the referees will be chosen from the authors' nominations, as the other referee will be selected independently.

I.4. Ethics of investigations

(a) Human subjects. Authors must state in the text of their paper that the research has been carried out in accordance with the Declaration of Helsinki (1989) of the World Medical Association, and has been approved by the Ethics Committee of the institution in which the work was performed. Consent **must** be obtained from each patient or subject after full explanation of the purpose, nature and risk of all procedures used, and the fact that such

consent has been given should be recorded in the paper.

(b) **Animals.** Care must always be taken to ensure that experimental animals do not suffer unnecessarily. Authors must state in the text the anaesthetic procedures used in full, and all precautions they took to ensure that the animals did not suffer unduly during and after the experimental procedure. Authors must confirm that the work was undertaken as required by the appropriate national legislation governing the use of animals, or, in the absence of such legislation, that the experimental procedures were carried out in accordance with the United States NIH guidelines [Guide for the care and use of laboratory animals, DHEW Publication no. (NIH) 85-23, Bethesda, MD: Office of Science and Health Reports, DRR/NIH, 1985].

The Editorial Board will not accept papers where the ethical aspects are, in the Board's opinion, open to doubt.

1.5. Originality of papers

Submission of a paper to *Clinical Science* implies that it has been approved by all the named authors, that all persons entitled to authorship have been so named, that it reports unpublished work that is not under consideration for publication elsewhere, that proper reference is made to the preceding literature, and that if the paper is accepted for publication the authors will transfer to the Biochemical Society the copyright of the paper, which will then not be published elsewhere in the same form, in any language, without the consent of the Society. Authors will be required to sign an undertaking to these effects. The restriction on previous publication does not usually apply to previous publication of oral communications in brief abstract form. In such cases authors should enclose three copies of the abstracts of previous publications. However, the restriction does apply to papers on the WWW. Requests for consent for reproduction of material published in *Clinical Science* should be addressed to the Managing Editor.

2. SUBMISSION OF MANUSCRIPTS: GENERAL INFORMATION AND FORMAT

2.1. General

Papers submitted for publication (together with correspondence about papers, proofs and requests for permission to reproduce material) should be sent to: The Managing Editor, *Clinical Science*, 59 Portland Place, London W1N 3AJ, U.K. [telephone: (UK) 0171-637 5873, (from overseas) +44 171-637 5873; fax: (UK) 0171-323 1136, (from overseas) +44 171-323 1136; e-mail: edit@portlandpress.co.uk]. The covering letter should include the author's telephone and fax numbers and e-mail address. Papers may be submitted electronically as an Adobe Acrobat PDF file either: (i) as uuencoded attachments to the e-mail address: edit@portlandpress.co.uk; or (ii) uploaded to the ftp site: ftp.portlandpress.co.uk in the directory /incoming/ClinSci [N.B. Authors using route (ii) are advised to alert us to any incoming documents via our regular e-mail address.]

Authors in North America should submit their papers to Professor A. E. Taylor, Regional Editor, Department of Physiology, University of South Alabama, College of Medicine, MSB 3024, Mobile, AL 36688-0002, U.S.A. (telephone +1 334 460 7004; fax +1 334 460 6464; e-mail ataylor@jaguar1.usouthal.edu). Authors in the Pacific Rim countries should submit their papers to Professor S. B. Harrap, Regional Editor, University of Melbourne, Department of Physiology, Parkville, Victoria 3052, Australia (telephone +61 3 9344 5836; fax +61 3 9349 4519; e-mail s.harrap@physiology.unimelb.edu.au).

The submission should contain four copies (of which three may be photocopies, except for half-tone figures) of the typescript, Tables, Figures, etc. The authors should retain one copy of the paper. The Editorial Board does not accept responsibility for damage or loss of papers submitted, although great care is taken to ensure safety and confidentiality of the typescript during the editorial process.

Papers should be presented so that they are intelligible to the non-specialist reader of the journal. This is particularly important in highly specialized fields and a very brief résumé of the current state of knowledge is usually helpful. Certain types of material, e.g. mathematical formulations requiring more than trivial derivations, should be given in a separate Appendix.

Where the reader is referred to previous works by the same author(s) for important details relevant to the present work, three copies or reprints of the publication (including papers on the WWW) should be sent with the typescript. This is of particular importance in relation to methodology.

The dates of receipt and acceptance of the paper will be published. If the paper has to be returned to the authors for revision and is not resubmitted within 1 month, the date of receipt will be revised accordingly and the revised paper may be treated as a new submission. It is emphasized that badly presented or unduly long papers will be returned for revision and delays in publication will be inevitable. Similar delays will be incurred if the typescript is not prepared strictly in accordance with the instructions detailed below.

Typescripts of rejected work will not be returned to authors unless a specific request for the return has been made at the time of submission.

2.2. Use of authors' diskettes

Authors should submit diskettes of revised papers to the editorial office. If the revised paper is acceptable every effort will be made to use the diskette during typesetting, but this cannot be guaranteed. Authors must ensure that files have been updated to incorporate all revisions, and hence that the version on the diskette matches the revised hard copy. Our preferred word-processing format is Microsoft Word for Windows version 6.0. Submission of papers in other word-processing formats may lead to delays in processing. The diskettes should be accompanied by a covering letter specifying manuscript number, operating system and software program.

(a) *Text*. Files should be formatted double-spaced with no hyphenation and automatic wordwrap (no hard returns within paragraphs). Please type your text consistently, e.g. take care to distinguish between '1'(one) and 'l' (lower case L), and '0' (zero) and 'O' (capital O), etc.

(b) *Tables*. Tables should be typed as text. The use of graphics programs and 'table editors' should be avoided.

(c) *Figures*. No artwork should be incorporated into the text files. Figures are normally handled conventionally, but artwork may be provided on disk either in TIFF or EPS format and saved as a separate file. We can also accept CorelDraw files. Hard copy of illustrations must also be supplied (see 3.10).

(d) *Mathematics*. In-line equations should be typed as text. The use of graphics programs and 'equation editors' should be avoided. Displayed equations (unless prepared by the 'MathType Equation Editor') are re-keyed by our printer.

2.3. Full Papers

These may be of any length that is justified by their content. Authors should, however, note that because of pressure for space in the journal, no paper, whatever its scientific merits, will be accepted if it exceeds the minimum length required for precision in describing the experiments and clarity in interpreting them. As a guide, most papers published in the journal are of between six and eight printed pages. A concise well-written paper tends to be published more rapidly. Extensive Tables of data can be deposited with the Royal Society of Medicine (see 2.8). *Guidance for Authors* is usually published in the January issue of the journal, and is revised periodically.

The authors should refer to a current issue of *Clinical Science* to make themselves familiar with the general layout. Typescripts should be, in general, arranged as follows:

(a) *Title page*. Title: this should be as informative as possible, since titles of papers are being increasingly used in indexing and coding for information storage and retrieval. The title should indicate the species in which the observations reported have been made. It should not contain any abbreviations. The numbering of parts in a series of papers is not permitted.

List of authors' names (degrees and appointments are not required).

Laboratory or Institute of origin, with full postal address.

Key words: for indexing the subject of the paper; they should, if possible, be selected from the current issues of 'Medical Subject Headings' (MeSH) produced by the *Index Medicus*.

Short title: for use as a running heading in the printed text; it should not exceed 45 characters and spaces and should not contain any abbreviations.

Author for correspondence: the name and address of the author to whom queries and requests for offprints should be sent.

(b) *Summary*. This should be a brief statement arranged in numbered paragraphs of what was done, what was

found and what was concluded, and should rarely exceed 250 words. Abbreviations should be avoided as far as possible and must be defined. Statistical and methodological details including exact doses should also be avoided unless they are essential to the understanding of the Summary.

(c) *Introduction*. This should be comprehensible to the general reader and should contain a clear statement of the reason for doing the work, but should not include either the findings or the conclusions.

(d) *Methods*. The aim should be to give sufficient information in the text or by reference to permit the work to be repeated without the need to communicate with the author.

(e) *Results*. This section should not include material appropriate to the Discussion section.

(f) *Discussion*. This should not contain results and should be pertinent to the data presented.

(g) *Acknowledgments*. These should be as brief as possible.

(h) *References*. See 3.18 for the correct format.

(i) *Figures and Tables*. See 3.10.

2.4. Rapid Communications

The passage of these papers through the editorial process will be expedited and contributors are encouraged to take advantage of this facility when data are novel and exciting, when rapid publication is of importance and when material can be presented concisely. Authors **must** include in their letter of submission a brief statement explaining the novelty of their work. Rapid Communications should describe completed work and should not be merely a preliminary communication.

Rapid Communications should be similar in format to full papers, except that they must occupy not more than four printed pages. This is about 3000 words, with appropriate deductions (at the rate of 1000 words/page) for Figures and Tables.

To achieve rapid publication, authors of accepted Rapid Communications will not be sent proofs. Rejection of a paper submitted as a Rapid Communication does not preclude its re-submission as a full paper for publication in *Clinical Science*, in which event the paper would be reviewed and reports provided with the editorial decision in the normal way.

2.5. Correspondence

Letters containing original observations or critical assessments of material published in *Clinical Science*, including Editorial Reviews, will be considered for the Correspondence section of the journal. All Letters received are subjected to the journal's peer-review procedure. Letters should be no longer than 750 words, with one Figure or Table and up to six references, or 1000 words maximum without a Figure or Table. Letters relating to material previously published in *Clinical Science* should be submitted within 6 months of the appearance of the article concerned. They will be sent to

the authors for comment and both the letter and any reply by the author will be published together. Further correspondence arising therefrom will also be considered for publication. Consideration will also be given to publication of letters on ethical matters.

2.6. Editorial Reviews

These are normally commissioned. However, unsolicited reviews will be considered. Prospective authors should first submit a synopsis of their proposed review rather than the full typescript.

► 2.7. Comments

These are normally commissioned by the Editorial Board.

2.8. Arrangements for large amounts of information

It is impracticable to publish very large sets of individual values or very large numbers of diagrams, and under these circumstances a summary of the information only should be included in the paper. The information from which the summary was derived should be submitted with the typescript and, if the latter is accepted, the Editors may ask for a copy of the full information and diagrams to be deposited with the Librarian, the Royal Society of Medicine, 1 Wimpole Street, London W1M 8AE, U.K., who will issue copies on request. Experience has shown that such requests are frequently received.

2.9. Proof corrections

These are expensive and corrections of other than printers' errors may have to be charged to the author.

► 2.10. Offprints

Offprints may be obtained at terms, based upon the cost of production, that will be given with the proofs. All offprints should be ordered when the proofs are returned (except for Rapid Communications, where they should be ordered when the subedited typescript is returned).

2.11. Availability on MEDLINE and from Adonis

Summaries of papers in *Clinical Science* are available on the MEDLINE system run by the National Library of Medicine, National Institutes of Health, Bethesda, MD, U.S.A.

Full text with illustrations of individual papers can be obtained from Adonis Document Delivery Service, PO Box 839, 1000 AV Amsterdam, The Netherlands.

3. MISCELLANEOUS NOTES

3.1. Abbreviations

Abbreviations should be avoided; if used they must be defined at the first mention; new abbreviations should be coined only for unwieldy names which occur frequently. Abbreviations, except those indicated by an asterisk in the list on pp. viii–x, should not appear in the title and short title nor, if possible, in the Summary. Numbers, not initials, should be used for patients and subjects.

3.2. Anatomical nomenclature

This should follow the recommendations of the International Anatomical Nomenclature Committee (*Nomina Anatomica*. 3rd ed. Amsterdam: Excerpta Medica Foundation, 1966).

3.3. Animals, plants and micro-organisms

The full binomial specific names should be given at first mention for all experimental animals other than common laboratory animals. The strain and, if possible, the source of laboratory animals should be stated. Thereafter in the text, single letter abbreviations may be given for the genus; if two genera with the same initial letter are studied, abbreviations such as *Staph.* and *Strep.* should be used.

3.4. Biochemical nomenclature

As far as possible authors should follow the recommendations of the Nomenclature Committee of IUBMB and IUPAC–IUBMB Joint Commission on Biochemical Nomenclature (see *Biochemical nomenclature and related documents*, 2nd ed., London: Portland Press, 1992; for corrections see *Eur J Biochem* 1993; 213: 1–3).

3.5. Buffers and salts

The acidic and basic components should be given, together with the pH. Alternatively, a reference to the composition of the buffer should be given. Further details are provided in *Biochem J* 1998; 329: 1–16.

When describing solutions containing organic anions and their parent acids, the salt designator (e.g. lactate, urate, oxalate) should be used in preference to the name of the acid (lactic, uric, oxalic) unless it is certain that virtually all of the acid is in the undissociated form.

The composition of incubation media should be described, or a reference to the composition should be given.

3.6. Computer modelling

Papers concerned primarily with computer modelling techniques are acceptable provided that use of such techniques leads to a clear choice between two or more alternative hypotheses, or to the formulation of a new hypothesis amenable to experimental challenge or verification, or provides some new insight into the behaviour of a particular physiological system. Extensive technical details of hardware and software should not be given.

3.7. Doses

Doses of drugs should be expressed in mass terms, e.g. milligrams (mg) or grams (g), and also (in parentheses) in molar terms, e.g. mmol, mol, where this appears to be relevant. Molecular masses of many drugs may be found in *The Merck Index*. 11th ed. Rahway, NJ, U.S.A.: Merck and Co. Inc., 1989.

3.8. Enzymes

Nomenclature should follow that given in *Enzyme Nomenclature* (San Diego: Academic Press, 1992); for

corrections and additions see *Eur J Biochem* 1994; **223**: 1–5 and *Eur J Biochem* 1995; **232**: 1–6. The Enzyme Commission (EC) number should be quoted at the first mention. Where an enzyme has a commonly used informal name, this may be employed after the first formal identification. A unit of enzyme activity can be expressed as that amount of material which will catalyse transformation of 1 μmol of the substrate/s under defined conditions, including temperature and pH. This gives the unit of the amount of enzyme named the katal (symbol kat). Alternatively, or when the natural substrate has not been fully defined, activity should be expressed in terms of units of activity relative to that of a recognized reference preparation, assayed under identical conditions. Activities of enzymes should normally be expressed as units/ml or units/mg of protein.

3.9. Evaluation of measurement procedures

When a new measuring procedure has been used, or when an established procedure has been applied in a novel fashion, an estimate of the precision of the procedure should be given. This should, as far as possible, indicate what sources of variation have been included in this estimate, e.g. variation of immediate replication, variation within different times of day, or from day to day, etc.

If the precision of measurement varies in proportion to the magnitude of the values obtained, it can best be expressed as the coefficient of variation; otherwise it should be expressed by an estimate of the (constant) standard error of a single observation, or by estimates of several points within the range of observed values.

When recovery experiments are described the approximate ratio of the amount added to the amount already present and the stage of the procedure at which the addition was made should be stated.

For methods or assays crucial to the understanding of the paper, information should normally be provided on the validity, accuracy and precision of those methods.

3.10. Figures and Tables

Their number should be kept to a minimum. Their appropriate position in the paper should be indicated in the margin of the text. References to Figures and Tables should be in arabic numerals, e.g. Figure 3, and they should be numbered in order of appearance. In general, the same data should not be presented in both a Figure and a Table.

Figures should be supplied in a form that can be reproduced directly by the printer, together with photocopies. All Figures should have their number and the authors' names written in pencil on the back; the top of the Figure should be indicated with a pencilled arrow. Figures are not routinely relettered. Authors should ensure that nomenclature, abbreviations, etc. used in lettering of Figures correspond to those used in the text. Separate panels within Figures should be clearly marked (a), (b), (c), etc. so that they can be referred to easily in the legend

and text. Acceptable symbols for experimental points are ●, ▲, ■, ○, △, □. The symbols × or + should be avoided. Symbols should not be generated by using tints or a graphics program. The same symbols must not be used for two curves where the points might be confused. For scatter diagrams, solid symbols are preferred. When a particular variable appears in more than one Figure, the same symbol should be used for it throughout, if possible.

Curves should not be drawn beyond the experimental points, nor should axes extend appreciably beyond the data. Only essential information that cannot readily be included in the legend should be written within the Figure.

The use of tints should be avoided; however, if tints are necessary, please ensure that a dot fill of 100 lines per inch or lower is used. Columns in histograms should be differentiated by the use of simple hatching etc.

Figures for half-tone reproduction should be submitted as glossy prints. **Four copies (not photocopies) of each print should be provided.** All lettering should be placed directly on to the Figure, not on a clear film overlay. Where the magnification is to be indicated (e.g. on electron micrographs), this should be done by adding a bar representing a stated length.

Colour figures are accepted when, in the opinion of the Editorial Board, they are essential to illustrate a particular scientific point. Authors will normally be required to pay the full cost of colour separation and printing (at 1998 prices, approximately £550 for the first Figure and £300 for each subsequent Figure).

Tables should be typed separately from the text. They should have an underlined title followed by any legend. Parameters being measured, with units if appropriate, should be clearly indicated in the column headings.

Captions for the Figures, and titles and legends for the Tables, should make them **readily understandable** without reference to the text. Adequate statistical information, including that on regression lines, should be included in Figure captions where appropriate.

Care is needed when using powers in Figure and Table headings to avoid numbers with too many digits (see 3.17).

3.11. Footnotes

These should be avoided as far as possible but where they are used in Tables they should be identified by the symbols *†‡§||¶, in that order.

3.12. 'Homology'

The term 'homologous' has a precise meaning in biology of 'having a common evolutionary origin', but it has often been used in work on protein and nucleic acid sequences to mean simply 'similar'. A group of experts has urged that the interests of clarity are best served by restricting use to the more precise definition (Reeck GR, et al. *Cell* 1987; **50**: 667; Lewin R. *Science* 1987; **237**: 1570). *Clinical Science* agrees with these arguments and seeks to preserve the distinction between 'homologous' and 'similar' in its pages.

3.13. Isotope measurements

Where possible radioactivity should be expressed in absolute terms; the SI unit for radioactivity is the becquerel (Bq), defined as 1 disintegration/s, but the Curie (Ci; $1 \text{ Ci} = 3.7 \times 10^{10} \text{ Bq}$) may also be used. Alternatively, radioactivity may be expressed as disintegrations (or counts) per unit of time, e.g. disintegrations/s (d.p.s.) or counts/min (c.p.m.).

3.14. Radionuclide applications in man

If new or modified radionuclide applications in man are described, an estimate of the maximal possible radiation dose to the body and critical organs should be given.

For the time being this can continue to be expressed in rems, but with the corresponding figure in sieverts (Sv) given in parentheses after it.

3.15. Methods

In describing certain techniques, namely centrifugation (when the conditions are critical), chromatography and electrophoresis, authors should follow the recommendations published by the Biochemical Society (currently, *Biochem J* 1998; **329**: 1–16).

3.16. Nomenclature of disease

This should follow the International Classification of Disease (9th revision, Geneva: World Health Organization, 1979) as far as possible. The correct abbreviation for insulin-dependent diabetes is type I diabetes (*not* IDDM), and for non-insulin-dependent diabetes is type II diabetes (*not* NIDDM).

3.17. Powers in Tables and Figures

Care is needed where powers are used in Table headings and in Figures to avoid numbers with an inconvenient number of digits. For example: (i) an entry '2' under the heading $10^3 k$ means that the value of k is 0.002; an entry '2' under the heading $10^{-3} k$ means that the value of k is 2000. (ii) A concentration 0.00015 mol/l may be expressed as 0.15 under the heading 'concn. (mmol/l)' or as 150 under heading 'concn. ($\mu\text{mol/l}$)' or as 15 under the heading ' $10^5 \times \text{concn. (mol/l)}$ ', but not as 15 under the heading 'concn. ($\text{mol/l} \times 10^{-5}$)'.

3.18. References

The 'Vancouver' system is used: references in the text are numbered consecutively in the order in which they are first mentioned, the numerals being given in brackets, e.g. [22]. References cited in Figure legends or Tables only should be numbered in a sequence determined by the position of the first mention in the text of the Figure or Table. References should be listed in numerical order and the names of all authors of a paper should be given (except where there are seven or more when only the first three should be listed and *et al.* added), with the full title of the paper and the source details in full including the first and last page numbers, e.g.

- Howard JK, Lord GM, Clutterbuck EJ, Ghatei MA, Pusey CD, Bloom SR. Plasma immunoreactive leptin concentration in end-stage renal disease. *Clin Sci* 1997; **93**: 119–26.

When the quotation is from a book, the following format should be used, giving the relevant pages or chapter number:

- Cornish-Bowden A. *Fundamentals of enzyme kinetics*. London: Portland Press Ltd, 1995.
- Hainsworth R, Drinkhill MJ. Regulation of blood volume. In: Jordan D, Marshall JM, eds. *Cardiovascular regulation*. London: Portland Press Ltd, 1995: 77–91.

References to 'personal communications' and 'unpublished work' should appear in the text only and not in the list of references. The name and initials of the source of information should be given. In the case of quotations from personal communications the authors **must** provide documentary evidence that permission for quotation has been obtained. When the reference is to material that has been accepted for publication but has not yet been published, this should be indicated in the list of references by 'In press' together with the name of the relevant journal and, if possible, the expected date of publication. If such a citation is of major relevance to the manuscript submitted for publication authors are advised that the editorial process might be expedited by the inclusion of a copy of such work.

3.19. Solutions

Concentration of solutions should be described where possible in molar terms (mol/l and subunits thereof), stating the molecular particle weight if necessary. Values should not be expressed in terms of normality or equivalents. Mass concentration should be expressed as g/l or subunits thereof, for example mg/l or $\mu\text{g/l}$. For solutions of salts, molar concentration is always preferred to avoid ambiguity as to whether anhydrous or hydrated compounds are used. Concentrations of aqueous solutions should be given as mol/l or mol/kg (g/l or g/kg if not expressed in molar terms) rather than % (w/v) or % (w/w). It should always be made clear whether concentrations of compounds in a reaction mixture are final concentrations or the concentrations in solutions added.

3.20. Spectrophotometric data

The general name for the quantity $\log(I_0/I)$ is attenuation, and it reduces to absorbance when there is negligible scattering or reflection. The more general term 'attenuance' should be used when scattering is considerable, e.g. when the quantity is measured to estimate the cell density of a culture. Otherwise the term absorbance should be used; neither should be called extinction or optical density. Symbols used are: A , absorbance; D , attenuation; a , specific absorption coefficient ($\text{litre} \cdot \text{g}^{-1} \cdot \text{cm}^{-1}$) (alternatively use $A_{1\text{cm}}^{1\%}$); ϵ , molar absorption coefficient (the absorbance of a molar solution in a 1 cm light-path) ($\text{litre} \cdot \text{mol}^{-1} \cdot \text{cm}^{-1}$, *not* $\text{cm}^2 \cdot \text{mol}^{-1}$).

3.21. Spelling

Clinical Science uses as standards for spelling the Concise or Shorter Oxford Dictionary of Current English

(Oxford: Clarendon Press) and Butterworth's Medical Dictionary (London: Butterworths).

3.22. Statistics

Papers are frequently returned for revision (and their publication consequently delayed) because the authors use inappropriate statistical methods. Two common errors are the use of means, standard deviations and standard errors in the description and interpretation of grossly non-normally distributed data and the application of *t*-tests for the significance of difference between means in similar circumstances, or when the variances of the two groups are non-homogeneous. In some circumstances it may be more appropriate to provide a 'scattergram' than a statistical summary. Authors are recommended to consult the statistical guidelines presented by Altman et al. in 'Statistical guidelines for contributors to medical journals' *Br Med J* 1983; **286**: 1489–93.

The type of statistical test used should be stated in the Methods section. A reference should be given for the less commonly encountered statistical tests. The format for expressing mean values and standard deviations or standard errors of the mean is, for example: mean cardiac output 10.4 litres/min (S.D. 1.2; *n* = 11). Degrees of freedom should be indicated where appropriate. Levels of significance are expressed in the form $P < 0.01$.

3.23. Trade names

The name and address of the supplier of special apparatus and of biochemicals should be given. Registered trademarks should be identified by the symbol ® where they appear in the text. In the case of drugs, approved names should always be given with trade names and manufacturers in parentheses.

4. UNITS: THE SI SYSTEM

The recommended *Système International* (SI) units (see Quantities, units and symbols in physical chemistry. Oxford: Blackwell Scientific Publications Ltd, 1988) are used by *Clinical Science*. **All papers submitted should use these units** except for blood pressure values, which should be expressed in mmHg, and gas partial pressures, where values at the author's discretion may be given in mmHg (with kPa in parentheses) or as kPa (with mmHg in parentheses). Airways pressure should be expressed in kPa. Where molecular mass is known, the amount of a chemical or drug should be expressed in mol or in an appropriate subunit, e.g. mmol. Energy should be expressed in joules (J).

The basic SI units and their symbols are as follows:

<i>Physical quantity</i>	<i>Name</i>	<i>Symbol</i>
length	metre	m
mass	kilogram	kg
time	second	s
electric current	ampere	A
thermodynamic temperature	kelvin	K
luminous intensity	candela	cd
amounts of substance	mole	mol

The following are examples of derived SI units:

<i>Physical quantity</i>	<i>Name</i>	<i>Symbol</i>	<i>Definition</i>
energy	joule	J	$\text{kg}\cdot\text{m}^2\cdot\text{s}^{-2}$
force	newton	N	$\text{kg}\cdot\text{m}\cdot\text{s}^{-2} = \text{J}\cdot\text{m}^{-1}$
power	watt	W	$\text{kg}\cdot\text{m}^2\cdot\text{s}^{-3} = \text{J}\cdot\text{s}^{-1}$
pressure	pascal	Pa	$\text{kg}\cdot\text{m}^{-1}\cdot\text{s}^{-2} = \text{N}\cdot\text{m}^{-2}$
electric charge	coulomb	C	A·s
electric potential difference	volt	V	$\text{kg}\cdot\text{m}^2\cdot\text{s}^{-2}\cdot\text{A}^{-1} = \text{J}\cdot\text{A}^{-1}\cdot\text{s}^{-1}$
electric resistance	ohm	Ω	$\text{kg}\cdot\text{m}^2\cdot\text{s}^{-3}\cdot\text{A}^{-2} = \text{V}\cdot\text{A}^{-1}$
electric conductance	siemens	S	$\text{kg}^{-1}\cdot\text{m}^{-2}\cdot\text{s}^3\cdot\text{A}^2 = \Omega^{-1}$
electric capacitance	farad	F	$\text{A}^2\cdot\text{s}^3\cdot\text{kg}^{-1}\cdot\text{m}^{-2} = \text{A}\cdot\text{s}\cdot\text{V}^{-1}$
frequency	hertz	Hz	s^{-1}
volume	litre	l	10^{-3} m^3

The word 'litre' has been accepted as a special name for cubic decimetre (1 litre = 1 dm³).

Both the basic and derived SI units, including the symbols of derived units that have special names, may be preceded by prefixes to indicate multiples and sub-multiples. The prefixes should be as follows:

<i>Multiple</i>	<i>Prefix</i>	<i>Symbol</i>	<i>Multiple</i>	<i>Prefix</i>	<i>Symbol</i>
10 ⁶	mega	M	10 ⁻³	milli	m
10 ³	kilo	k	10 ⁻⁶	micro	μ
10 ²	hecto	h*	10 ⁻⁹	nano	n
10	deka	da	10 ⁻¹²	pico	p
10 ⁻¹	deci	d*	10 ⁻¹⁵	femto	f
10 ⁻²	centi	c*			

*To be avoided where possible (except for cm).

Compound prefixes should not be used, e.g. 10⁻⁹ m should be represented by 1 nm, not 1 mμm.

Notes

- (i) Full stops are not used after symbols.
- (ii) Minutes (min), hours (h), days and years will continue to be used in addition to the SI unit of time [the second(s)].
- (iii) The solidus may be used in a unit as long as it does not have to be employed more than once, e.g. mmol/l is acceptable, but ml/min/kg is not, and should be replaced by $\text{ml}\cdot\text{min}^{-1}\cdot\text{kg}^{-1}$.

5. ABBREVIATIONS, CONVENTIONS, DEFINITIONS, SYMBOLS AND SPECIAL COMMENTS

Standard symbols and abbreviations that can be used without definition are indicated by an asterisk; this list also shows selected abbreviations in the form of groups of capital letters (e.g. ALA, ECF, MCHC) which when used must be defined in the text as indicated on p. iv. The standard abbreviations for amino acids are only for use in Figures and Tables or for peptide sequences.

absorbance	<i>A</i>	cycle/s	Hz
acceleration due to gravity	<i>g</i>	cysteine	Cys
adenosine 3':5'-cyclic mono-phosphate (cyclic AMP)	cAMP*	dates	e.g. 11 August 1998
adenosine 5'-phosphate	AMP*	dead-space minute ventilation	\dot{V}_D
adenosine 5'-diphosphate	ADP*	dead-space volume	V_D
adenosine triphosphatase	ATPase*	degrees, Celsius or centigrade	°C
adenosine 5'-triphosphate	ATP*	deoxy (prefix)	<i>not</i> desoxy
adrenoceptor (<i>see also</i> blocking agents)		deoxycorticosterone	DOC
adrenocorticotrophic hormone	ACTH	deoxycorticosterone acetate	DOCA
alanine	Ala	deoxyribonucleic acid	DNA*
alternating current	a.c.*	complementary	cDNA*
alveolar minute ventilation	\dot{V}_A	deoxyribonuclease	DNase*
alveolar to arterial oxygen partial pressure difference	$(P_{AO_2} - P_{aO_2})$	diabetes, insulin-dependent	type I (<i>not</i> IDDM)
aminolaevulinic acid	ALA	diabetes, non-insulin-dependent	type II (<i>not</i> NIDDM)
ampere	A	diethylaminoethylcellulose	DEAE-cellulose*
angiotensin	ANG; reference amino acid abbreviations are used as prefix within brackets: e.g. [Sar ¹ , Val ⁵ , Ala ⁸]ANG	differential of <i>x</i> with respect to time	\dot{x} (= dx/dt)
ångstrom	Å (1 ångstrom = 10 ⁻¹ nm)	1,25-dihydroxycholecalciferol	1,25-(OH) ₂ D ₃
antidiuretic hormone	ADH (when referring to the physiological secretion)	dilute	dil.
arginine	Arg	dimethyl sulphoxide	DMSO*
arteriovenous	a - v: <i>permitted</i> in Figures and Tables	2,3-diphosphoglycerate	2,3-DPG
asparagine	Asn	direct current	d.c.*
aspartic acid	Asp	disintegrations/min	d.p.m.*
atmosphere (unit of pressure)	<i>not used</i> ; express in kPa (1 atmosphere = 101.325 kPa)	disintegrations/s	d.p.s.*
attenuance	<i>D</i>	dissociation constant	
base pair	bp*	acidic	<i>K_a</i>
becquerel	Bq (1 d.p.s.)	apparent	e.g. <i>K'_a</i>
blocking agents	e.g. β-adrenoceptor antagonists preferred	basic	<i>K_b</i>
blood pressure	express in mmHg	minus log of	p <i>K</i>
blood urea nitrogen	<i>not used</i> ; recalculate as urea, express in mmol/l	doses	avoid Latin designations such as b.d. and t.i.d.
blood volume	BV	dyne	dyn
body temperature and pressure, saturated	BTPS*	elastance	<i>E</i> ; express in Pa·m ⁻³
bovine serum albumin	BSA*	electrocardiogram	ECG*
British Pharmacopoeia calculated	write in full and give edition calc. (in Tables only)	electroencephalogram	EEG*
'Calorie' (= 1000 cal)	<i>not used</i> ; recalculate as kilojoules (1 'Calorie' = 4.184 kJ)	electromotive force	e.m.f.*
carbon dioxide output (in respiratory physiology)	\dot{V}_{CO_2} ; express in ml STP/min	electron paramagnetic (or spin) resonance	EPR*, ESR*
cardiac frequency	<i>f_c</i> in beats/min	electronvolt	eV (or radiation energies)
cardiac output	express in l/min	enzyme-linked immunosorbent assay	ELISA*
centimetre	cm	equation	eqn.
clearance of <i>x</i>	<i>C_x</i>	equivalents (amount of a chemical)	<i>not used</i> ; recalculate in molar terms
coenzyme A and its acyl derivatives	CoA* and acyl-CoA*	erythrocyte count	express as 10 ¹² cells/l
compare	cf.	ethanol, ethanolic	<i>not</i> ethyl alcohol or alcoholic
complement components	C1-C9*	ethylenediaminetetra-acetate	EDTA*
compliance (respiratory physiology)	<i>C</i> ; express in l·kPa ⁻¹	'ethyleneglycolbis(aminoethyl-ether)tetra-acetate'	EGTA*
concentrated	conc.	exchangeable	Na _c , K _c etc. for total exchangeable sodium, potassium etc.
concentration	concn.; may be denoted [], e.g. plasma [HCO ₃ ⁻]	Experiment (with reference numeral)	Expt.; plural, Expts.
concentration giving half-maximal response	EC ₅₀ *	expired minute ventilation	\dot{V}_E
concentration giving half-maximal inhibition	IC ₅₀ *	extinction	<i>use</i> absorbance
conductance (respiratory physiology)	<i>G</i> ; express in l·s ⁻¹ ·kPa ⁻¹	extracellular fluid	ECF
correlation coefficient	<i>r</i>	extracellular fluid volume	ECFV
counts/min, counts/s	c.p.m.*, c.p.s.*	extraction ratio of <i>x</i> (renal)	<i>E_x</i>
cubic centimetres	<i>use</i> ml	fast protein liquid chromatography	FPLC*
curie	Ci (1 Ci = 3.7 × 10 ¹⁰ d.p.s.)	filtered load of <i>x</i> (renal)	<i>F_x</i>
		flavin-adenine dinucleotide	FAD*
		flavin mononucleotide	FMN*
		follicle-stimulating hormone	FSH
		forced expiratory volume in 1.0 s	FEV _{1.0}
		fractional concentration in dry gas	<i>F</i>
		fractional disappearance rate	<i>k</i> (as in $A = A_0 e^{-kt}$)
		frequency of respiration	<i>f_R</i> ; in breaths/min
		functional residual capacity	FRC
		gas-liquid chromatography	GLC*
		gas transfer factor	<i>T</i> ; in mmol·min ⁻¹ ·kPa ⁻¹

glomerular filtration rate	GFR	luteinizing hormone	LH
glutamic acid	Glu	lysine	Lys
glutamine	Gln	mass spectrometry	MS*
glutathione	GSH (reduced); GSSG (oxidized)	maximum	max.
glycine	Gly	mean corpuscular haemoglobin	MCH; express in pg
gram	g	mean corpuscular haemoglobin concentration	MCHC; express in g/dl
gravitational field, unit of (9.81 m·s ⁻¹)	g	mean corpuscular volume	MCV; express in fl (1 μm ³ = 1 fl)
gray	Gy (100 rads)	melting point	m.p.
growth hormone	GH; if human, hGH	<i>meta</i> -	<i>m</i> -
guanine-nucleotide-binding regulatory protein	G-protein*	methanol, methanolic	<i>not</i> methyl alcohol
haematocrit	Hct; no units	methionine	Met
haemoglobin	Hb*; express in g/dl	metre	m
half-life	t _{1/2}	Michaelis constant	K _m
hertz (s ⁻¹)	Hz	micromole	μmol
high-pressure (or high- performance) liquid chromatography	HPLC*	micron (10 ⁻⁶ m)	μm; <i>not</i> μ
histidine	His	milliequivalent	<i>not used</i> ; give amount in mmol
hour	h	millilitre	ml
human chorionic gonadotropin	hCG	millimetre of mercury	mmHg; for blood pressure and, at authors' discretion, for gas partial pressures: see p. vii (1 mmHg = 0.133 kPa)
human placental lactogen	hPL	millimolar (concentration)	mmol/l or mM
hydrocortisone	<i>use</i> cortisol	millimole	mmol
hydrogen ion activity	aH; express in nmol/l	minimum	min.
minus log of	pH	minute (60 s)	min
25-hydroxycholecalciferol	25-(OH)D ₃	molal	mol/kg
4-(2-hydroxyethyl)-1- piperazine-ethanesulphonic acid	Hepes*	molar (concentration)	mol/l or M
hydroxyproline	Hyp	molar absorption coefficient	ε (the absorbance of a molar solution in a 1 cm light-path)
immunoglobulins	IgA, IgD, IgE, IgG, IgM*	mole	mol
infrared	IR*	molecular mass	express in Da or kDa
injection routes:	<i>use</i> abbreviations only in Figures	molecular mass (relative)	M _r (no units)
intra-arterial	i.a.	4-morpholine	Mops*
intramuscular	i.m.	propanesulphonic acid	
intraperitoneal	i.p.	nicotinamide-adenine dinucleotide	NAD if oxidation state not indicated*
intravenous	i.v.		NAD ⁺ if oxidized*
subcutaneous	s.c.	nicotinamide-adenine dinucleotide phosphate	NADH if reduced*
international unit	i.u. (definition and reference should be given for uncommon or ambiguous applications, e.g. enzymes)		NADP if oxidation state not indicated*
intracellular fluid	ICF	normal	NADP ⁺ if oxidized*
intracellular fluid volume	ICFV		NADPH if reduced*
ionic strength	I	normal temperature and pressure	should not be used to denote the concentration or osmolarity of a solution
isoleucine	Ile	normal magnetic resonance	<i>use</i> standard temperature and pressure (STP*)
isotonic	specify composition of fluid, e.g. 150 mmol/l NaCl	nuclear magnetic resonance	NMR*
isotopically labelled compounds	e.g. [U- ¹⁴ C]glucose, [1- ¹⁴ C]glucose, sodium [1- ¹⁴ C]acetate; <i>use</i> ¹³¹ I-labelled albumin, <i>not</i> [¹³¹ I]albumin for simple molecules: ¹⁴ CO ₂ , ³ H ₂ O	number (in enumerations)	no. (in Tables only)
joule	J	observed	obs. (in Tables only)
katal	kat	ohm	Ω
kilobases	kb*	ornithine	Orn
kilogram	kg	<i>ortho</i> -	<i>o</i> -
lactate dehydrogenase	LDH	orthophosphate (inorganic)	P _i
leucine	Leu	osmolarity	express in osmol (or mosmol)/l
leucocyte count	express as 10 ⁹ cells/l	oxygen uptake per minute (in respiratory physiology)	V _{O₂} ; express in ml STP/min
lipoproteins (serum)		packed cell volume	PCV; express in %
high density	HDL	page, pages	p., pp.
low density	LDL	<i>para</i> -	<i>p</i> -
very low density	VLDL	<i>para</i> -aminohippurate	PAH
litre	l (write in full if confusion with the numeral 1 is possible)	partial pressure	P; express in either kPa or mmHg (see p. vii)
logarithm (base 10)	log		P _{AO₂}
logarithm (base e)	ln	e.g. alveolar, of O ₂	P _{ACO₂}
		arterial, of CO ₂	P _{capO₂}
		capillary, of O ₂	P _{ETCO₂}
		end-tidal, of CO ₂	P _{VCO₂}
		mixed venous, of CO ₂	
		pascal	Pa
		per	/
		per cent	%

petroleum ether	<i>not used; use light petroleum and give boiling range</i>	sodium dodecyl sulphate species	SDS* sp., plural spp.
phenylalanine	Phe	specific activity	sp. act. Confusion must be avoided between e.g. specific radioactivity and the specific activity of an enzyme
phenylmethanesulphonyl fluoride	PMSF*		sGaw; express in $s^{-1} \cdot kPa^{-1}$
phosphate-buffered saline	PBS*	specific conductance of airways	S.D.*
plasma renin activity	express as pmol of angiotensin $I \cdot h^{-1} \cdot ml^{-1}$	standard deviation	S.E.M.*
plasma volume	PV	standard error of the mean	STP
poise	1 poise = $10^{-1} N \cdot s \cdot m^{-2}$	standard temperature and pressure	
polyacrylamide-gel electrophoresis	PAGE*	steroid nomenclature	see Eur J Biochem 1989; 186 : 429–58 and Eur J Biochem 1993; 213 : 1–3
potential difference	p.d.	sulphydryl	<i>use</i> thiol or SH
power output	W (1 kpm/min = 0.1635 W)	sum	Σ
precipitate	ppt.	Svedberg unit	S
pressure	<i>P</i> ; express in kPa (except for blood pressures and gas partial pressures: see p. vii); 1 kPa = 7.5 mmHg	temperature (absolute) (empirical)	<i>T</i>
probability of an event being due to chance alone	<i>P</i>	temperature, thermodynamic	<i>t</i>
proline	Pro	thin-layer chromatography	K
pulmonary capillary blood flow	\dot{Q}_c	threonine	TLC*
pyrophosphate (inorganic)	PP _i *	thyrotrophic hormone	Thr
rad (radiation dose; 10^{-5} J absorbed/g of material)	not abbreviated (100 rads = 1 Gy)	thyrotrophin-releasing hormone	TSH
radioimmunoassay	RIA*	tidal volume	TRH
red blood cell	<i>use</i> erythrocyte; express counts as 10^{12} cells/l	time (symbol)	V_T
relative band speed (partition chromatography)	R_F	time of day	<i>t</i>
rem	100 ergs/s \times quality factor	torr	e.g. 18.15 hours
renin	<i>see</i> plasma renin activity	tryptophan	not used; <i>use</i> kPa (1 torr = 0.133 kPa)
residual volume	RV	tubular maximal reabsorptive capacity for x	Trp
resistance (rheological)	<i>R</i> ; express in $kPa \cdot l^{-1} \cdot s$	tyrosine	$T_{m,x}$
respiratory exchange ratio (pulmonary)	<i>R</i>	ultraviolet	Tyr
respiratory quotient (metabolic)	RQ	urinary concentration of x	UV*
revolutions	rev.	valency	U_x
rev./min	<i>not</i> r.p.m.; <i>use</i> g if possible (see p. ix)	valine	e.g. Ca^{2+} , <i>not</i> Ca^{++}
ribonucleic acid messenger transfer	RNA*	variance ratio	Val
ribonuclease	mRNA*	vascular resistance	<i>F</i>
röntgen	tRNA*	velocity	express in $kPa \cdot l^{-1} \cdot s$ (with value in $dyn \cdot s \cdot cm^{-5}$ in parentheses); primary values of differential vascular pressure (mmHg) and flow (l/min) should always also be given in Tables or text as appropriate
saline	RNase*	venous admixture	v ; express as $m \cdot s^{-1}$
saturation	R	viscosity, dynamic	Q_{va}
	define at first mention [e.g. NaCl solution (154 mmol/l)]	viscosity, kinematic	η
	<i>S</i> , e.g. S_{aO_2} for arterial oxygen saturation (see partial pressure for other analogous abbreviations)	vital capacity	<i>v</i>
		volt	VC
second (time)	s	volume of blood (in cardio-respiratory physiology)	V
serine	Ser	watt	Q ; <i>use</i> \dot{Q} for blood flow rate
sievert	Sv (1 J/kg \times quality factor)	wavelength	W
solvent systems	e.g. butanol/acetic acid/water (4:1:1, by vol.), butanol/acetic acid (4:1, v/v)	weight	λ
		white blood cell	wt.
			<i>use</i> leucocyte; express counts as 10^9 cells/l

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