



US00RE37534E

(19) **United States**
(12) **Reissued Patent**
Hider et al.

(10) **Patent Number:** **US RE37,534 E**
(45) **Date of Reissued Patent:** **Jan. 29, 2002**

(54) **PHARMACEUTICAL COMPOSITIONS**
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(21) Appl. No.: **08/792,456**
(22) Filed: **Jan. 31, 1997**
(Under 37 CFR 1.47)

Related U.S. Patent Documents

Reissue of:
(64) Patent No.: **4,834,983**
Issued: **May 30, 1989**
Appl. No.: **07/054,086**
Filed: **May 26, 1987**

U.S. Applications:

(63) Continuation of application No. 08/521,281, filed on Aug. 30, 1995, now abandoned, which is a continuation of application No. 08/064,800, filed on May 21, 1993, now abandoned, which is a continuation of application No. 07/708,043, filed on May 30, 1991, now abandoned, which is a continuation of application No. 06/601,485, filed on Apr. 18, 1984, now abandoned, which is a continuation-in-part of application No. 06/542,976, filed on Oct. 18, 1993, now Pat. No. 4,575,502.

(30) **Foreign Application Priority Data**

Oct. 22, 1982 (GB) 8230150
Oct. 14, 1983 (EP) 83306246
Oct. 14, 1983 (GB) 8327612
Jan. 31, 1984 (IE) 84/0222
(51) **Int. Cl.⁷** **A61K 31/555**
(52) **U.S. Cl.** **514/184; 514/814**
(58) **Field of Search** 514/184, 814

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(57) **ABSTRACT**

Pharmaceutical compositions containing an iron complex of 3-hydroxy-4-pyrone or of a 3-hydroxy-4-pyrone in which one or more of the hydrogen atoms attached to ring carbon atoms are replaced by an aliphatic hydrocarbon group of 1 to 6 carbon atoms, are of value for the treatment of iron deficiency anaemia.

40 Claims, 6 Drawing Sheets

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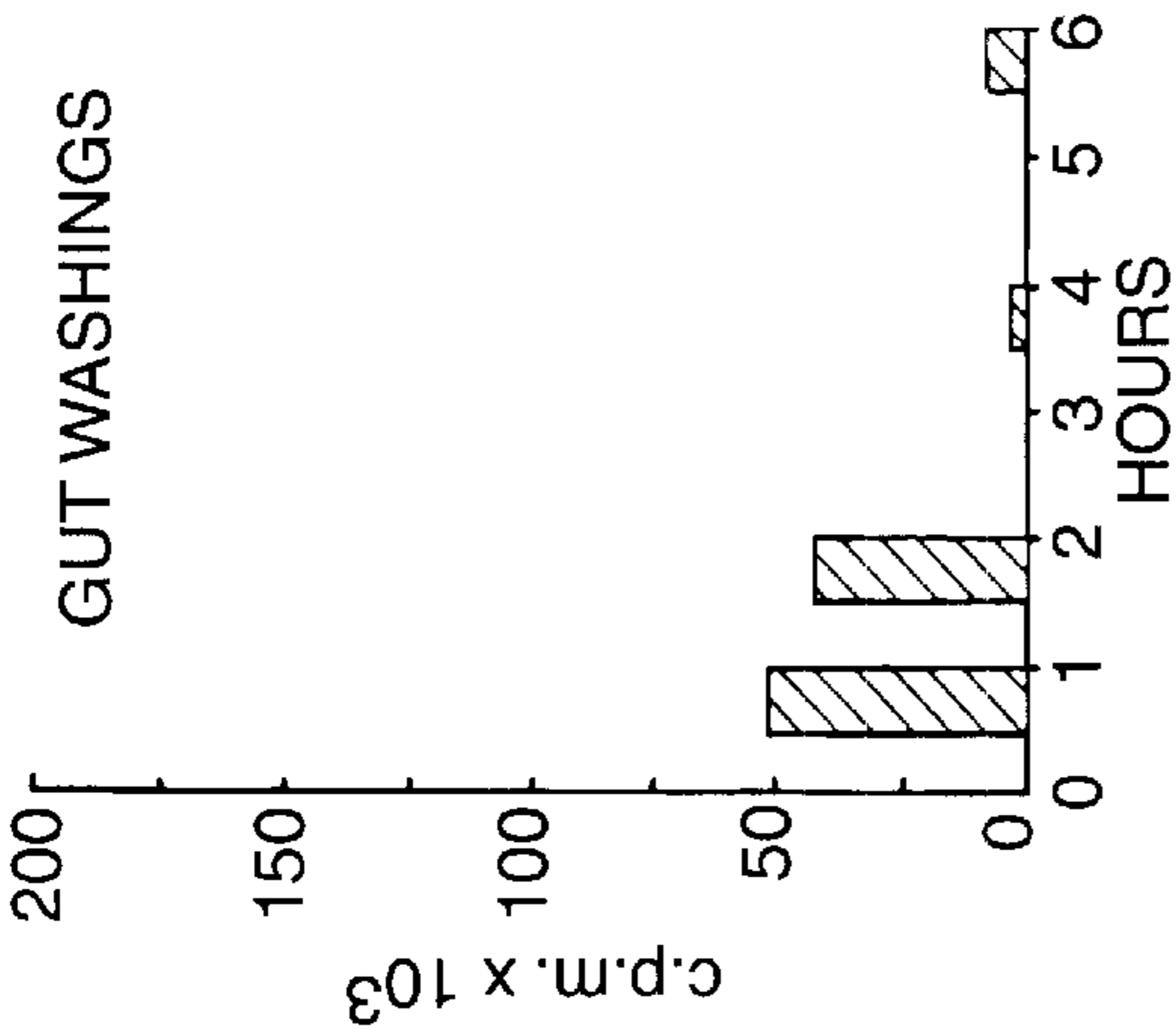


Fig. 1A

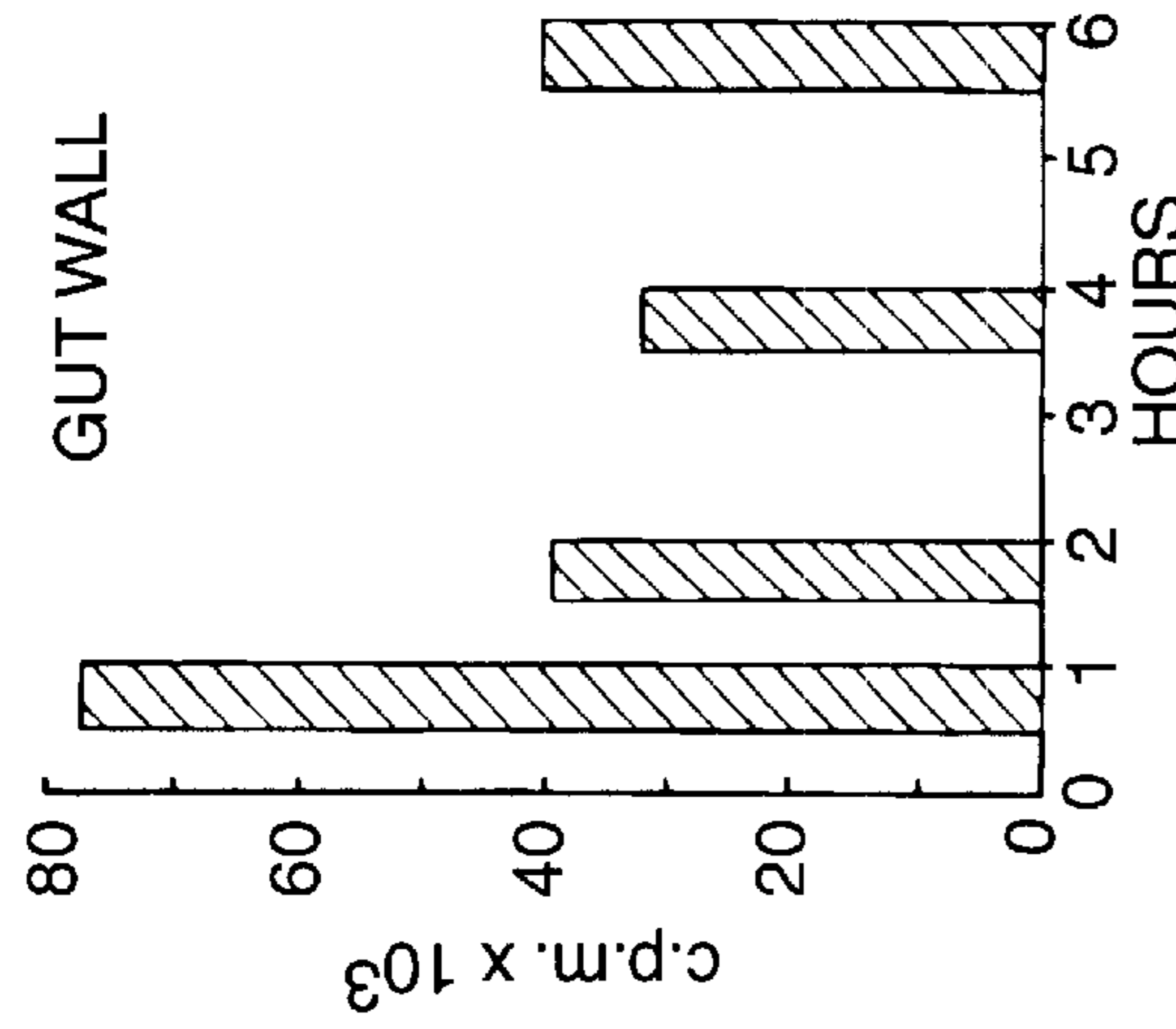


Fig. 1B

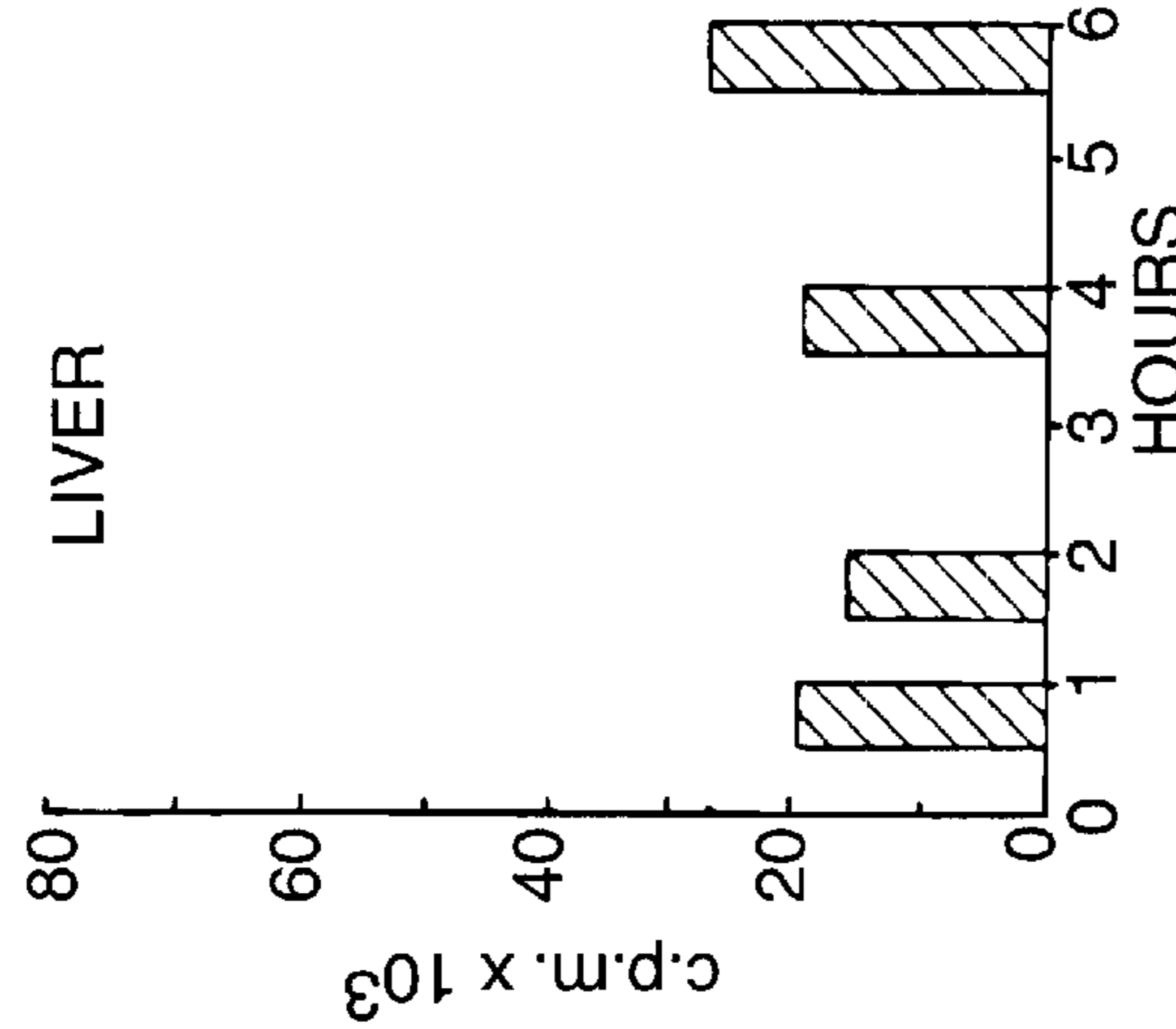


Fig. 1C

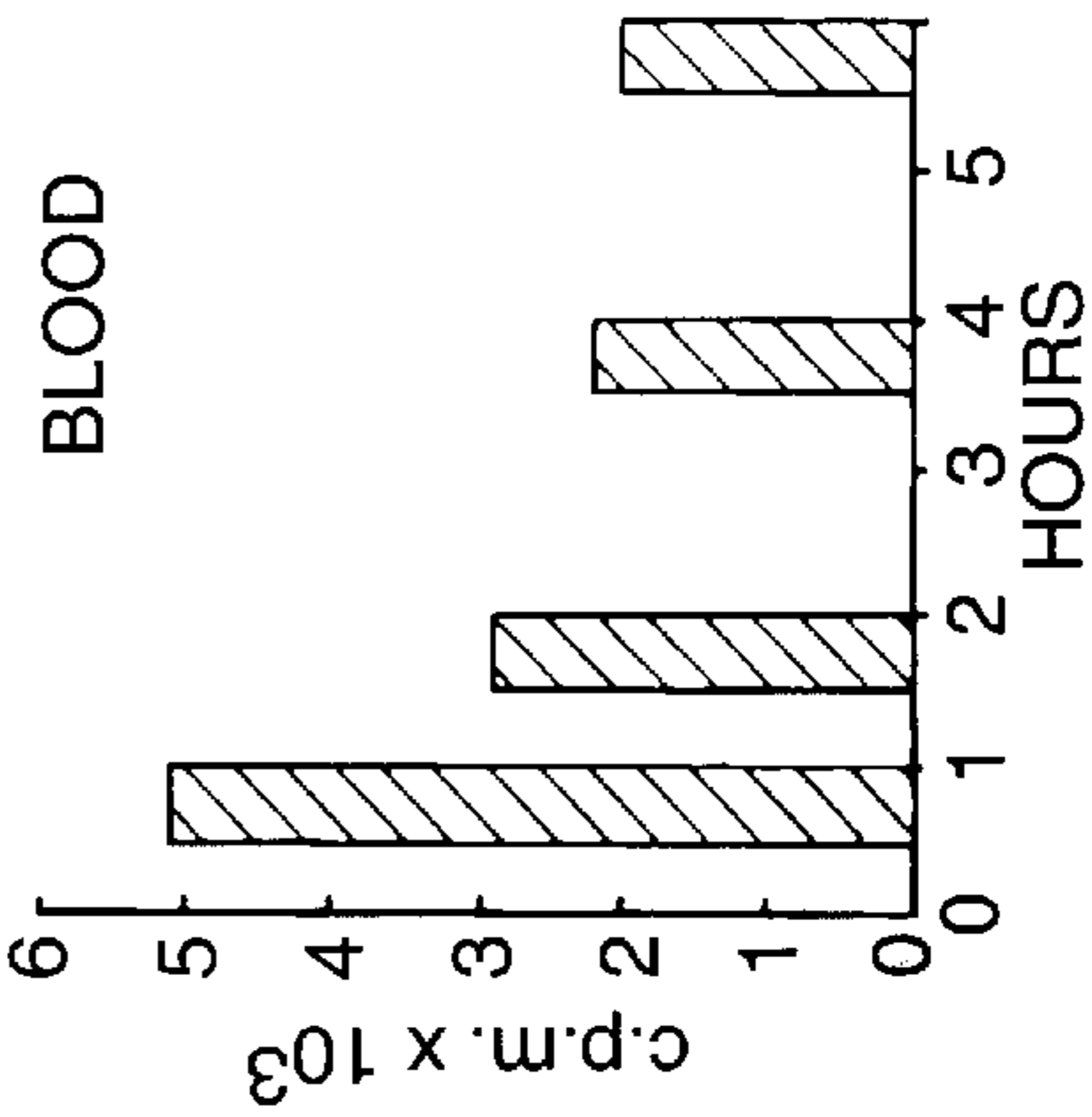


Fig. 1D

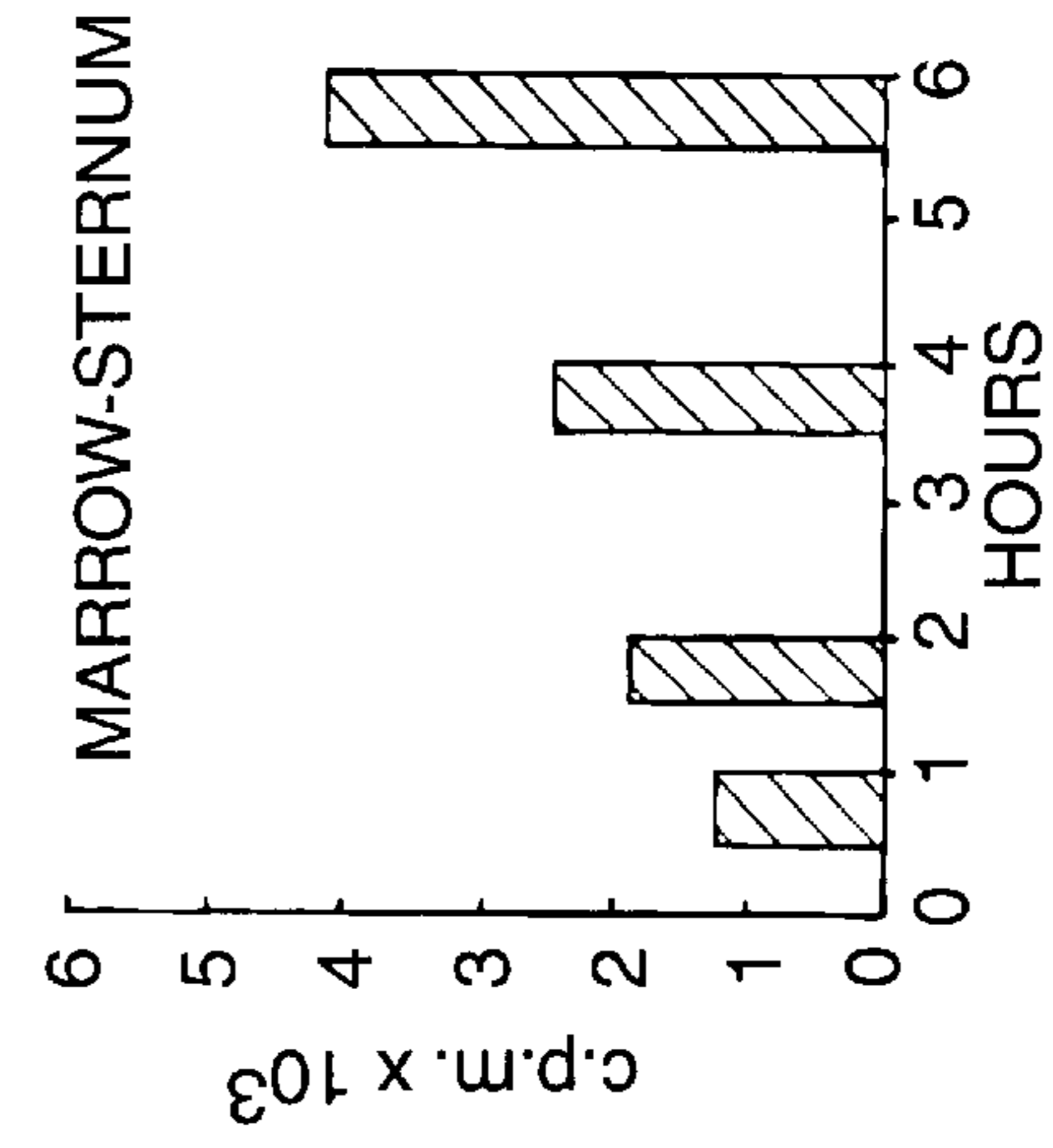


Fig. 1E

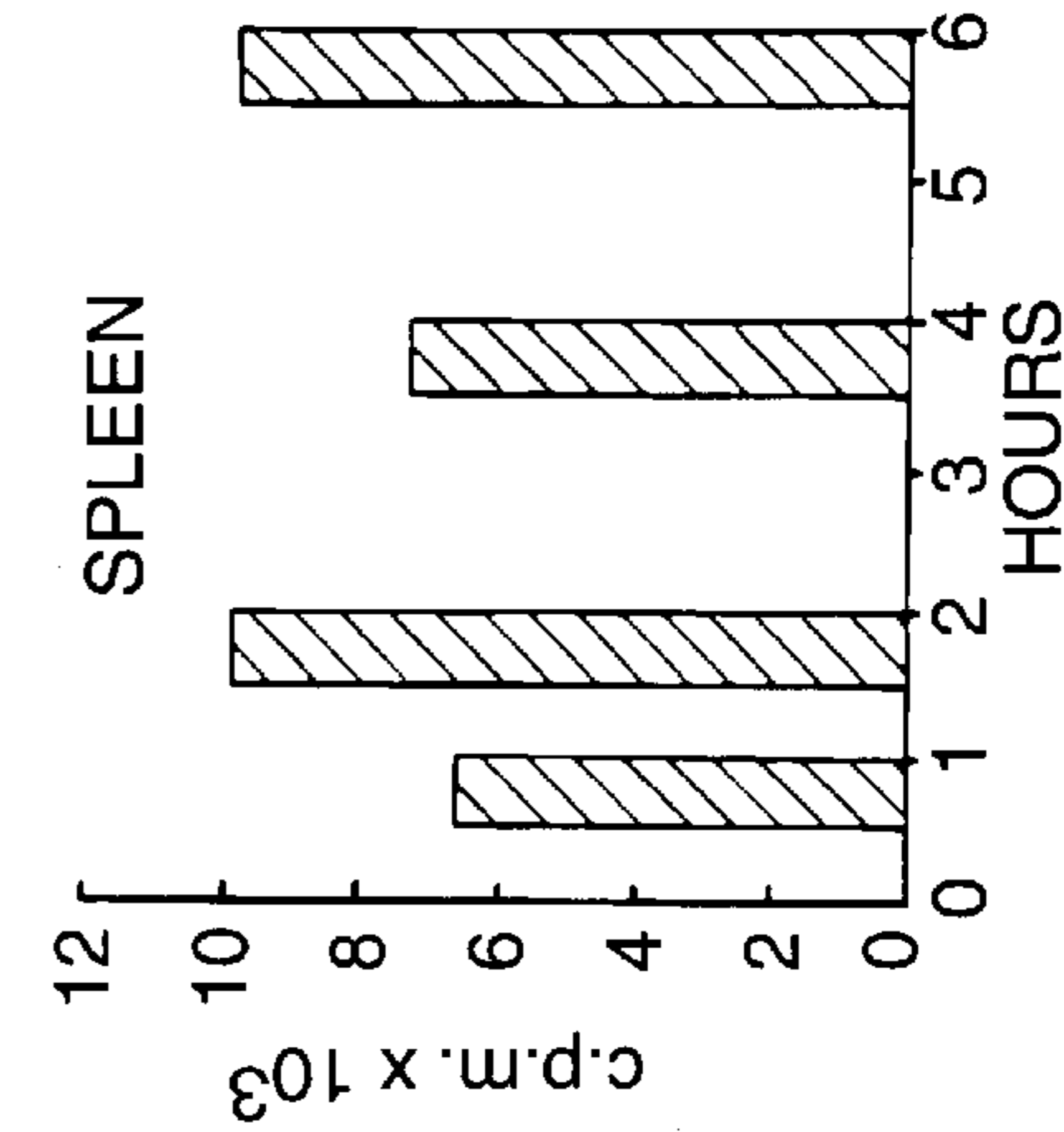


Fig. 1F

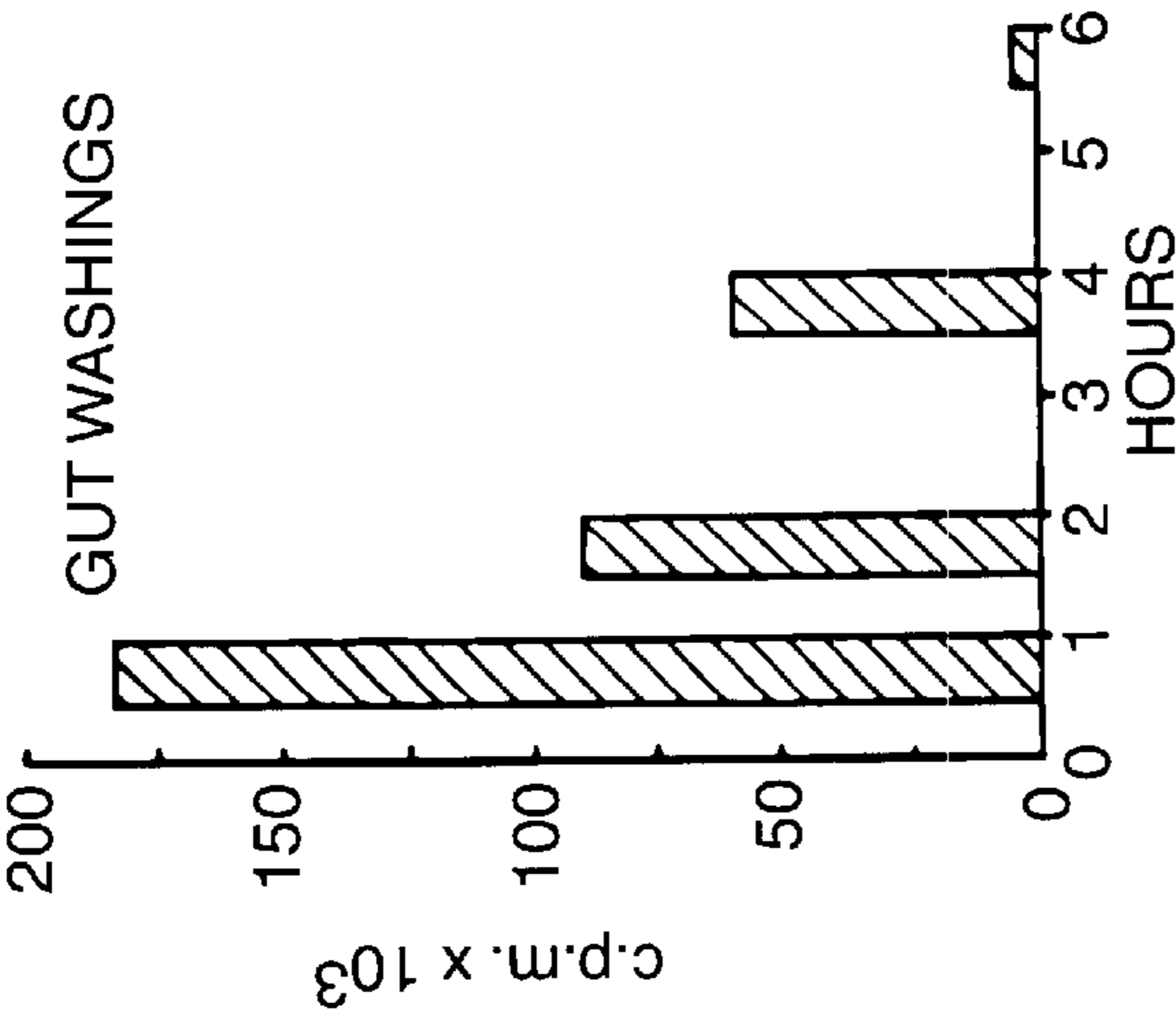


Fig. 2A

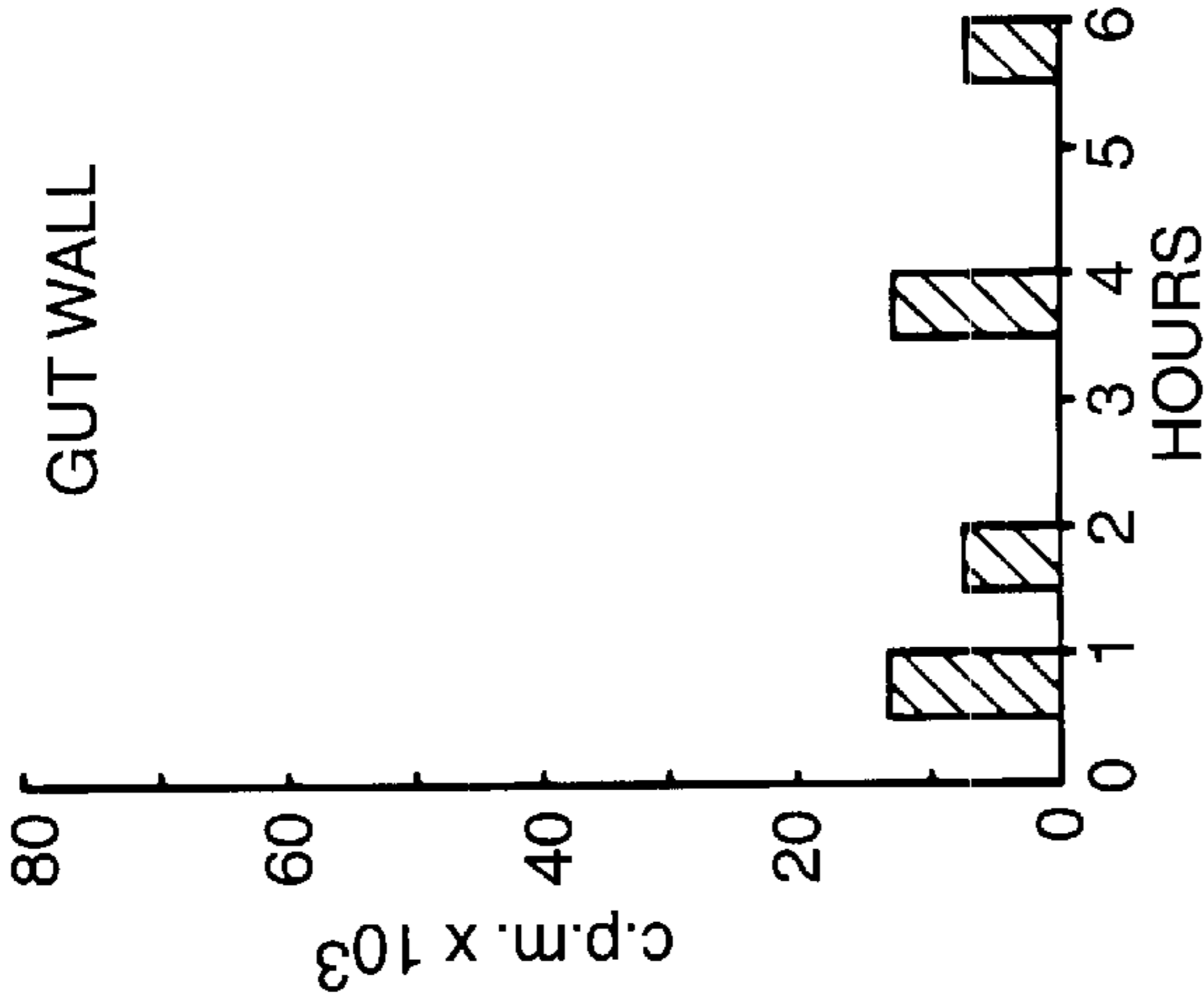


Fig. 2B

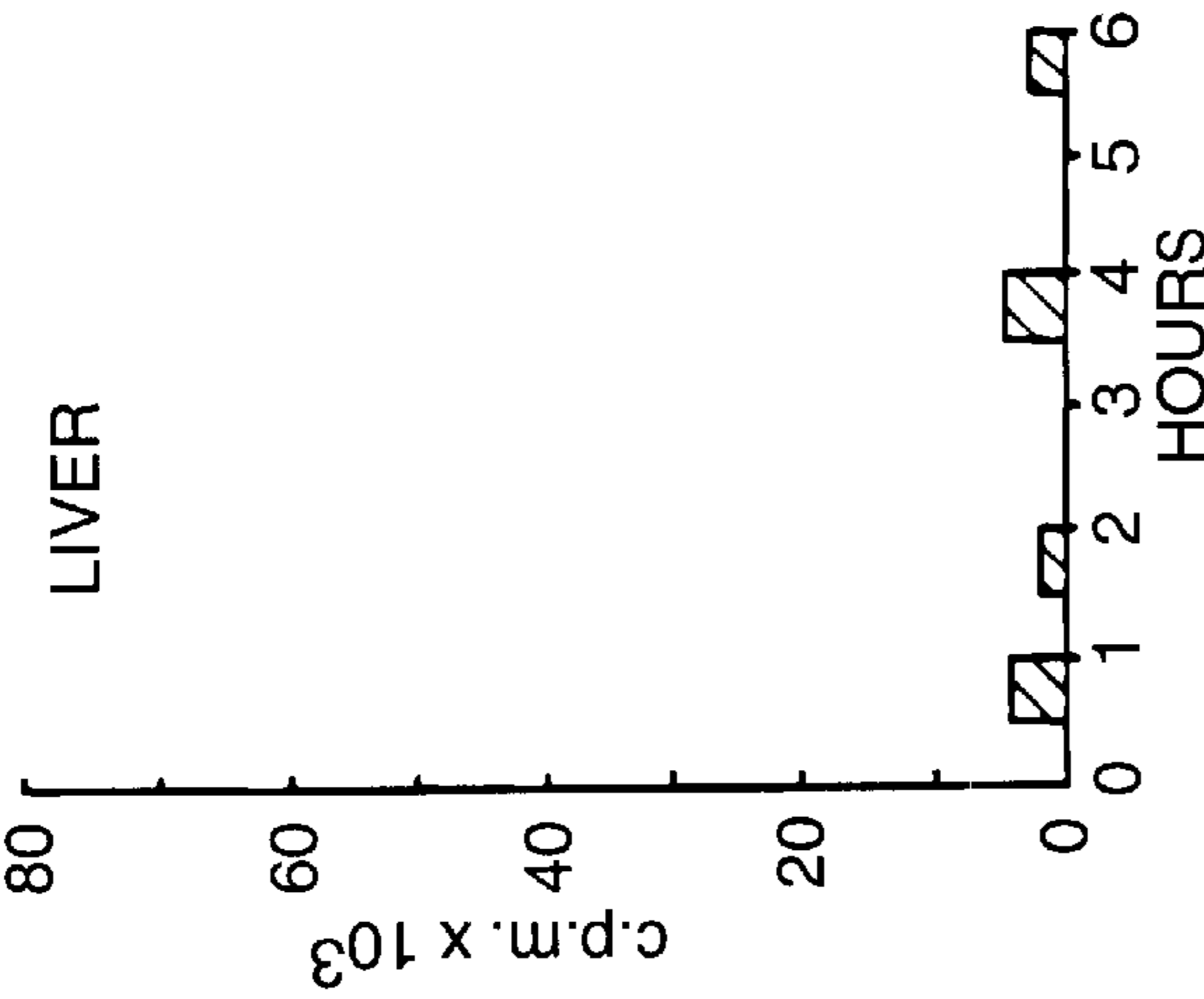


Fig. 2C

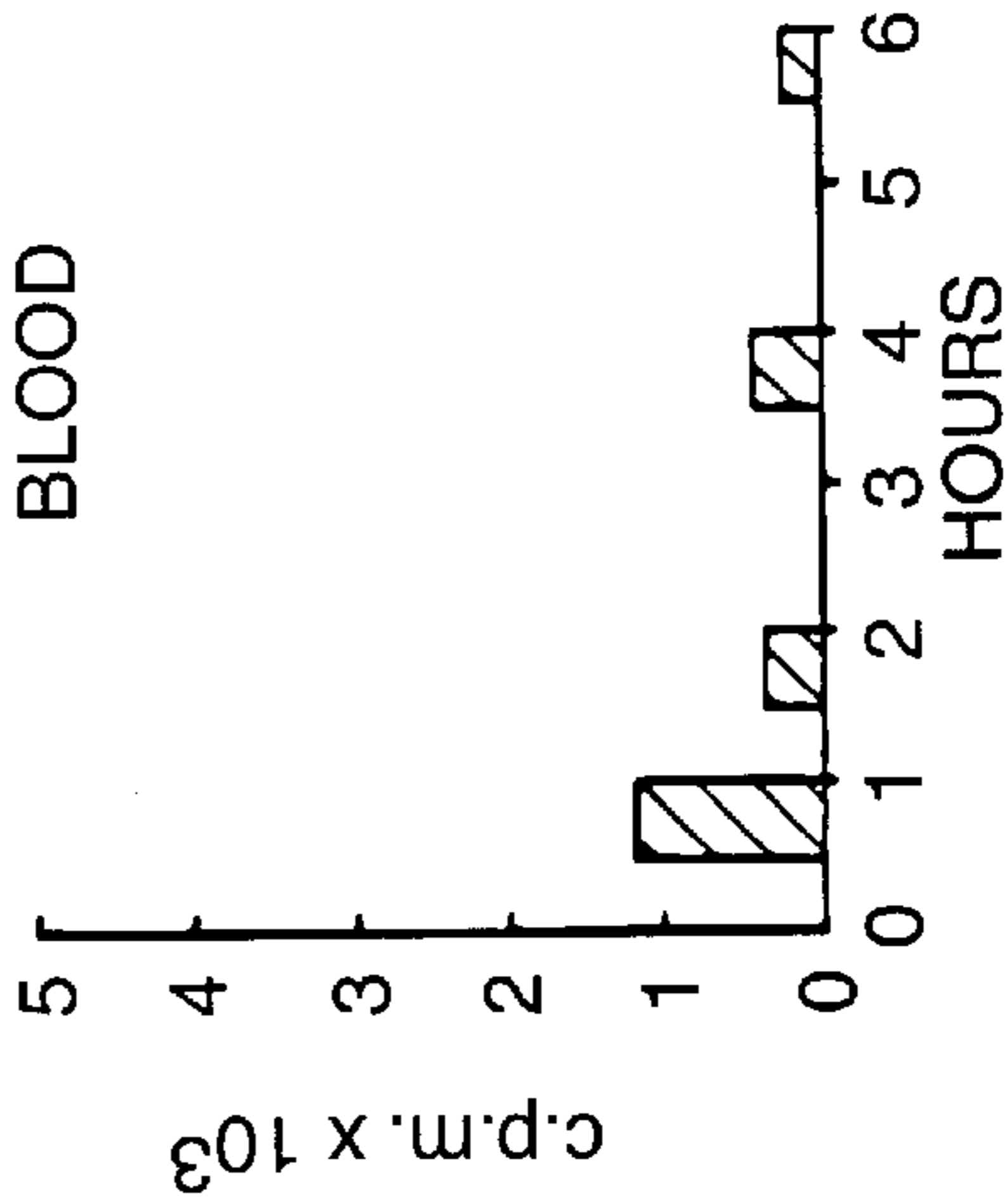


Fig. 2D

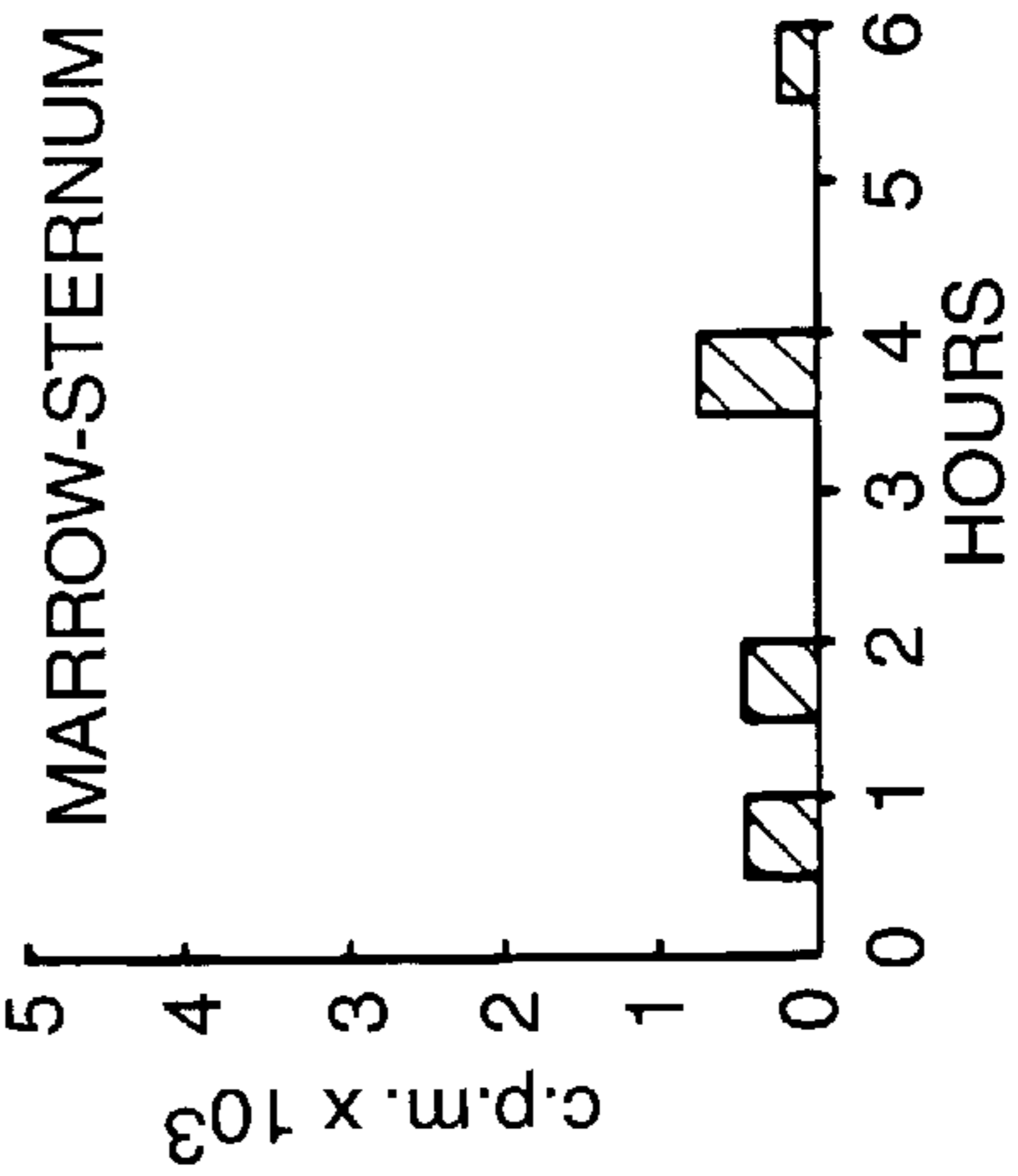


Fig. 2E

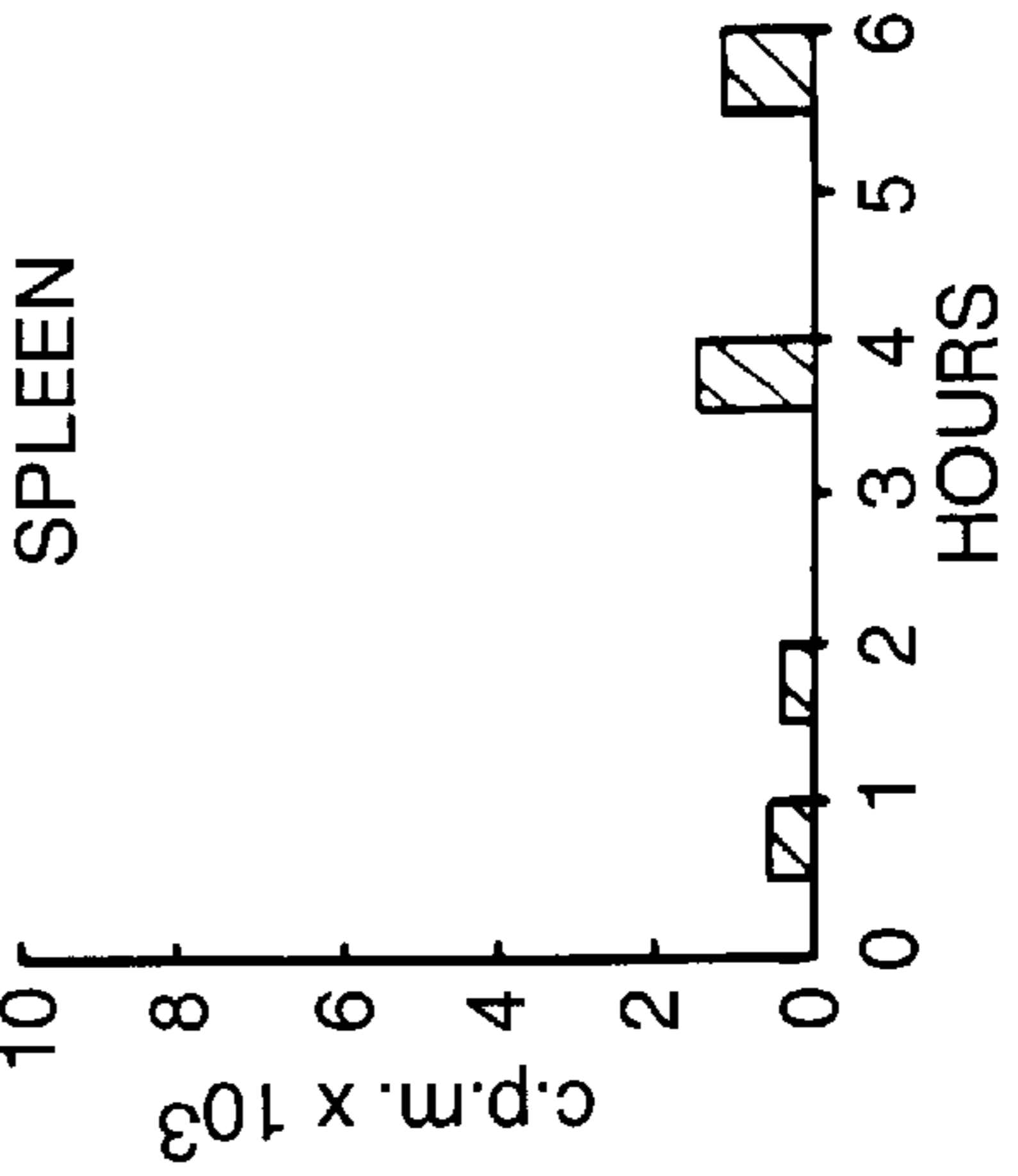
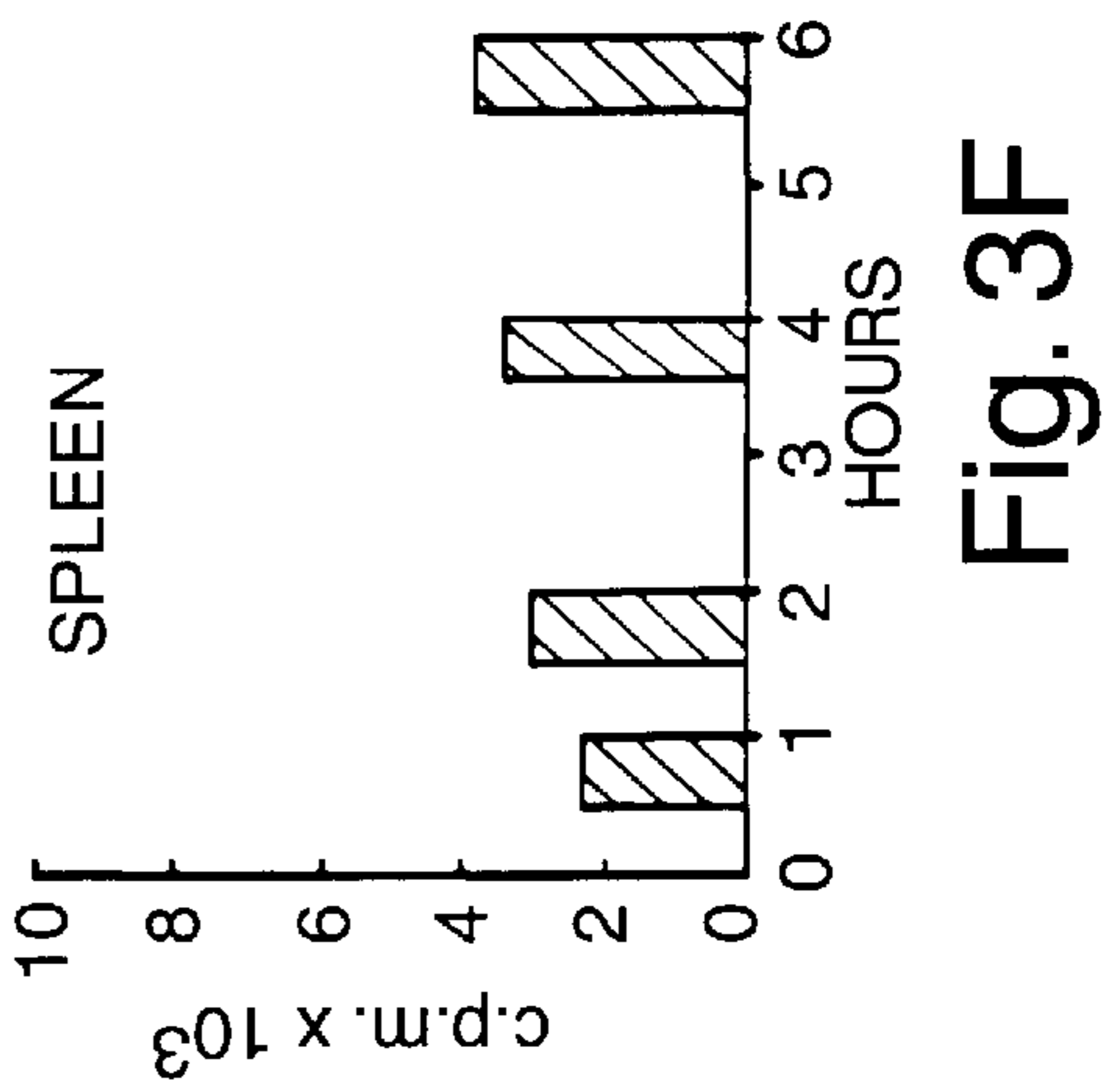
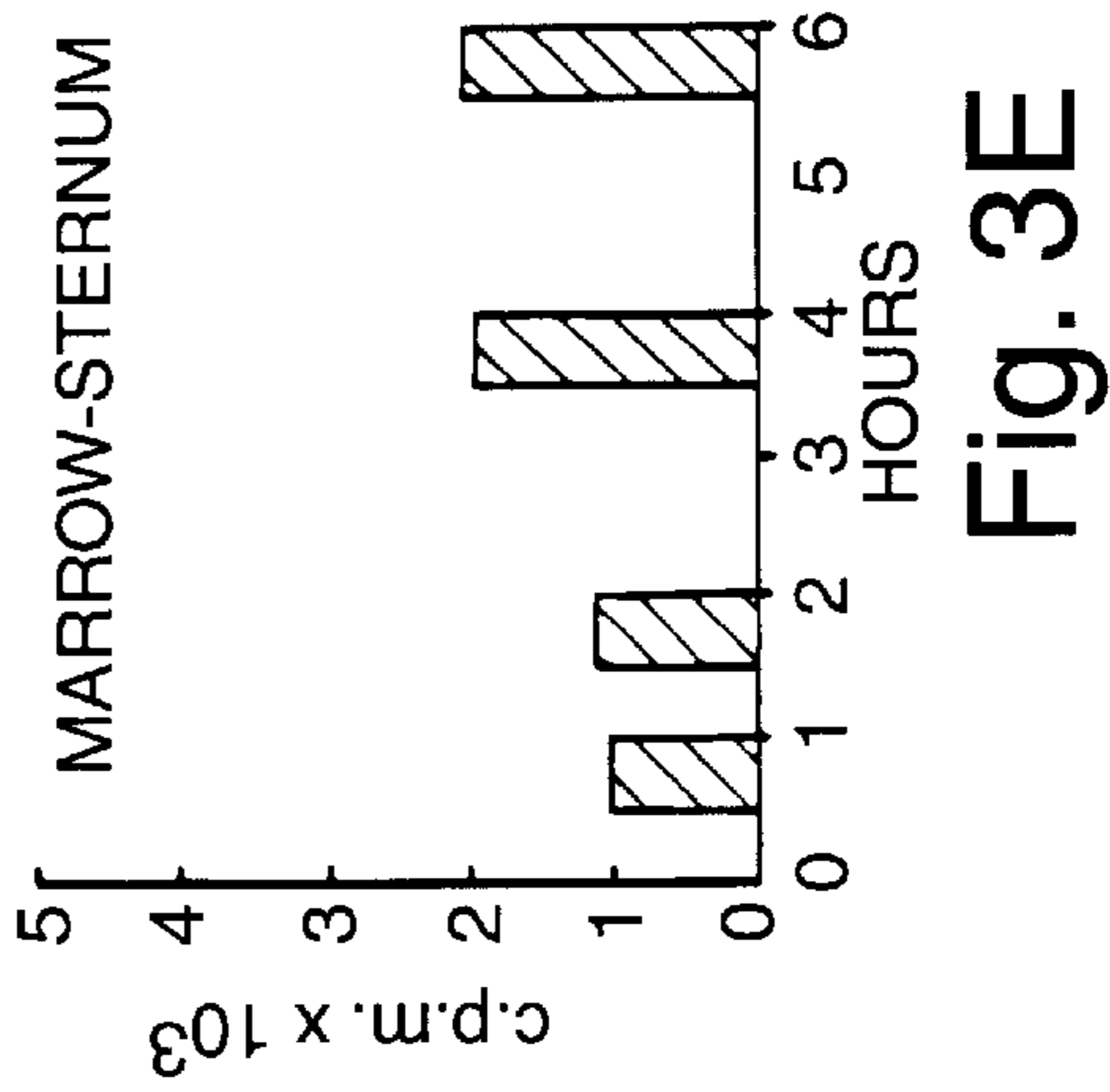
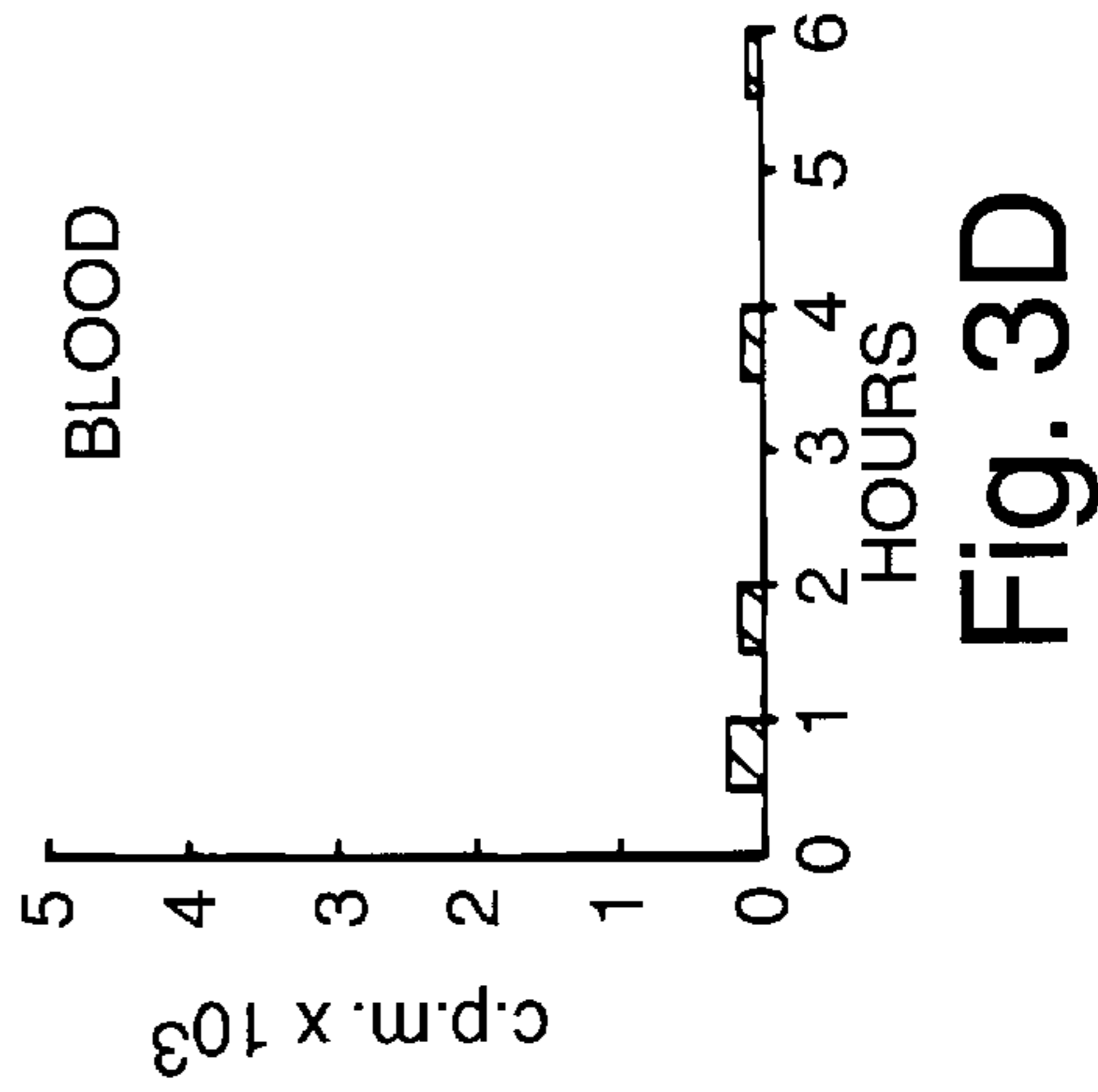
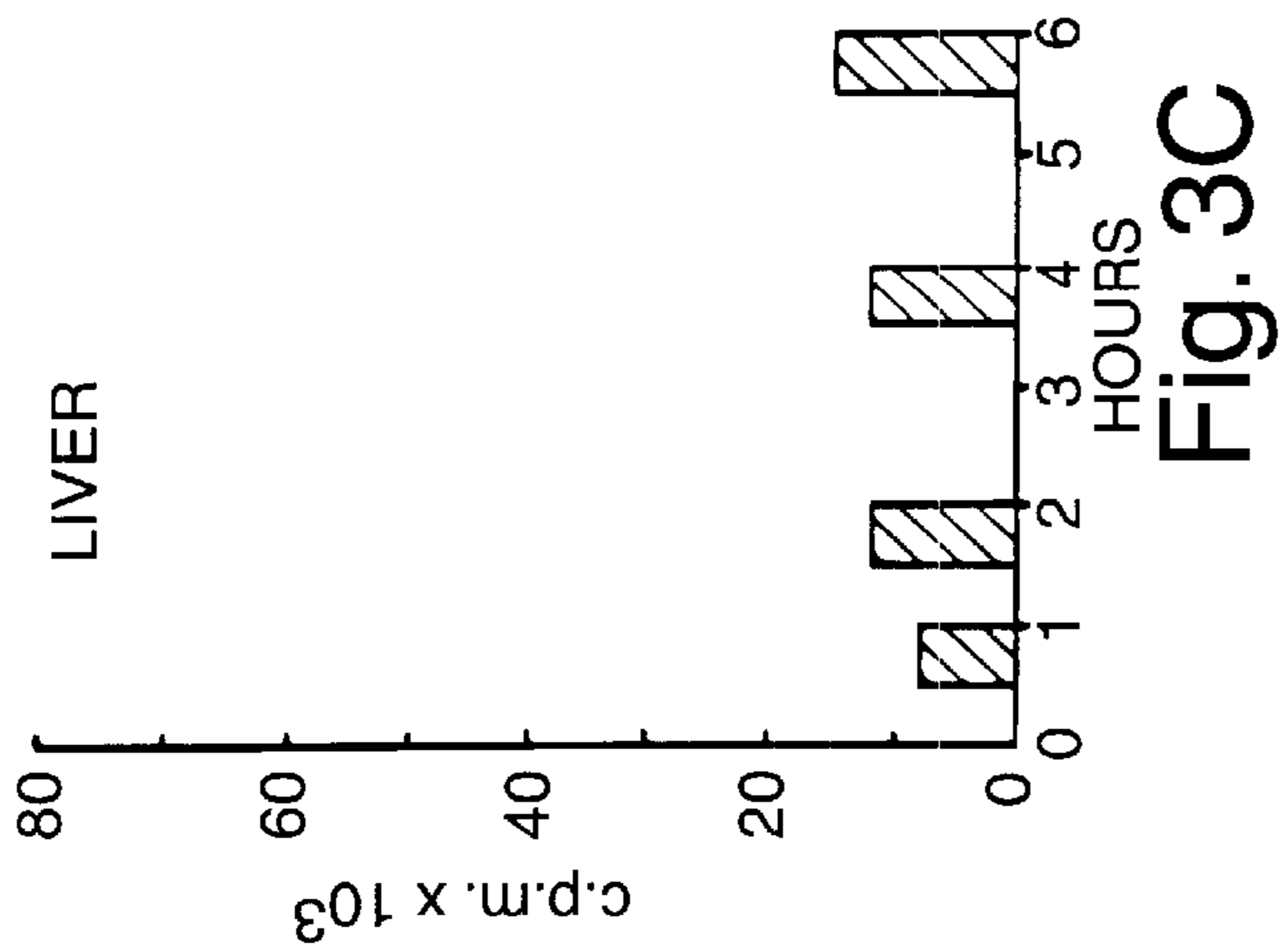
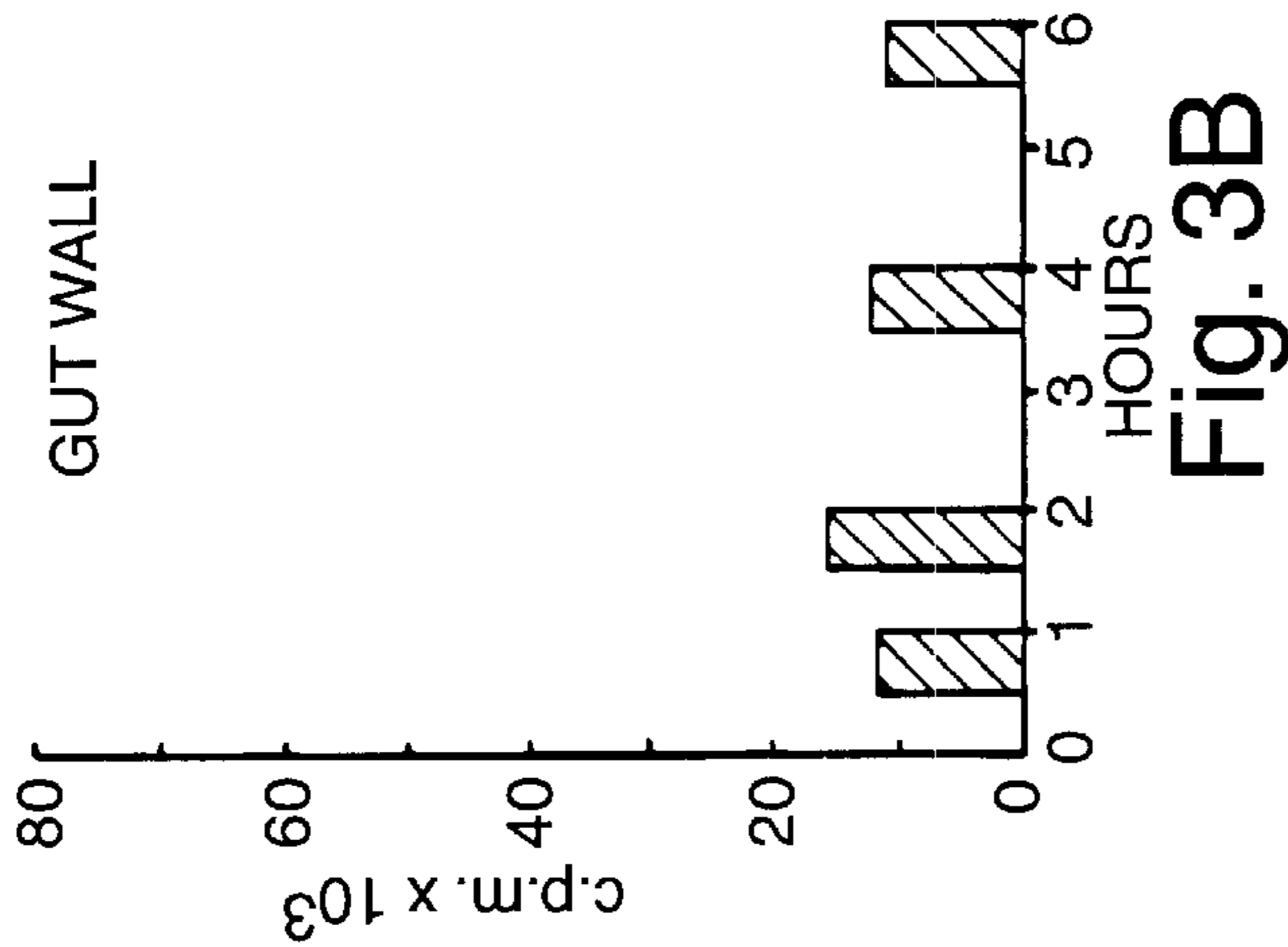
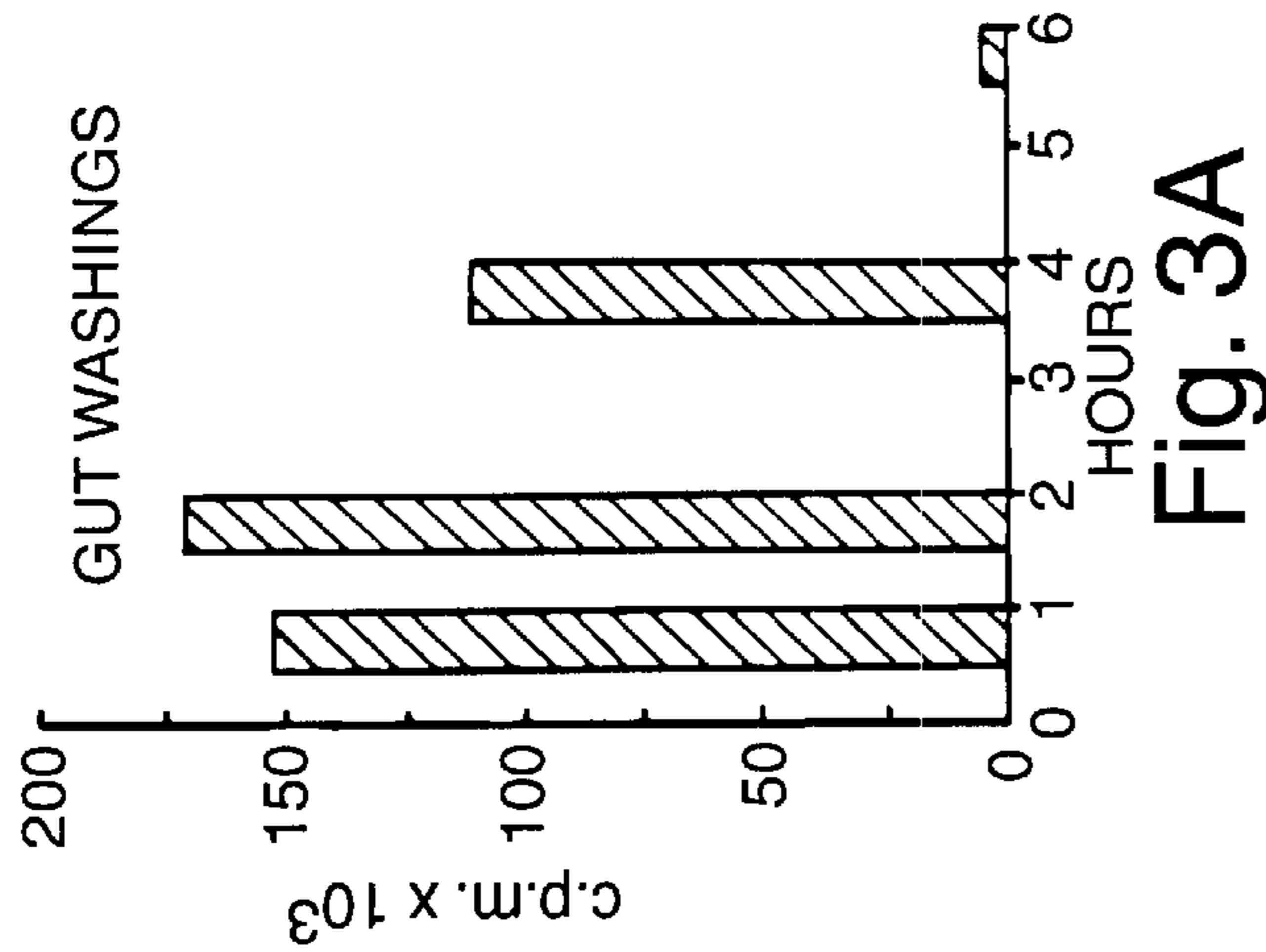
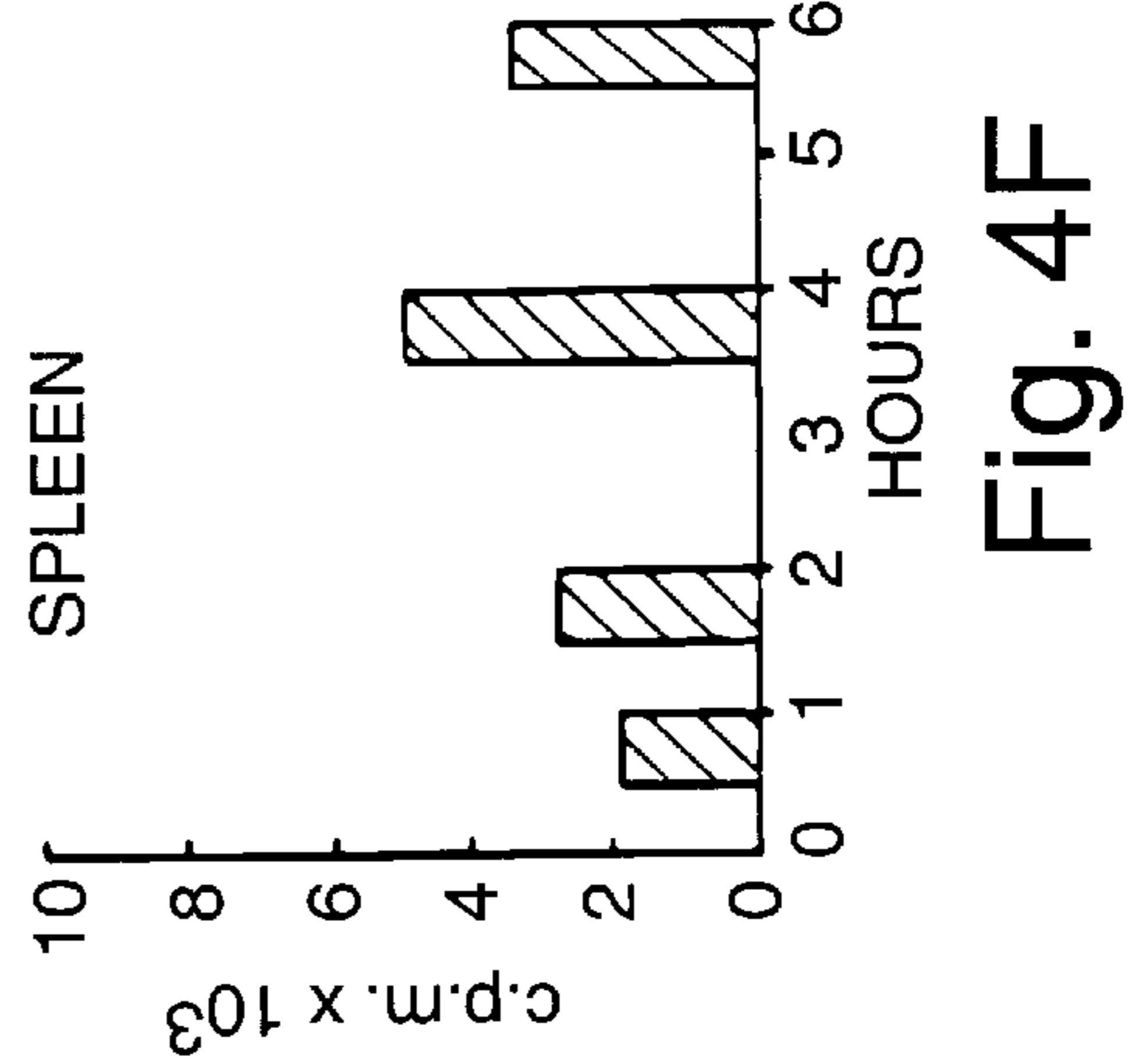
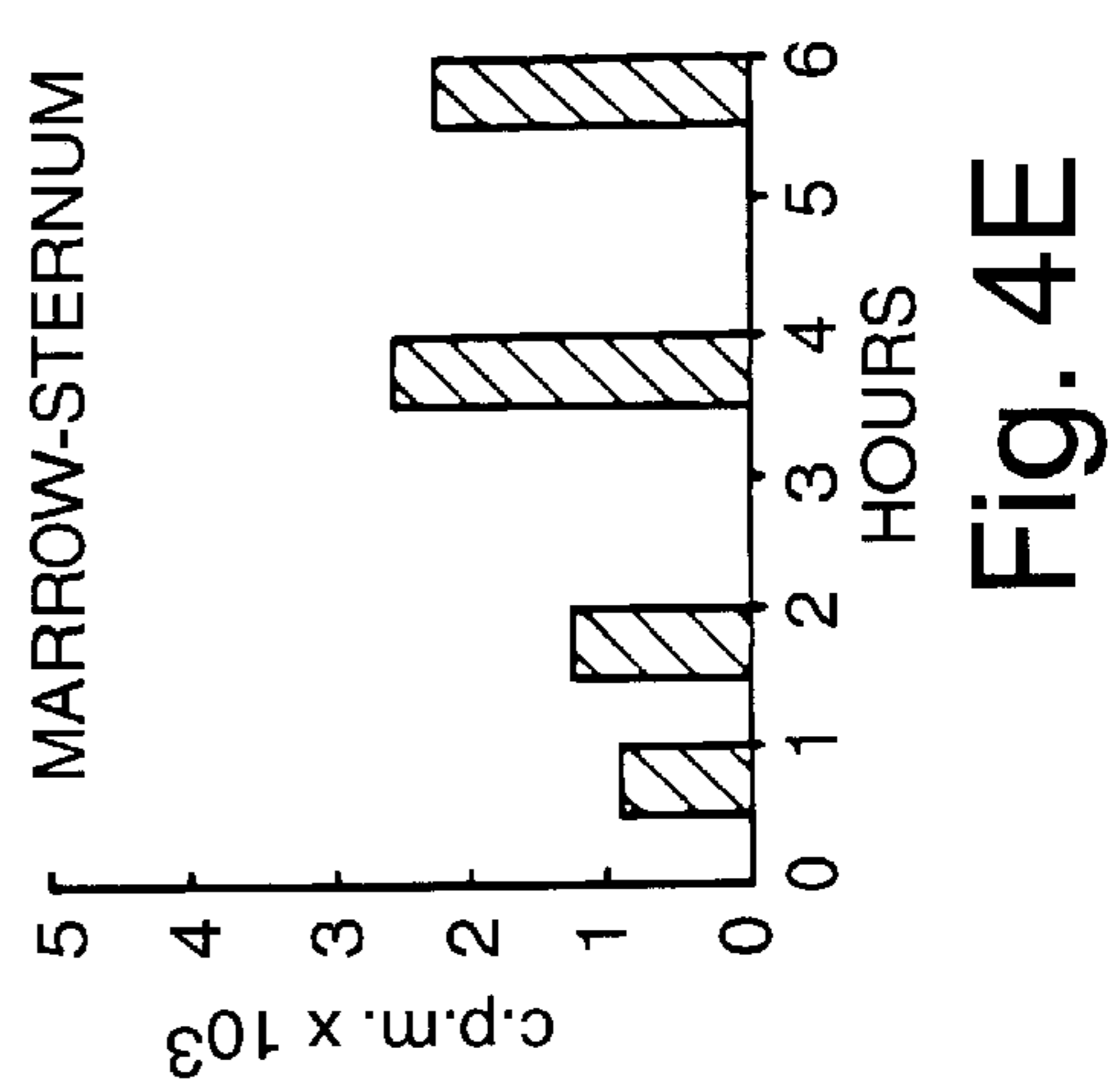
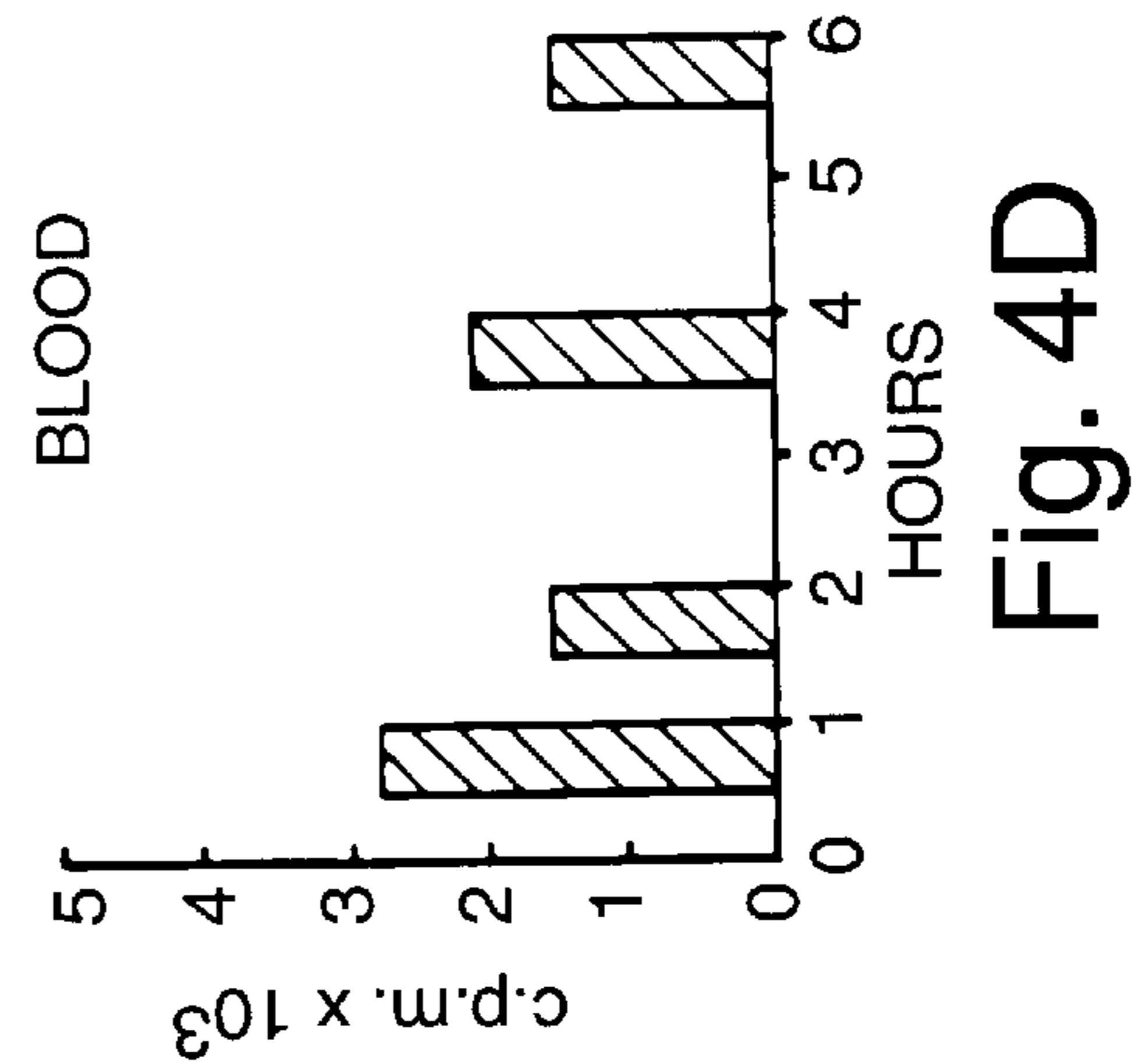
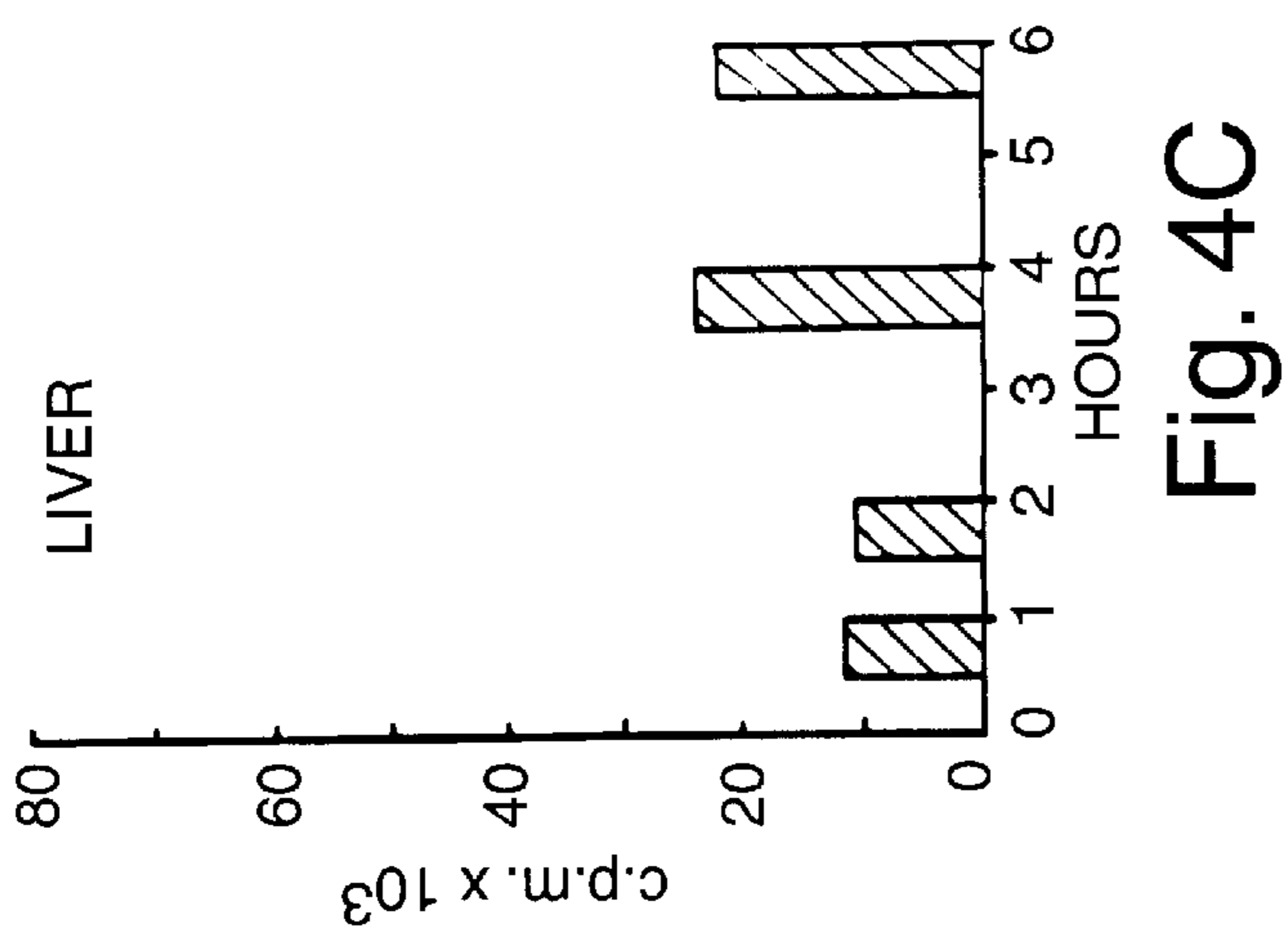
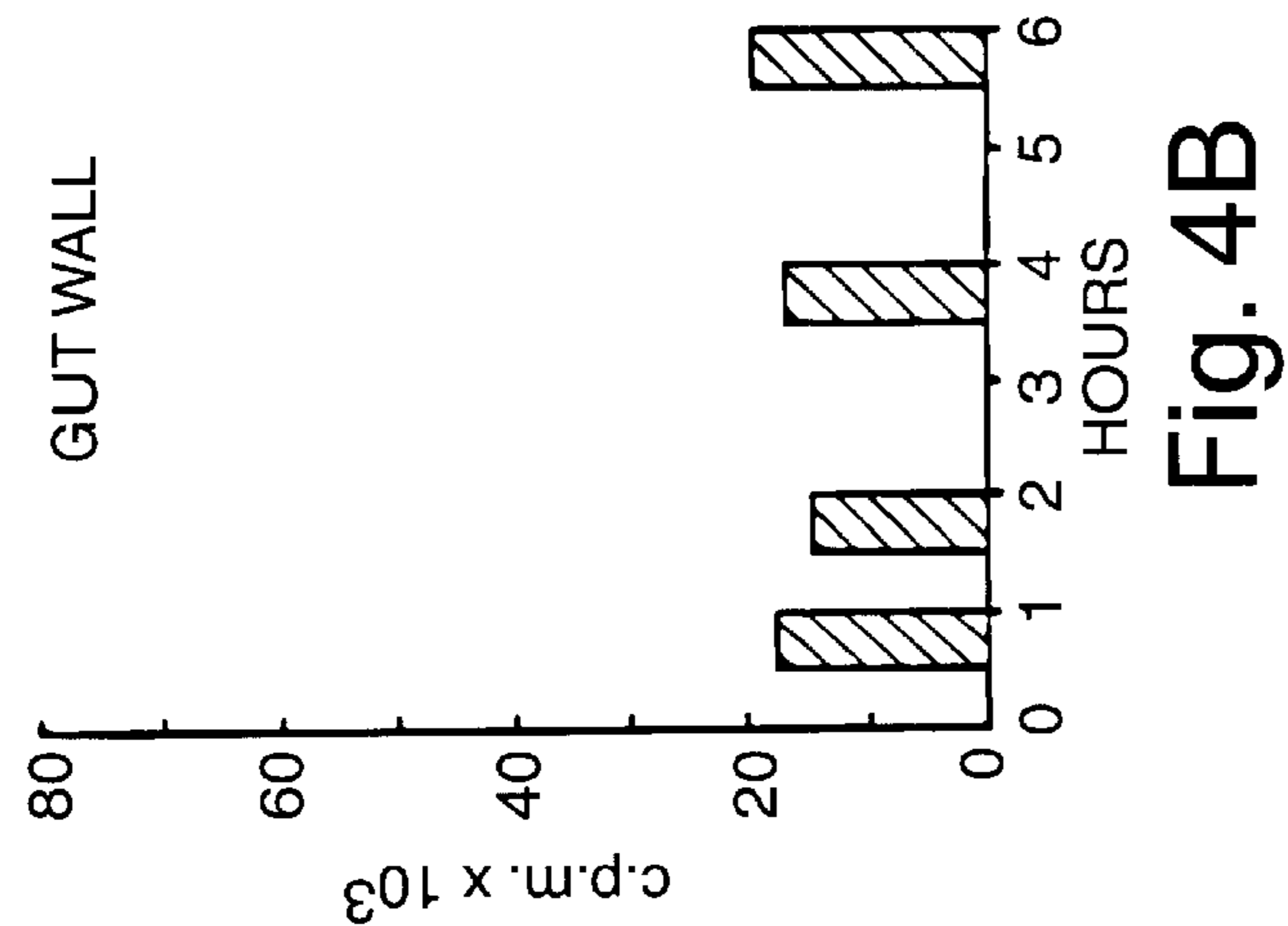
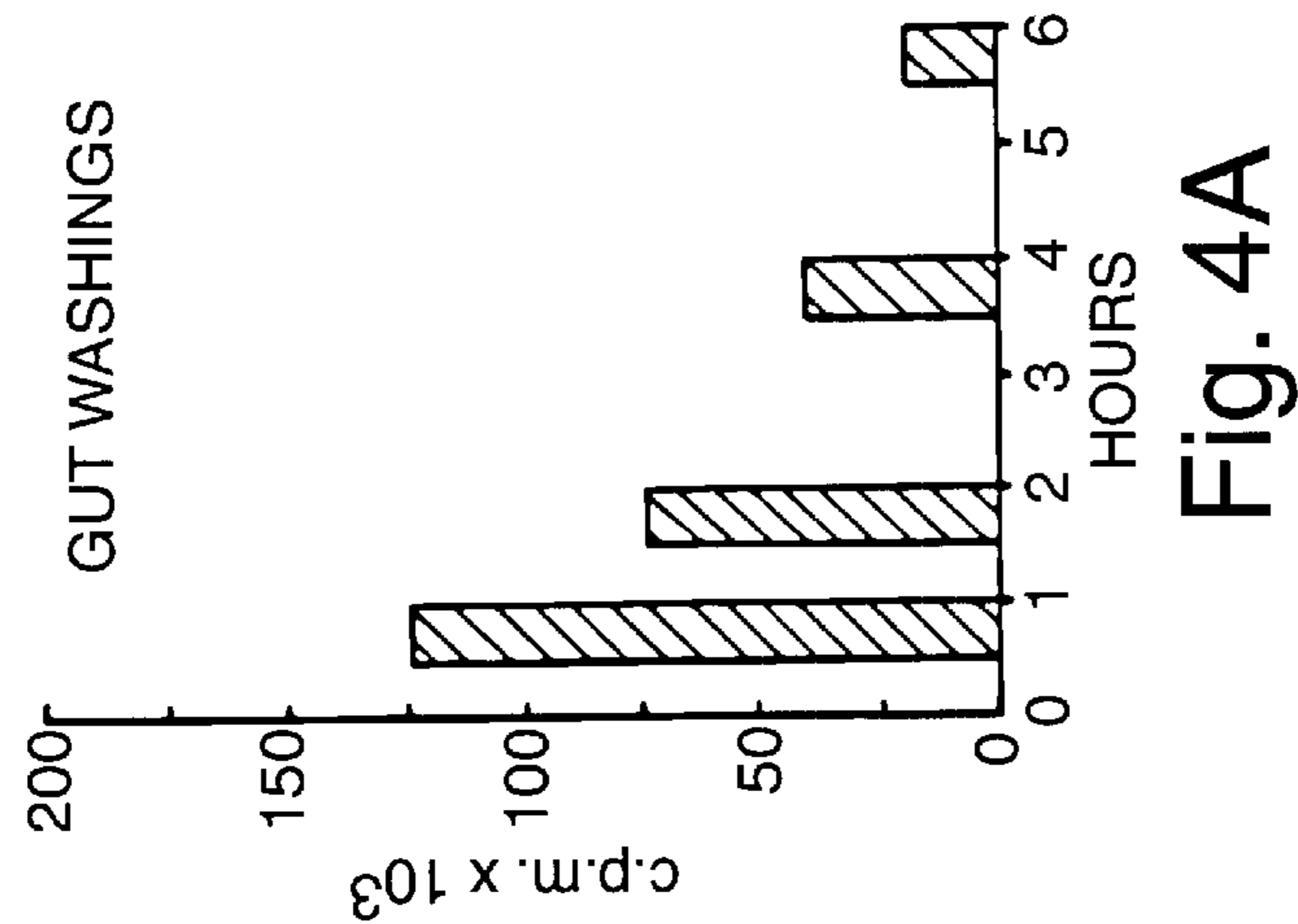


Fig. 2F





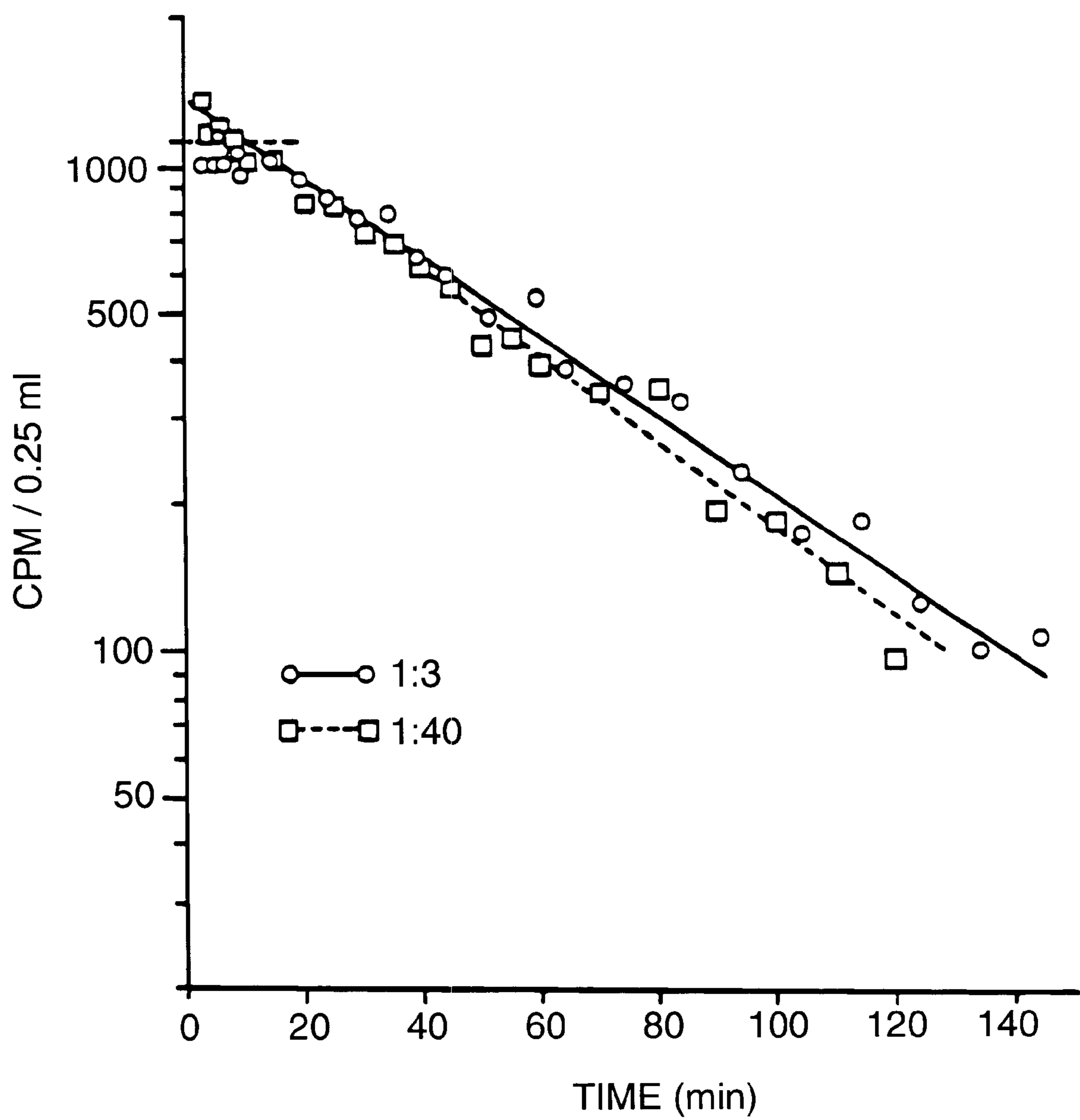


Fig. 5

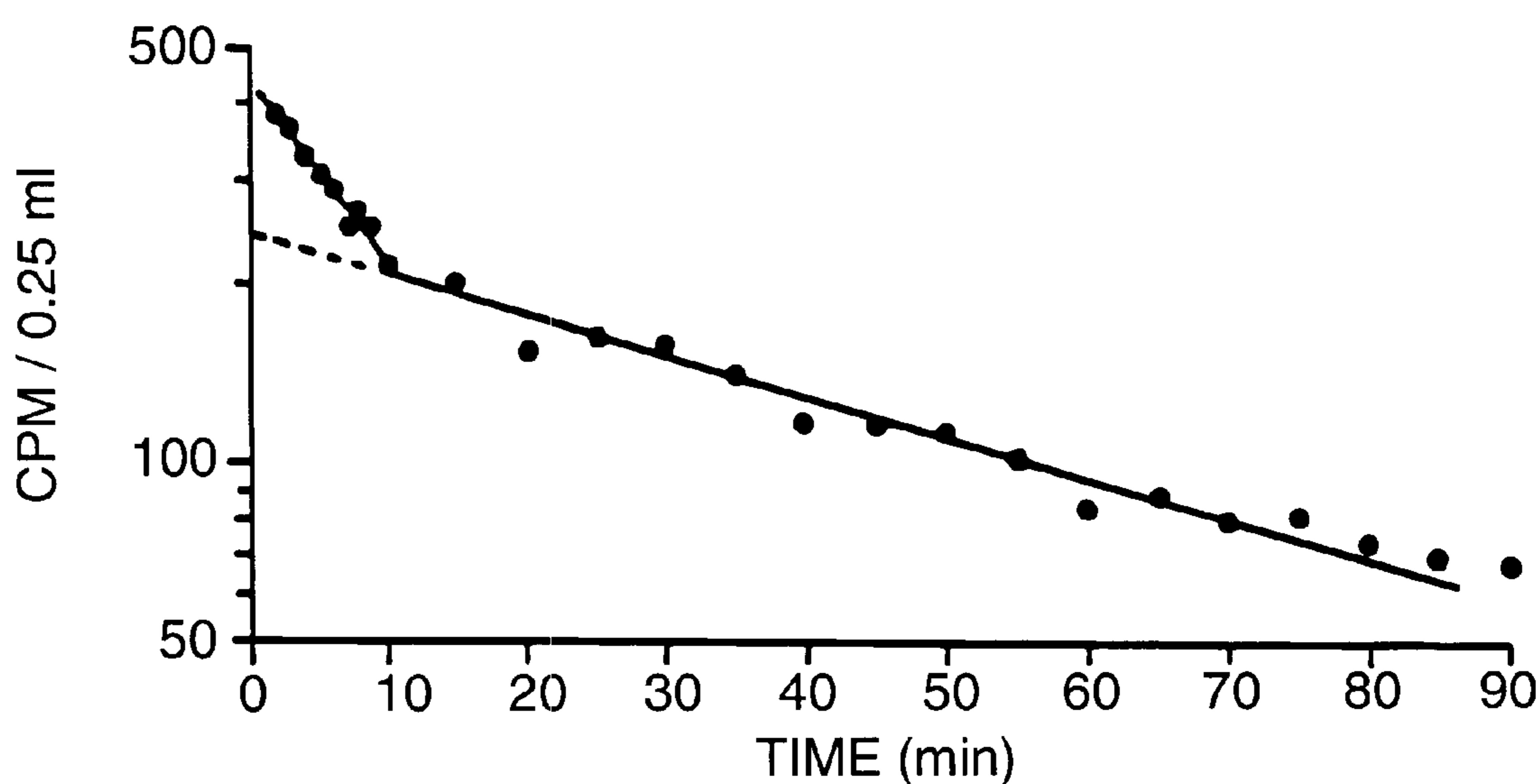


Fig. 6

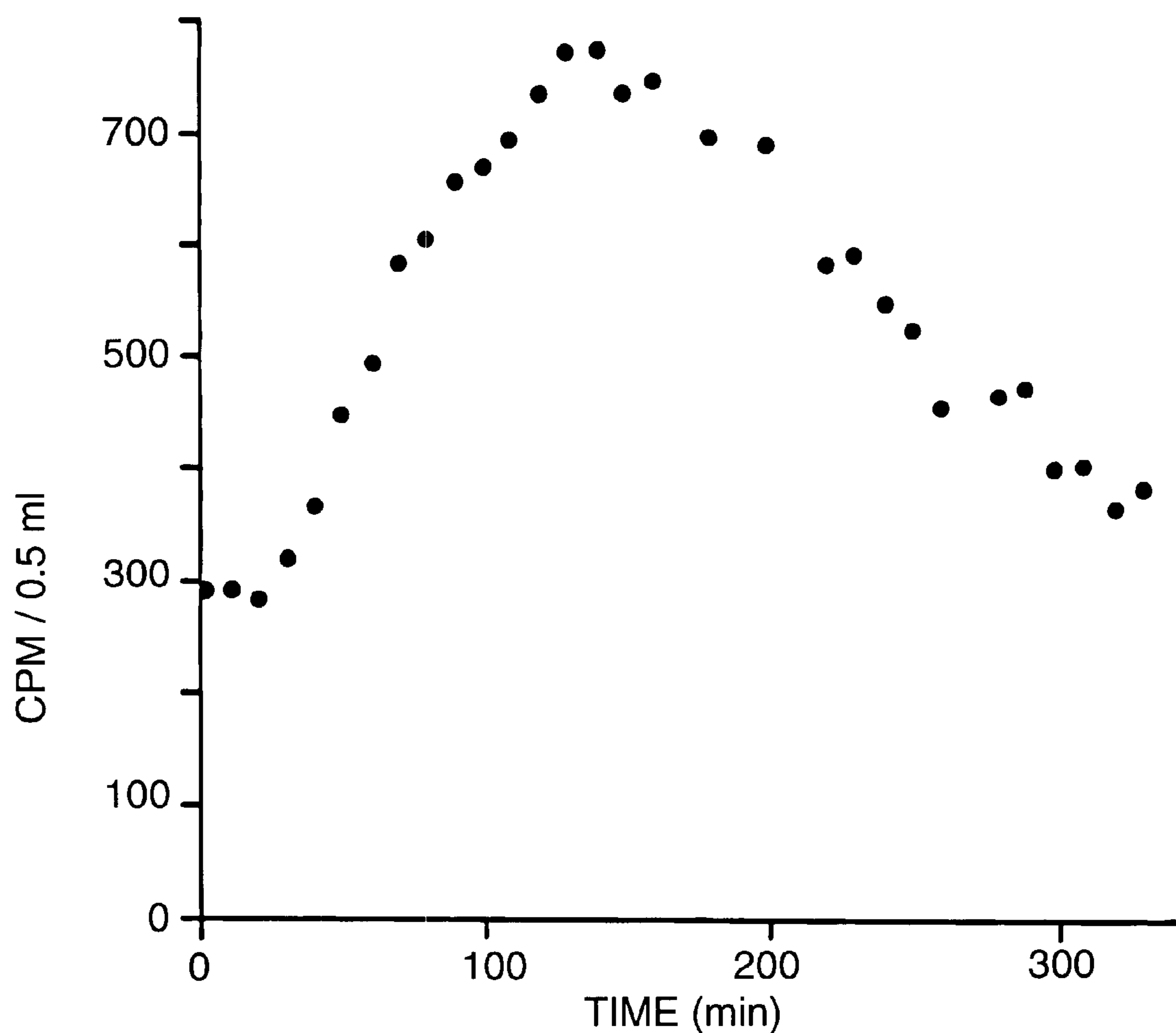


Fig. 7

PHARMACEUTICAL COMPOSITIONS

Matter enclosed in heavy brackets [] appears in the original patent but forms no part of this reissue specification; matter printed in italics indicates the additions made by reissue.

This a rule 62 continuation of application Ser. No. 08/521,281, filed Aug. 30, 1995, now abandoned, which is a Rule 62 continuation of application Ser. No. 08/064,800, filed May 21, 1993, now abandoned, which is a Rule 62 continuation of application Ser. No. 07/708,043, filed May 30, 1991, now abandoned, which is a Reissue application of U.S. Pat. No. 4,834,983, issued May 30, 1989, [This application] which is a continuation of application of Ser. No. 601,485, filed Apr. 18, 1984, now abandoned, which is a continuation-in-part of application Ser. No. 542,972, filed Oct. 18, 1983, now U.S. Pat. No. 4,575,502.

This invention relates to iron compounds for use in pharmaceutical compositions for the treatment of iron deficiency anaemia.

An adequate supply of iron to the body is an essential requirement for tissue growth in both man and animals. Although there is normally an ample amount of iron in the diet, the level of absorption of iron from food is generally low so that the supply of iron to the body can easily become critical under a variety of conditions. Iron deficiency anaemia is commonly encountered in pregnancy and may also present a problem in the newly born, particularly in certain animal species such as the pig. Moreover, in certain pathological conditions there is a mal distribution of body iron leading to a state or chronic anaemia. This is seen in chronic diseases such as rheumatoid arthritis, certain haemolytic diseases and cancer.

Although a wide range of iron compounds is already marketed for the treatment of iron deficiency anaemia, the level of iron uptake by the body of these compounds is often quite low thereby necessitating the administration of relatively high dosage levels of the compound. The administration of high dose, poorly absorbed, iron complexes may cause siderosis of the gut wall and a variety of side effects such as nausea, vomiting, constipation and heavy malodorous stools.

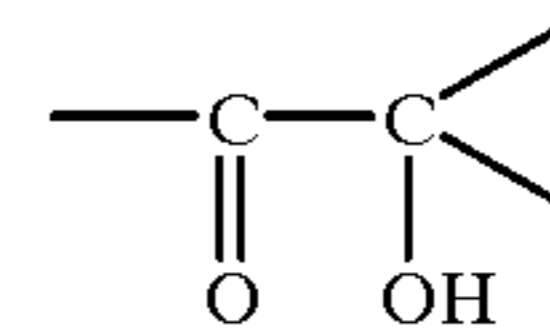
The present invention relates to a group of iron complexes which we have identified as being of particular value for use at relatively low dosage levels in the treatment of iron deficiency anaemia. The hitherto unrecognised value of these complexes in such a context, as shown by in vivo experiments, is unexpected in view of the well known need for improved iron compounds for the treatment of iron deficiency anaemia. This is particularly so as among the compounds whose iron complexes are of the most interest for use in pharmaceutical compositions according to the present invention is a significant number of compounds which are naturally occurring materials, or are readily derivable from such materials, and which have been known for some time to be capable of forming iron complexes. Furthermore, several of these compounds have previously been used in foodstuffs thereby indicating their non-toxic nature and the consequent suitability of pharmaceutical use of their iron complexes.

According to the present invention a pharmaceutical composition comprises an iron complex of 3-hydroxy-4-pyrone or of a 3-hydroxy-4-pyrone in which one or more of hydrogen atoms attached to ring carbon atoms are replaced by an aliphatic hydrocarbon group of 1 to 6 carbon atoms, together with a physiologically acceptable diluent or carrier.

The iron complex present in the pharmaceutical compositions according to the present invention preferably contain iron in the ferric state. Although the use of complexes containing iron in the ferrous state may be considered, such complexes tend to be less stable and are thus of less interest.

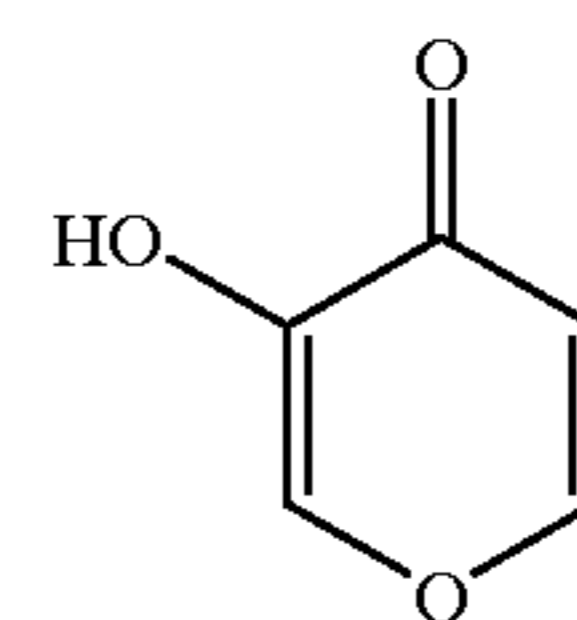
The iron complexes are preferably neutral and this is conveniently achieved by complexing with the iron cation the appropriate number of anions derived from the hydroxypyrrone (through the conversion $\text{OH} \rightarrow \text{O}^-$) necessary to produce neutrality. Preferred iron complexes of use in the present invention are thus of the 3:1 form, containing three hydroxypyrrone anions complexed with a ferric cation. It will be appreciated, however, that the invention does not exclude the use of complexes of the 1:1 or particularly the 2:1 form, although the 3:1 neutral ferric complexes are of especial interest.

The substituted 3-hydroxy-4-pyrones may carry more than one type of aliphatic hydrocarbon group but this is not usual and, indeed, substitution by one rather than two or three aliphatic hydrocarbon groups is preferred. The aliphatic hydrocarbon groups may be cyclic or acyclic, having a branched chain or especially a straight chain in the latter case, and may be unsaturated or especially saturated. Groups of from 1 to 4 carbon atoms are particularly of 1 to 3 carbon atoms are of most interest. Alkyl groups are preferred, for example cyclic groups such as cyclopropyl and especially cyclohexyl but, more particularly preferred are acyclic alkyl groups such as n-propyl and isopropyl, and especially ethyl and methyl. Substitution at the 2- or 6-position is of especial interest although, when the ring is substituted by the larger aliphatic hydrocarbon groups, there may be an advantage in avoiding substitution on a carbon atom alpha to the

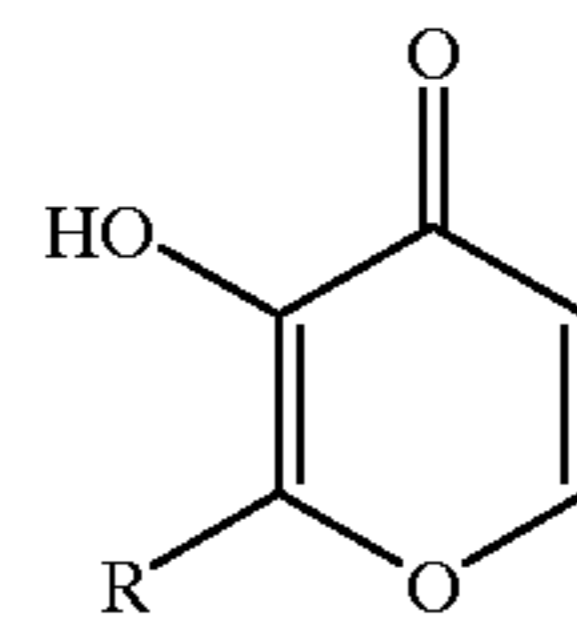


system. The system is involved in the complexing with iron and the close proximity of one of the larger aliphatic hydrocarbon groups may lead to steric effects which inhibit complex formation.

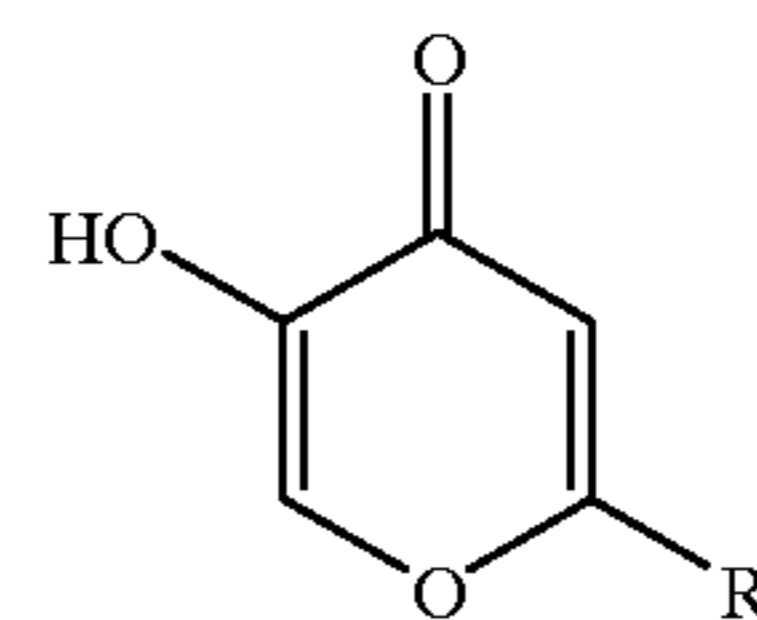
Examples of specific compounds whose iron complexes may be used in compositions according to the present invention are shown by the following formulae (I), (II) and (III);



(I)



(II)



(III)

in which R is an alkyl group, for example methyl, ethyl, n-propyl or isopropyl. Among these compounds 3-hydroxy-2-methyl-4-pyrone (maltol; II, $\text{R}=\text{CH}_3$) is of most interest, whilst 3-hydroxy-4-pyrone (pyromeconic acid; I), 3-hydroxy-6-methyl-4-pyrone (isomaltol; III, $\text{R}=\text{CH}_3$) and

particularly 2-ethyl-3-hydroxy-4-pyrone (ethylpyromeconic acid; II, $R=CH_2H_5$) are also of especial interest.

In the case of certain of the hydroxypyrones referred to above, for example maltol, ethylpyromeconic acid and isomaltol, the formation of an iron complex of the compound has been referred to in the literature, although it should be noted that the procedures described in the literature for the production of such complexes often would not provide complexes of a form which is preferred for use in the pharmaceutically compositions according to the present invention. In the case of other hydroxypyrones, the iron complexes are novel and are included, per se, by the present invention.

The iron complexes are conveniently prepared by the reaction of the hydroxypyrene and iron ions, the latter conveniently being derived from an iron salt, particularly a ferric halide and especially ferric chloride. The reaction is conveniently effected in a suitable mutual solvent and water may often be used for this purpose. If desired, however, an aqueous/organic solvent mixture may be used or an organic solvent, for example ethanol, methanol or chloroform and mixtures of these solvents together and/or with water where appropriate. In particular, methanol or especially ethanol may be used where it is desired to effect the separation of at least a major part of a by-product such as sodium chloride by precipitation whilst the iron complex is retained in solution.

It should be appreciated that the nature of the iron complex obtained by the reaction of a hydroxypyrene and iron ions will depend both on the proportion of these two reactants and upon the pH of the reaction medium. Thus, for the preparation of the 3:1 ferric complex, for example, the hydroxypyrene and the ferric salt are conveniently mixed in solution in a 3:1 molar proportion and the pH adjusted to a value in the range of 6 to 9, for example 7 or 8. If a similar excess of hydroxypyrene:iron is employed but no adjustment is made of the acidic pH which results on the admixture of the hydroxypyrene and an iron salt such as ferric chloride, then a mixture of the 2:1 and 1:1 complex will instead be obtained.

Reaction to form the iron complex is generally rapid and will usually have proceeded substantially to completion after 5 minutes at about 20° C., although a longer reaction time may be used if necessary. Following separation of any precipitated by-product, such as sodium chloride in the case of certain solvent systems, the reaction mixture may conveniently be evaporated on a rotary evaporator or freeze dried to yield the solid iron complex. This may, if desired, be crystallised from a suitable solvent, for example water, an alcohol such as ethanol, or a solvent mixture, including mixtures containing ether. The present invention further includes a process for the preparation of an iron complex of 3-hydroxy-4-pyrone or of a 3-hydroxy-4-pyrone in which a hydrogen atom attached to one or more ring carbon atoms is replaced by an aliphatic hydrocarbon group of 1 to 6 carbon atoms, which comprises reacting said hydroxypyrene with iron ions and isolating the resultant complex; such a process being restricted, however, to exclude processes for the preparation of those particular forms of iron complex of particular hydroxypyrones which have already been described in the literature.

Whilst for some uses it may be appropriate to prepare the iron complex in substantially pure form, i.e. substantially free from by-products of manufacture, in other cases, for example with a solid oral formulation as described hereinafter, the presence of by-products such as sodium chloride may be quite acceptable. In general, however, the

neutral 3:1 hydroxypyrene:iron(III) complex is of particular interest in a form which is substantially free at least from those by-products which are complexes containing different proportions of hydroxypyrene and iron, in particular the 2:1 and 1:1 complexes. Accordingly the present invention includes an iron complex, for example the 3:1 hydroxypyrene:iron(III) complex, of 3-hydroxy-4-pyrone or of a 3-hydroxy-4-pyrone in which one or more of the hydrogen atoms attached to ring carbon atoms are replaced by an aliphatic hydrocarbon group of 1 to 6 carbon atoms, but excluding 3-hydroxy-2-methyl-4-pyrone, when in a form substantially free from iron complexes of the pyrene containing other proportions of iron. As indicated hereinafter, it may be advantageous under some circumstances for the iron complex to be used in admixture with the free pyrene. It is possible to produce such a mixture by mixing the two components either in the solid form or in solution, followed by isolation of a solid mixture in the latter case when a solid composition is required. However, it may be more convenient to obtain such a mixture by reacting a molar proportion of the pyrene and iron ions of greater than 3:1. It should be stressed, however, that the conditions as well as the proportion of reactants used in the reaction are of importance if a mixture of the free pyrene and the preferred neutral 3:1 complex is to be obtained. In particular, as indicated previously, the pH of the reaction mixture is particularly important and, because of this fact, certain prior art procedures concerned with the use of iron pyrene complexes in food coloring, for example as described in U.S. Pat. No. 4,018,907, substantially fail to yield the 3:1 complex even though an excess of the pyrene is present, owing to the lack of pH control.

Certain hydroxypyrones, such as maltol, are available commercially. With others, a convenient starting material in many instances consists of pyromeconic acid which is readily obtainable by the decarboxylation of meconic acid. Thus, for example, pyromeconic acid may be reacted with an aldehyde to insert a 1-hydroxy-alkyl group at the 2-position, which group may then be reduced to produce a 2-alkyl-3-hydroxy-4-pyrone. The preparation of 2-ethyl-3-hydroxy-4-pyrone, etc., by this route is described in U.S. application Ser. No. 310,141 (series of 1960).

It will be appreciated that these are not the only routes available to these compounds and their iron complexes and that various alternatives may be used as will be apparent to those skilled in the art.

The iron complexes may be formulated for use as pharmaceuticals for veterinary or human use by a variety of methods. For instance, they may be applied as an aqueous, oily or emulsified composition incorporating a liquid diluent, which composition most usually will be employed for parenteral administration, for example intramuscularly, and therefore may conveniently be sterile and pyrogen free. Oral administration is, however, generally to be preferred for the treatment of iron deficiency anaemia in humans and the complexes of the present invention may be given by such a route. Although compositions incorporating a liquid diluent may be used for oral administration, it is preferred to use compositions incorporating a solid carrier, for example a conventional solid carrier material such as starch, lactose, dextrin or magnesium stearate. The iron complex will of course be present in such a preferred composition in solid form, which form is accordingly a preferred one for the complex, and such a solid composition may conveniently be presented as some type of formed composition, for example as tablets, capsules (including spansules), etc.

Although solid compositions are preferred in many applications, liquid compositions are of interest in certain

particular instances, for example human and veterinary intramuscular administration and veterinary oral administration as discussed hereinafter. It is often desirable to produce liquid compositions containing a higher concentration than is readily obtainable with a purely aqueous composition or indeed one containing organic solvents such as simple monohydric alcohols. It has been found that this may be done by the use of solvents containing two or more hydroxy groups or a hydroxy and an ether group, especially of glycols or glycol ethers, either in admixture with water or, for better solubilisation, alone. The glycol ethers of particular interest are the mono-ethers containing as an etherifying group an aliphatic hydrocarbon group of 1 to 6 carbon atoms as described above, for example a methyl group, such a glycol mono-ether being methyl ethylene glycol. In general, however, the glycols themselves are preferred. Examples of such glycols are the simple dihydroxy alkanes such as ethylene glycol as well as those more complex compounds comprising two hydroxy groups attached to a chain containing both carbon and oxygen atoms, such as triethylene glycol, tetraethylene glycol and polyethylene glycol, for example of 4,000 daltons molecular weight. Triethylene glycol and especially tetraethylene glycol are of particular interest in view of their very low toxicity. By using such glycols and glycol ethers it is possible to increase solubility for many complexes to 10 to 20 mg/ml.

In the case of animals, compositions for parenteral administration are of greater interest than with humans. The problems of iron deficiency anaemia in newly born pigs arise primarily during the first three weeks or so of their life when a very rapid weight gain takes place. The unusual routes for administration of the iron complexes of the present invention to young piglets are parenteral, for example intramuscular, or oral, for example as a liquid preparation "injected" into the mouth. However, an alternative approach is to enhance the iron content of the milk on which the piglets are feeding by treating the mother pig using oral or parenteral administration, for example with an injectable slow release preparation (such an approach may also be of interest in a human context). When it is applicable to feed piglets on foodstuffs other than the milk of the mother pig, it may also be possible to effect the pharmaceutical administration of the iron complex in this other foodstuff.

Other forms of administration than by injection or through the oral route may also be considered in both human and veterinary contexts, for example the use of suppositories or pessaries for human administration.

Compositions may be formulated in unit dosage form, i.e. in the form of discrete portions containing a unit dose, or a multiple or sub-unit dose. Whilst the dosage of hydroxypyron iron complex given will depend on various factors, including the particular compound which is employed in the compositions, it may be stated by way of guidance that maintenance at a satisfactory level of the amount of iron present in the human body will often be achieved using a daily dosage, in terms of the iron content of the compound, which lies in a range from about 0.1 to 100 mg and often in a range from 0.5 to 10 mg, for example 1 or 2 mg, veterinary doses being on a similar g/Kg body weight ratio. However, it will be appreciated that it may be appropriate under certain circumstances to give daily dosages either below or above these levels. In general, the aim should be to provide the amount of iron required by the patient without administering any undue excess and the properties of the pharmaceutical compositions according to the present invention are particularly suited to the achievement of this aim. Similarly, the

concentration of iron in the pharmaceutical composition in the form of the hydroxypyron complex may vary quite widely, for example over a range from about 0.001 to about 20% w/w. However, it is more usual for the concentration to exceed 0.01% w/w and it may often exceed 0.05 or 0.1% w/w, whilst a more usual limit for the upper end of the range is about 13% w/w. A common range of concentration is 0.05 to 5% w/w, for example 0.2 to 0.5, 1 or 2% w/w.

Where desired, more than one hydroxypyron iron complex as described above may be present in the pharmaceutical composition or indeed other active compounds having the ability to facilitate the treatment of anaemia, such as folic acid. Another additional component which may be included in the composition, if desired, is a source of zinc. Iron compounds used in the treatment of iron deficiency anaemia can inhibit the mechanism of zinc uptake in the body and this can cause serious side effects in the foetus when treating anaemia is in a pregnant female. It is believed, however, that the iron complexes of the present invention have a further advantage in that they either do not have this effect or exhibit the effect at a lower level than the compounds at present used in the treatment of anaemia. Accordingly, it may often be the case that the level of zinc providing compound added to the composition may not require to be high or, with preferred formulations of the iron complexes, may be dispensed with altogether.

Although certain of the iron complexes, for example iron maltol, have previously been proposed for use as colouring agents in foodstuffs, it had never previously been appreciated that they have any therapeutic use and the conditions proposed for the use of such complexes as colouring agents would not generally be such as to lead to any significant physiological effect. Accordingly the present invention includes an iron complex of 3-hydroxy-4-pyrone or of a 3-hydroxy-4-pyrone in which one or more of the hydrogen atoms attached to ring carbon atoms are replaced by an aliphatic hydrocarbon group of 1 to 6 carbon atoms, for use in medicine, particularly in the treatment of iron deficiency anaemia.

We have found that the iron complexes described herein are particularly suited to the treatment of iron deficiency anaemia, both in humans and also in a veterinary context and particularly for the treatment of various mammalian species, especially pigs. Thus, the chelating agents which they contain, and particularly maltol, have a high affinity for iron ($\log \beta_3=30$ for maltol) but a lower affinity for copper (II), zinc (II), calcium and magnesium. Both the high affinity of maltol for iron and its low affinity for calcium are reflected in its K_{sol} value $\{\log K_{sol}$ is defined as being equal to $\log \alpha_{Fe(L)n} + 21 - [pK_{sp} + n \log a_L(H^+) + m \log a_L(Ca^{++})]$ where $\log \alpha_{Fe(L)n}$ is the cumulative affinity constant of the ligand in question for iron (III), pK_{sp} is the negative logarithm of the solubility product for $Fe(OH)_3$ and has a value of 39, n and m are the number of hydrogen and calcium ions, respectively, which are bound to the ligand, and $a_L(H^+)$ and $a_L(Ca^{++})$ are the affinities of the ligand for hydrogen ions and calcium ions, respectively}. In order to solubilise iron(III) hydroxide, $\log K_{sol}$ must be greater than 0. The value of K_{sol} for maltol is 0.8 and this is also sufficiently large to prevent appreciable competition from phytate, phosphate, thiols and other potential ligands likely to occur in the intestinal lumen. In order to exchange iron efficiently with transferrin, the $\log K_{sol}$ value should be close to that of apotransferrin, which is 6.0, so that maltol is also suitable in this respect. Moreover, although the neutral 3:1 maltol:iron(III) complex is thermodynamically stable (thermodynamic stability constant=30) it is also extremely labile and is therefore able to donate iron

to high affinity sites, such as those found in apotransferrin. The half life for the exchange of iron(III) between the maltol complex and apotransferrin is 1 minute whereas, by contrast, the corresponding figure for the complex of EDTA with iron(III) is 4 days.

It will be appreciated, however, that in addition to possessing properties such as those described above for iron maltol, a compound which is to act as a source of iron though oral administration is required to show a high level of membrane permeability. An indication of the properties of a compound in this respect is provided by the value of the partition coefficient (K_{part}) obtained on partition between n-octanol and Tris hydrochloride (20 mM, pH 7.4; Tris representing 2-amino-2-hydroxymethylpropane 1,3-diol) at 20° C. and expressed as the ratio (concentration of compound in organic phase)/(concentration of compound in aqueous phase). The value of K_{part} for the neutral 3:1 maltol:iron(III) complex is 0.5, which is well placed in the preferred range of 0.2 to 1.0 and compares favourably with the figures of 0.001 and 0.0015 for the EDTA:iron(III) complex and iron(III) ascorbate, respectively.

The value of the iron complexes of the present invention is configured by various in vitro and in vivo tests. Thus, their ability to permeate biological membranes is confirmed in practice by tests of the ability of the ^{59}Fe labelled iron complexes to permeate erythrocytes. Moreover, iron complexes of the present invention have been found to exhibit a high level of efficiency in promoting iron uptake, as measured in the rat small intestine, as compared with a range of other iron complexes currently marketed for the treatment of iron deficiency anaemia. In vivo experiments in the cat and rat have confirmed the value of iron maltol compounds as a source of iron, the iron uptake obtained either on intravenous administration or on direct administration into the small intestine being markedly superior to that obtained with commercially available iron compounds such as iron sulphate, iron EDTA and iron gluconate. It was found from these experiments that the iron was not excreted to any significant extent in the urine but became generally distributed throughout the body, the complexes donating iron to transferrin to an equilibrium level one they are present in the bloodstream.

Certain aspects of their formulation may enhance the activity of the complexes in particular contexts. Thus, although the neutral 3:1 ferric complexes are stable over a wide pH range from about 4 or 5 up to 10, they will dissociate at the pH values of less than 4 prevailing in the stomach to form a mixture of the 2:1 and 1:1 complex together with the free hydroxypyrrone, and it has been found that the blood levels of ^{59}Fe achieved on administration of the 3:1 complex into the small intestine are much higher than when administration is made into the stomach. However, when the stomach contents is flushed to the small intestine in in vivo cat experiments an increase of iron uptake occurs almost immediately. The undesirable effects of this dissociation on iron uptake may be countered by using one or more of the following procedures in the formulation of the iron complex. Firstly, one of several variations may be employed which avoid or reduce exposure of the iron complex to the acidic conditions of the stomach. Such approaches may involve various types of controlled release system, ranging from one, which may for example be based on a polymer, which simply provides a delayed release of the complex with time, through a system which is resistant to dissociation under acidic conditions, for example by the use of buffering, to a system which is biased towards release under conditions such as prevail in the small intestine, for

example a pH sensitive system which is stabilised towards a pH of 1 to 3 such as prevails in the stomach but not one of 7 to 9 such as prevails in the small intestine. Since the pH of the stomach is higher after a meal, it may be advantageous, whatever method of formulation is used, to administer the iron complexes at such a time.

A particularly convenient approach to a controlled release composition involves encapsulating the iron complex by a material which is resistant to dissociation in the stomach but which is adapted towards dissociation in the small intestine (or possibly, if the dissociation is slow, in the large intestine). Such encapsulation may be achieved with liposomes, phospholipids generally being resistant to dissociation under acidic conditions. The liposomally entrapped 3:1 iron(III) complexes can therefore survive the acid environment of the stomach without dissociating to the 2:1 and 1:1 complexes, and the free hydroxypyrrone. On entry into the small intestine, the pancreatic enzymes rapidly destroy the phospholipid-dependent structure of the liposomes thereby releasing the 3:1 complex. Liposome disruption is further facilitated by the presence of bile salts. However, it is usually more convenient to effect the encapsulation, including microencapsulation, by the use of a solid composition of a pH sensitive nature.

The preparation of solid composition adapted to resist dissociation under acidic conditions but adapted towards dissociation under non-acidic conditions is well known in the art and most often involves the use of enteric coating, whereby tablets, capsules, etc, or the individual particles or granules contained therein, are coated with a suitable material. Such procedures are described, for example, in the article entitled "Production of enteric coated capsules" by Jones in *Manufacturing Chemist and Aerosol News*, May 1970, and in such standard reference books as "Pharmaceutical Dosage Forms", Volume III by Liebermann and Lackmann (published by Marcel Decker). One particular method of encapsulation involves the use of gelatine capsules coated with a cellulose acetate phthalate/diethylphthalate layer. This coating protects the gelatin capsule from the action of water under the acid conditions of the stomach where the coating is protonated and therefore stable. The coating is however destabilised under the neutral/alkaline conditions of the intestine where it is not protonated, thereby allowing water to act on the gelatin. Once released in the intestine the rate of permeation of the intestine wall by the water soluble 3:1 iron(III) complex is relatively constant irrespective of the position within the intestine, i.e. whether in the jejunum, ileum or large intestine. Other examples of methods of formulation which may be used include the use of polymeric hydrogel formulations which do not actually encapsulate the iron complex but which are resistant to dissociation under acidic conditions.

A second approach to countering the effect of the acidic conditions prevailing in the stomach involves formulation of the complex in the pharmaceutical composition together with the metal-free hydroxypyrrone from which it is derived. The dissociation of the neutral 3:1 ferric complex, for example, involves various equilibria between this complex, the 2:1 and 1:1 complexes, and the metal-free compound, so that the presence of the latter will inhibit this dissociation. Any proportion of the free compound can be advantageous in this context but little further advantage accrues from increasing the proportion beyond a certain level. A preferred range for the molar proportion of the free compound present in compositions according to the present invention is thus from 0 to 100 moles of free hydroxypyrrone:1 mole of iron complex, particularly of the neutral 3:1 iron(III) complex.

Conveniently, a proportion of up to no more than 20, 30 or 50 moles:1 mole is used with a lower level of 0.5, 1 or 2 moles:1 mole. Although to obtain a marked effect upon dissociation of the iron complex a proportion of at least 5 or 10 moles:1 mole is usually employed, it should be emphasised that even a molar ratio such as 1:1 will achieve a noticeable degree of acid stabilisation of the iron complex. Thus although a range of, for example, from 10 moles:1 mole to 20 moles:1 mole of metal-free hydroxy pyrone:iron complex will often be suitable to produce a marked effect, a range of, for example, 3 or even 1 mole:1 mole to 10 moles:1 mole will still produce a worthwhile effect without requiring administration of the larger amounts of the hydroxy pyrone. It will be appreciated that a mixture of the hydroxypyrrone and iron complex in at least certain of the proportions described above is novel for the various hydroxypyrrones, irrespective of the particular form of the iron complex, and that such novel mixtures are included by the present invention. The use of such a mixture is an important feature of the present invention since in principle it can enable one to obtain almost quantitative uptake of iron from the complex.

The use of an uncompleted hydroxypyrrone in admixture with its iron complex may also have another advantage in addition to the prevention of dissociation of the iron complex under acidic conditions. Thus, in certain pathological conditions there may be an excess of iron deposited at certain sites even though the patient exhibits an overall anaemia. In these patients the use of such a mixture has the advantage that the iron complex will remedy the overall anaemia whilst the free hydroxypyrrone will act to remove iron from pathological to physiological sites. However, although it is preferable for the hydroxypyrrone present in an iron donor to be rapidly metabolized in order that iron may be efficiently transferred to the binding proteins and eventually to the iron requiring mechanisms within the body, it is preferable for a hydroxypyrrone being used as an iron remover not to be rapidly metabolized so that it remains in the system, taking up iron, for an extended period. Thus, for example, maltol is rapidly metabolized and is therefore particularly suited for use as an iron complex, but for this same reason it is not appropriate for use in the free form. It is also the case that different compounds may function more efficiently either in the free form as an iron remover or in complex form as an iron donor for quite other reasons. Accordingly the present invention includes a mixture of an iron complex of a 3-hydroxy-4-pyrone or of a 3-hydroxy-4-pyrone in which one of the hydrogen atoms attached to ring carbon atoms are replaced by an aliphatic hydrocarbon group of 1 to 6 carbon atoms, together with a different 3-hydroxy-4-pyrone. Alternatively, the different 3-hydroxy-4-pyrone may be replaced by a quite different form of iron chelating agent. Examples of such order iron chelating agents which may be used include the substituted 3-hydroxypyrid-2-ones and -4-ones, and 1-hydroxypyrid-2-one and substituted 1-hydroxypyrid-2-ones (and salts of these various pyridones with a physiologically acceptable cation) described in copending U.S. patent application Ser. Nos. 592,271 (now U.S. Pat. No. 4,585,780) 478,493 and 651,772 (now U.S. Pat. No. 4,587,240).

When a free hydroxy-4-pyrone, hydroxypyrid-2-one, hydroxypyrid-4-one, or other iron chelating agent is present in admixture with the iron complex of a hydroxy-4-pyrone for the purpose of acting as an iron remover, then the amount of this agent used may be different than when a free hydroxypyrrone necessarily corresponding to that present in the iron complex is present primarily to prevent dissociation.

Thus the daily dosage of the iron complex may be as above and the daily dosage of the free iron chelating agent, particularly when this is a hydroxypyrid-2- or -4-one or a 1-hydroxy-pyrid-2-one, may be that quoted in the copending applications referred to above, i.e. about 0.1 g to 5 g for human use, particularly 0.5 g to 2 g, from which it will be seen that the proportion of iron complex and free iron chelating agent used in such a context may extend across a wide range but referred amounts of the free iron chelating agent to be higher than when this is necessarily a hydroxypyrrone.

It will be appreciated that the present invention also includes a method for the treatment of a human or other mammalian patient which comprises administering to said patient an iron complex of 3-hydroxy-4-pyrone or of a 3-hydroxy-4-pyrone in which one or more of the hydrogen atoms attached to ring carbon atoms are replaced by an aliphatic hydrocarbon group of 1 to 6 carbon atoms in order to effect an increase in the levels of iron in the patient's bloodstream.

In addition to the pharmaceutical uses of the iron complexes discussed above they are also of potential interest as a source of iron in various other contexts including cell and bacterial growth, plant growth, and the control of iron transport across membranes.

This invention is illustrated by the following Examples:

EXAMPLES

Example 1

The Preparation of Iron Maltol

A chloroform solution of maltol is mixed with a 1M solution of ferric chloride in ethanol to provide a 3:1 molar ratio of maltol:iron in the mixture. After 5 minutes at 20° C., a 10 molar excess of solid sodium carbonate is added and the mixture is stirred for 10 minutes. The mixture is then filtered and the solvent evaporated to give the neutral complex containing maltol and the ferric cation in 3:1 proportion. Recrystallisation of the 3:1 complex from the ethanol gives wine red needle crystals in an essentially quantitative yield, m.p. 275°, ν_{max} (nujol) 1600 cm^{-1} .

The use of an excess of maltol above the 3:1 molar ratio leads to an essentially quantitative yield of a solid mixture of the excess maltol and the 3:1 iron maltol complex on rotary evaporation, this mixture not being deliquescent.

The partition coefficient K_{part} (concentration in n-octanol/concentration in aqueous phase) between n-octanol and Tris hydrochloride (20 mM, pH 7.4) of maltol and of its 3:1 iron complex is measured at 10^{-4} M by spectrophotometry. Acid washed glassware is used throughout and, following mixing for 1 minute, the aqueous/n-octanol mixture is configured at 1000 g for 30 seconds. The two resulting phases are separated for a concentration determination by spectrophotometry on each. For maltol, the range 220–340 nm is used for the concentration determination whilst for the complex the range 340–640 nm is used. Typically, a value of 0.66 is obtained for maltol and of 0.50 for its iron complex, whilst comparative experiments on iron(III) EDTA and iron(III) ascorbate give much smaller values of 0.001 and 0.0015, respectively.

The ability of the iron complex of maltol to bind to haemoglobin is investigated by studying the elution profile of a ^{59}Fe label when a mixture of haemoglobin and the ^{59}Fe -labeled complex (at 1 mM concentration) in NaCl (130 mM) buffered to pH 7.4 by Tris hydrochloride is applied to a PH-10 column (Sephadex G-10 gel permeation column—

Pharmacia). Typically, no evidence is found for binding of the complex to haemoglobin which is an advantageous finding since such binding reduces availability of the iron.

The ability of the iron complex of maltol to bind to bovine serum albumen (BSA) is investigated through a similar procedure in which the complex is applied to a column with BSA rather than haemoglobin. The iron complex also shows little ability to bind to BSA.

Example 2

In vitro Tests on Permeation of Iron Complexes into Human Erythrocytes

The accumulation of iron by human erythrocytes which are associated with the iron complexes of maltol described in Example 1, and various other iron compounds by way of comparison, was studied by incubating the erythrocytes for 1 hour at 37° C. in a medium consisting of the ⁵⁹Fe labelled iron compound in aqueous sodium chloride (130 mM) buffered to a pH of 7.4 by Tris hydrochloride. Following this period of incubation an aliquot of the erythrocyte/medium mixture was placed above a layer of silicone oil (ρ=1.06) and the erythrocytes separated by centrifugation through the oil. The ⁵⁹Fe levels associated with the erythrocytes and the incubation medium were then counted and presented as a distribution ratio (concentration in erythrocytes/concentration in medium). The ratios obtained for the various iron compounds after incubation for 1 hour are shown in Table 1 where it will be seen that the uptake of iron is clearly much greater with the iron maltol complex than with the other compounds. Values of less than 0.1 are probably associated with binding to the external surface and do not represent transmembrane movement of iron. Moreover, although a period of 1 hour was employed in order to facilitate monitoring of the more slowly permeating iron compounds, the uptake of iron maltol complex reached equilibrium at the level shown after about 15 minutes.

TABLE 1

Compound	Concentration (mM)	Distribution ratio
Fe ^{III} (maltol) ₃	3	1.60
Fe ^{II} gluconate	1	0.08
Fe ^{III} ascorbate	1	0.12
Fe ^{III} citrate	1	0.05
Fe ^{III} EDTA	1	0.05

When the above described procedure was applied using ratios of maltol to iron of less than 3:1 larger apparent distribution ratios were observed than 1.60. However, this is explained by the non-specific binding of the positively charged 2:1 and 1:1 maltol:iron complexes to the surface of the erythrocytes which possesses a net negative charge, being rich in both phosphate and sulfate moieties. Experiments to determine the percentage of ⁵⁹Fe associated with erythrocyte ghosts after lysis confirm this hypothesis. In one experiment, lysis was initiated by a small volume of 10% v/v Triton X100 and in a second experiment by a 10 fold excess of water. In each case the resulting ghosts were centrifuged through silicon oil (ρ=1.02) and, as will be seen from Table 2, very little of the 3:1 maltol:iron complex was found to be bound to the membranes, in contrast with the situation with the 2:1 and 1:1 complexes. Such binding is of course undesirable as the complex is likely to remain tightly bound to the membrane by electrostatic interactions and not be transmitted across it.

TABLE 2

Ratio of maltol:iron	Iron associated with with ghosts (%)	
	Triton lysis	Hypotonic lysis
0:1	100	—
1:1	55	63
2:1	22	39
3:1	<1	<5

Example 3

In vitro Tests on Permeation of Rat Jejunal Sac by Iron Complexes

The iron uptake into the several space of the rat jejunal sac was compared for the iron complex of maltol described in Example 1 and various other iron compounds by way of comparison. Rats (male Sprague Dawley, 60 g) were killed and the jejunum removed, everted and cut into three segments (4 cm length). The segments were tied at both ends, filled with Krebs Ringer buffer (0.2 ml) and incubated in Krebs Ringer buffer containing the appropriate ⁵⁹Fe compound at 37° C. for periods up to 90 minutes. The contents of the sac were counted for ⁵⁹Fe and measured spectrophotometrically.

The results obtained for the iron maltol complex and for 6 other iron compound which are each contained in preparations marketed for the treatment of iron deficiency anaemia are shown in Table 3, the iron uptake for each at 15 and 60 minutes after the initiation of the experiment being shown relative to that for ferric chloride as 1. It will be seen that the iron maltol complex provides a level of iron uptake which is significantly higher than the levels observed for any of 6 compounds in current use for the treatment of iron deficiency anaemia. The uptake of the iron maltol complex was linear for a period 90 minutes. Moreover, the uptake increased linearly as the concentration of the complex was increased over a range of 0.5 to 10 mM, so it does not show saturation kinetics and the process is thus non-facilitated and therefore should occur in all natural membranes.

TABLE 3

Compound	Relative iron uptake	
	15 minutes	60 minutes
FeCl ₃	1	1
Fe ^{III} (maltol) ₃	40	5.8
Fe ^{II} sulphate	2.4	1.4
Fe ^{II} fumarate	4.0	1.8
Fe ^{II} gluconate	1.6	0.8
Fe ^{II} succinate	2.0	1.0
Fe ^{III} ascorbate	0.4	0.8
Fe ^{III} citrate	2.0	1.8

The procedure described above was used to compare the uptake of iron from buffer containing differing molar proportions of maltol:iron. The results obtained are presented in Table 4 which shows the amount of time transferred via the maltol complex into the serosal contents of the sac, the basal uptake of iron measured in a control experiment being subtracted in each case. It will be seen that the amount of iron transferred in the case of a 3:1 molar proportion of maltol:iron(III) is much higher than in the other two cases and, moreover the low, but significant level of iron uptake

observed in the case of a 2:1 ratio is substituted to the proportion of the 3:1 complex (containing 13% of the total iron) present under these conditions.

TABLE 4

Maltol/iron (molar ratio)	Iron uptake (n mole)
1:1	1.6
2:1	4.0
3:1	30.0

Example 4

In vivo Test of Action of Iron Compounds in the Rat

The action of the iron complex of maltol described in Example 1 was compared with that of iron(II) sulphate, iron(III) EDTA (1:1 gmolar ratio) and iron(II) gluconate.

Groups of rats (300–350 g) were anaesthetised with nembutal (0.25 ml) and then with ether. A mid-line incision was made and the ⁵⁹Fe labelled sample (100 µg Fe 10 µCi) was passed into the lumen of the duodenum via a small incision. The abdominal well was then closed with a suture. The animals were sacrificed 1, 2, 4 and 6 hours after the administration of the compound and the various organs were monitored for their ⁵⁹Fe content. The data is presented as histograms in FIGS. 1 to 4 which relate to iron maltol, iron sulphate, iron EDTA and iron gluconate, respectively, and shown the levels of ⁵⁹Fe in cpm after various time intervals for the different organs, the data in each case representing a mean of the results for three individual animals. In the case of the data for blood and sternum (bone marrow) the counts given are cpm/ml and cpm/g respectively, whilst in all other cases they are the total cpm counts. The various histograms have been normalised and consequently are directly comparable.

A comparison between FIGS. 1 and 2 shows that the neutral 3:1 maltol:iron(III) complex is markedly superior to iron(II) sulphate for the introduction of iron via the rat intestine. The gut washings (which contain non-absorbed iron) show a much lower level of counts for the maltol complex, and the counts associated with the gut wall, liver, blood, bone marrow and spleen are correspondingly greater. It is clear from FIG. 1 that ⁵⁹Fe associated with maltol enters the intestine wall very rapidly and from there it is efficiently removed by the blood supply. Iron is deposited in the bone marrow continuously throughout the 6 hour period at an apparently constant rate.

The maltol complex is also more efficient than iron(III) EDTA as shown by FIG. 3. With the later complex, the gut washings remain high for 4 hours and may be pressured to decrease only due to the effect of natural bowel movements translocating material from the portion under investigation to lower portions of the intestine. The levels in the intestine wall and blood are extremely low. Although iron is transferred to both bone marrow and spleen, this is at reduced rates as compared to those obtained with the maltol complex. As shown by FIG. 4, iron(II) gluconate proved more effective than the sulphate or the EDTA complex, although deposition in the gut wall was less than that observed with the maltol complex. The decrease was reflected in the lower levels of ⁵⁹Fe in both bone marrow and the spleen, the difference being particularly marked after 6 hours. In view of the much higher levels of ⁵⁹Fe trapped in the intestine

wall in the case of the maltol complex, it may be predicted that this compound facilitates a more prolonged supply of iron than iron(II) gluconate.

This test illustrates the superiority of the neutral 3:1 maltol:iron(III) complex as compared with three commonly used “soluble iron” preparations for the movement of iron across the rat jejunal wall into the blood circulation, the iron maltol being very rapidly removed from the lumen of the intestine.

Example 5

In vivo Test of Action of Iron Complex in the Cat

The action of the iron complex of maltol described in Example 1 was compared with that of iron(III) EDTA (1:1 molar ratio) which is one of the iron compounds currently marketed for the treatment of iron deficiency anaemia. Cats were anaesthetised with chloralase (60 mg/kg) and pentobarbitone sodium (60 mg/kg) (i.p.), having been kept free of food for 18 hours. In each animal the trachea was cannulated to maintain a clear airway and to allow positive pressure artificial respiration if necessary. The left femoral vien was cannulated for the intravenous administration of drugs and physiological saline solution. Arterial blood pressure was monitored by a Washington pressure transducer through a cannula inserted into the femoral artery of the right hind leg. Arterial blood samples were taken at appropriate intervals from a short cannula inserted into an external carotid artery. Body temperature was monitored with a rectal thermometer. Each animal was given heparin (1000 iu/kg) as anticoagulant and additional small amounts of pentobarbitone sodium if needed to maintain a satisfactory level of anaesthesia.

If those animals where the iron compounds were to be administered into the duodenum, a mid-line incision was made in the abdomen to reveal the intestines. A cannula was then inserted through a small cut such that its tip rested approximately 5 cm below the opening of the bile duct. The cannula was then sutured in place and the abdominal wall closed with stitches.

The iron maltol complex (100 µg Fe) alone (3:1 molar ratio of maltol:iron) and together with a large excess of maltol (40:1 molar ratio of maltol:iron) was injected intravenously in separate experiments and 0.25 ml samples of blood were taken in intervals. The apparent volume of distribution of the compound was calculated by extrapolation of the log-linear blood concentration curve to zero. (The volume corresponds to a value between that of the total extracellular space of the animal and the blood volume.) Elimination of ⁵⁹Fe from the blood followed first order kinetics with a rate constant of -0.022/minute in the presence and absence of excess maltol, as illustrated in FIG. 5 which shows the ⁵⁹Fe level in the blood in cpm/0.25 ml plotted against time in the case of one typical experiment of each type in an individual cat.

The distribution of ⁵⁹Fe in the tissues of the animal after the same intravenous experiment to which FIG. 5 relates (the lower end of the ordinate in this Figure represents the background level) was investigated and the typical results are shown in Table 5. The amount of ⁵⁹Fe administration in this experiment was 4 µCi or 2.2×10⁶ cpm. It will be seen that approximately 10% of the dose was located in the combined tissue of the heart, liver and spleen. As less than 0.2% of the dose was located in the urine, the bulk (approximately 90% of the ⁵⁹Fe was almost certainly directed to the bone marrow and extremely high levels were found to be located in the sternum.

As indicated previously, the maltol complex is able to donate iron rapidly to transferrin and it is hypothesised that such an exchange occurs as soon as the complex is delivered to the plasma; and that the initial plateau (represented in FIG. 5 by a dotted line) represents saturation of the plasma transferrin pool with ⁵⁹Fe. When there is net donation of ⁵⁹iron from the plasma into the organ is of the animal, the blood levels of radioactivity begin to fall, the major route of transfer of iron bound to transferrin being to the bone marrow, liver and spleen. Binding of ⁵⁹Fe to transferrin prevents its excretion in the urine.

TABLE 5

(Iron maltol, i.v.)				
Tissue	Total tissue weight (g)	Sample weight (g)	Net ⁵⁹ Fe content (cpm/g)	Net total ⁵⁹ Fe content (cpm)
Heart	14.4	0.91	490	7,056
Liver	105	1.3	510	53,550
Spleen	8.4	0.86	14,890	125,076
Kidney	12.2	1.05	546	6,661
Skeletal muscle	—	1.85	0	0
Sternum (bone marrow)	—	1.2	3,200	—
Urine	—	1	152	<3,000

Identical experiments carried out with ⁵⁹Fe labelled iron (III) EDTA gave a entirely different picture as will be seen for the results of a typical experiment illustrated in FIG. 6 (in which the lower end of the ordinate represents the background level) and Table 6 (the amount of ⁵⁹Fe administered in this experiment was 2 μ Ci but the figures given in the table have been adjusted to correspond to a dosage of 2.2 \times 10⁶ cpm in order to facilitate comparison with Table 5). In this experiment the radioactivity in the blood showed no initial plateau. Instead, loss of radioactivity followed at least a two-component process such that a large amount found its way to the urine rather than to the issues. The rate constant of the elimination from the blood to the linear phase of the regression was 0.023/minute. The concentration of radioactivity in the kidney and urine, and not in the bone marrow or spleen, would indicate that iron in this form does not appear to be able to attach to transferrin in the plasma and protect itself from urinary excretion. The combined tissue of heart, liver and spleen contained only 1% of the original dose at the end of the experiment, whereas the urine contained over 50%. This is in accord with the fact that EDTA does not exchange iron with transferrin rapidly.

TABLE 6

(Iron EDTA, i.v.)				
Tissue	Total tissue weight (g)	Sample weight (g)	Net ⁵⁹ Fe content (cpm/g)	Net total ⁵⁹ Fe content (cpm)
Heart	15.5	1.01	209	3,248
Liver	75	1.21	261	19,600
Sternum (bone marrow)	—	0.28	1,164	—
Spleen	11.1	0.89	162	1,814
Kidney	19.2	1.47	1,134	21,770
Skeletal muscle	—	2.59	95	—
Urine	19 ml	2 ml	62,156	1,180,900

The iron maltol complex (100 μ g Fe) was also administered to the duodenum of the cat in the presence of a 40 fold excess of maltol followed by 5 ml of 150 ml tris hydrochlo-

ride buffer (pH 7.4). In this case the ⁵⁹Fe content of the blood, as shown in FIG. 7, reaches a maximum level 2 hours after the initial administration (the readings start at about 300 cpm/0.5 ml which represents the background reading). The distribution of ⁵⁹Fe in the tissues of the animal after the same duodenal experiment to which FIG. 7 relates were investigated and the typical results are shown in Table 7. The amount of ⁵⁹Fe and administered in this experiment was 10 μ Ci or 5.327 \times 10⁶ cpm into a 2.9 kg cat. It will be seen that the distribution of the ⁵⁹Fe after 4 hours was similar to that after intravenous infusion, with low levels in the kidney and urine and high levels in both the spleen and bone marrow.

TABLE 7

(Iron maltol, per duodenum)				
Tissue	Total tissue weight (g)	Sample weight (g)	Net ⁵⁹ Fe content (cpm/g)	Net total ⁵⁹ Fe content (cpm)
Heart	14	0.633	50	1,106
Liver	81	1.45	400	32,400
Spleen	12.7	1.19	3,783	48,047
Kidney	14.4	0.835	79	1,138
Sternum (bone marrow)	10	1.26	790	7,905
Bile	~5 ml	1 ml	2,200	~11,000
Urine	~10 ml	1 ml	22	~220

When ⁵⁹Fe labelled iron(III) EDTA was administered duodenally in the same manner, the plasma levels of radioactivity hardly exceeded the background level and are therefore not illustrated in a Figure. The distribution of ⁵⁹Fe in the tissues of the animal after the same duodenal experiment were investigated and the typical results are shown in Table 8. The amount of ⁵⁹Fe administered in this experiment was 10 μ Ci or 2.65 \times 10⁶ cpm into a 2/9 kg cat. It will be seen that, although some ⁵⁹Fe entered the tissues, rather low levels were detected in the spleen and bone marrow (sternum) whereas a large proportion of the dose was located in the urine.

TABLE 8

(Iron EDTA, per duodenum)				
Tissue	Total tissue weight (g)	Sample weight (g)	Net ⁵⁹ Fe content (cpm/g)	Net total ⁵⁹ Fe content (cpm)
Heart	15.3	1.18	188	2,878
Liver	59.3	0.78	499	29,574
Kidney	11.3	0.90	1,762	19,913
Spleen	4.4	0.42	200	880
Sternum (bone marrow)	—	0.78	917	—
Skeletal muscle	—	1.48	117	—
Urine	15 ml	5 ml	36,306	544,596

Example 6

Polymer Formulation of Iron Maltol

A solution of ferric chloride (concentration between 1 and 5% w/v), together with maltol in a weight ratio of 8 parts by weight of maltol to 1 part by weight of ferric chloride, is prepared in a 4.5:4.5:1 v/v/v mixture of chloroform:methanol:water. Sodium carbonate is added in a 10 molar excess over the iron content in order to remove hydrochloric acid and the precipitated NaCl and Na₂CO₃ are filtered off. The preparation is contacted with a cross-linked polyethylene

glycol hydrogel material to effect take up the solution by the polymer and provide a polymer formulation of iron maltol in the presence of excess maltol.

Example 7

Capsule Formulation of Iron Maltol

A preparation of iron(III) maltol in admixture with maltol (containing 1 part by weight of iron to 10 parts by weight of maltol) is obtained by the addition of a 1M ethanolic solution of ferric chloride to a methylene chloride solution of appropriate amount of maltol, followed after 5 minutes at 20° C. by treatment with a 10 molar excess of solid carbonate, stirring for 10 minutes, filtration and evaporation of the solvents.

The resulting solid iron(III) maltol preparation is divided into 50 mg quantities and added to standard gelatine capsules (16x5 mm), each capsule containing 5 mg of iron. The capsules are then coated with a cellulose acetate phthalate/diethylphthalate layer (6 mg coat per cm² of capsule surface) in a small scale procedure analogous to the procedure described by Jones, *ibid*. A proportion of the capsules are treated to add a second similar coating.

Such capsules are resistant to dissociation in the stomach but will undergo dissociation in the intestine. Thus, when treated at 37° C. with dilute aqueous hydrochloric acid (pH 2.0) the singly coated capsules are typically stable for 30 minutes but in Krebs Ringer bicarbonate solution (pH 7.4) at 37° C. they dissociate to release the iron complex within 1 minute. The doubly coated capsules are typically stable at pH 2.0 for 20 hours, again dissociating within 1 minute at pH 7.4.

Example 8

Liposome Formulation of Iron Maltol

(A) a solution of egg yolk phosphatidyl chloride⁽¹⁾ (40 mg) and cholesterol (40 mg) in chloroform (1 ml) is rotary evaporated in a 50 ml round bottomed flask to form a thin lipid film. An aqueous solution of the 3:1 neutral iron(III) maltol complex (6 ml, 1 mg/ml) is added to the flask and the mixture is vibrated for 15 minutes. Centrifugation (3,000 rev/minute for 10 minutes) yields multilammellar liposomes containing iron(III) maltol.

(1) In a modification of this procedure the phospholipid may be varied among egg yolk phosphatidyl chloride, dimyristoyl phosphatidylcholine and dipalmitoyl phosphatidylcholine together with a preparation of cholesterol varying from 0 to 1 moles of cholesterol per mole of phospholipid.

(B) A chloroform solution (2 ml) of the 3:1 neutral iron(III) maltol complex (5 mg/ml) is added to egg yolk phosphatidylcholine (100 mg) and a cholesterol (50 mg). The solution is rotary evaporated to yield a deep red skin on the surface of a round bottomed flask. Addition of 6 ml of a buffered solution of sodium chloride (100 mM, Tris.HCl: 20 mM, pH 7.4) followed by shaking for 15 minutes leads to a finely dispersed lipid-iron(III) maltol preparation. Centrifugation at 3000 revs/minute for 100 minutes yields a liposome preparation which can be readily freeze dried. The entrapment of iron(III) maltol using this method is particularly efficient.

Liposomes produced by either method are resistant to dissociation in the stomach but will undergo dissociation in the intestine.

Example 9

Tests with Iron Maltol in Human Subjects

Capsules containing 3:1 iron(III) maltol, together with an excess of free maltol were prepared according to the pro-

cedure of Example 7⁽¹⁾ with the modification that at 3:1 molar ratio of maltol:iron was used (no free maltol therefore being present in the capsules) and that a proportion of the ferric chloride used in the preparation of the iron maltol was radiolabelled as ⁵⁹Fe. The capsules each contained 10 mg of iron, this amount being labelled with 1 microcurie of ⁵⁹Fe, and were subjected to a single coating procedure.

(1) When preparing 3:1 iron(III) maltol on a large scale for inclusion in bulk preparation of capsules or other forms of compositions, it may be advantageous to modify the procedure described in Example 7, and in Example 1, by carrying out the reaction in aqueous ethanol (4:1 v/v C₂H₅OH:H₂O) as the solvent and by adjusting the pH to 8.5 by the addition of 2M aqueous sodium hydroxide rather than with solid sodium carbonate. Evaporation of the solvent after filtration will give a product containing some sodium chloride but this can be removed through recrystallisation of the 3:1 iron(III) maltol from water.

The test involved two normal subjects, i.e. not suffering from iron deficiency, one of whom took a 10 mg dose of iron (one capsule) and the other whom took a 20 mg dose of iron (two capsules). The capsule or capsules were taken after fasting overnight and fasting was then continued for a further 2 hours. The take-up of iron by the subjects was followed by means of whole body counting, using a calibrated body counter, reading being taken 4 to 8 hours after ingestion of the capsules to establish a base level and then after 7 and 14 days. Excretion of iron by the subjects was followed through the counting, in a large volume counter calibrated for iron, of daily urine samples over 3 days and daily stool samples over 10 days.

The following results were obtained. The levels of take-up of iron as indicated by the whole body counting were substantially the same at 7 days and at 14 days (apart from a small direction in the latter figure due to decay of the ⁵⁹Fe rather than excretion thereof). For one subject (10 mg dose) the 14 day figure was 19% whilst for the other (20 mg dose) it was 11%. These figures were confirmed by the counting of stool samples (an essentially zero loss of iron being observed in the urine).

In an exactly similar experiment employing 10 mg and 20 mg of ⁵⁹Fe labelled ferrous sulphate with the same subjects (at the same dosage level of iron) the level of take-up of iron, measured by whole body counting at 7 days, was 13% for one subject (10 mg dose) and 6% for the other (20 mg dose).

This test therefore showed a significantly increased uptake of iron in these normal subjects from 3:1 iron(III) maltol as compared with ferrous sulphate. An even greater difference would be anticipated for iron deficiency subjects.

We claim:

1. A pharmaceutical composition comprising a compound being a tissue permeable, neutral 3:1 hydroxypyruone:iron (III) complex of 3-hydroxy-4-pyrone or of a 3-hydroxy-4-pyrone in which at least one of the hydrogen atoms attached to ring carbon atoms is replaced by an aliphatic hydrocarbon group of 1 to 6 carbon atoms, wherein said complex is encapsulated by a material resistant to dissociation under aqueous acidic conditions.

2. A pharmaceutical composition according to claim 1, in which the iron complex is encapsulated by a solid material which is resistant to dissociation under acidic aqueous conditions but which is adapted for dissociation under non-acidic aqueous conditions.

3. A pharmaceutical composition according to claim 2 in capsule form.

4. A pharmaceutical composition according to claim 2, in which each aliphatic hydrocarbon group is an acyclic group of 1 to 4 carbon atoms.

5. A pharmaceutical composition according to claim 2, in which each aliphatic hydrocarbon group is an alkyl group.

6. A pharmaceutical composition according to claim 2, in which the compound is an iron complex of a 3l-hydroxy-4-pyrone in which at least one of the hydrogen atoms attached to ring carbon atoms is replaced by a substituent selected from the group consisting of methyl, ethyl, n-propyl and isopropyl.

7. A pharmaceutical composition according to claim 6, in which each substituent is a methyl group.

8. A pharmaceutical composition according to claim 6, in which the substituted 3-hydroxy-4-pyrone has a single substituent at the 2- or 6-position.

9. A pharmaceutical composition according to claim 2, in which the compound is the 3:1 iron complex of 3-hydroxy-4-pyrone, 3-hydroxy-2-methyl-4-pyrone, 3-hydroxy-6-methyl-4-pyrone or 2-ethyl-3-hydroxy-4-pyrone.

10. A pharmaceutical composition according to claim 2, in which the compound is the 3:1 iron complex of 3-hydroxy-2-methyl-4-pyrone.

11. A pharmaceutical composition according to claim 2, in which the compound is the 3:1 iron complex of 2-ethyl-3-hydroxy-4-pyrone.

12. A pharmaceutical composition according to claim 2, in which the neutral 3:1 complex is substantially free from complexes containing other proportions of the hydroxypyrrone and iron(III).

13. A pharmaceutical composition according to claim 2, in which the iron complex is in substantially pure form.

14. A pharmaceutical composition according to claim 2 which additionally contains an iron chelating agent.

15. A pharmaceutical composition according to claim 14, in which the iron chelating agent is uncomplexed 3-hydroxy-4-pyrone or an uncomplexed 3-hydroxy-4-pyrone in which at least one of the hydrogen atoms attached to ring carbon atoms is replaced by an aliphatic hydrocarbon group of 1 to 6 carbon atoms, or a salt thereof containing a physiologically acceptable cation.

16. A pharmaceutical composition according to claim 15, which contains a complexed hydroxypyrrone together with the same hydroxypyrrone or a salt thereof in uncomplexed form.

17. A pharmaceutical composition according to claim 16, which comprises the 3:1 iron complex of 3-hydroxy-2-methyl-4-pyrone and uncomplexed 3-hydroxy-2-methyl-4-pyrone or a salt thereof containing a physiologically acceptable cation.

18. A pharmaceutical composition according to claim 2 which additionally contains folic acid.

19. A method for the treatment of a patient to effect an increase in the level of iron in the patient's bloodstream, which comprises administering to said patient in need thereof a composition according to claim 1, in an amount which is effective to achieve said increase.

20. A method for the treatment of a patient to effect an increase in the level of iron in the patient's bloodstream, which comprises administering to said patient in need thereof a composition according to claim 2, in an amount which is effective to achieve said increase.

21. A method according to claim 20, in which the compound is the 3:1 iron complex of a 3-hydroxy-4-pyrone, 3-hydroxy-2-methyl-4-pyrone, 3-hydroxy-6-methyl-4-pyrone or 2-ethyl-3-hydroxy-4-pyrone.

22. A method according to claim 20 in which the compound is the 3:1 iron complex of 3-hydroxy-2-methyl-4-pyrone.

23. A method for the treatment of a patient to effect an increase in the level of iron in the patient's bloodstream

which comprises administering to said patient in need thereof a compound being a 2:1 hydroxypyrrone:iron (III) complex of 3-hydroxy-4-pyrone or of a 3-hydroxy-4-pyrone in which at least one of the hydrogen atoms attached to the ring carbon atoms is replaced by an aliphatic hydrocarbon group of 1 to 6 carbon atoms, in an amount which is effective to achieve said increase.

24. A method according to claim 23, in which each aliphatic hydrocarbon group is an acyclic group of 1 to 4 carbon atoms.

25. A method according to claim 23, in which each aliphatic hydrocarbon group is an alkyl group.

26. A method according to claim 23, in which the compound is an iron complex of a 3-hydroxy-4-pyrone in which at least one of the hydrogen atoms attached to ring carbon atoms is replaced by a substituent selected from the group consisting of methyl, ethyl, n-propyl and isopropyl.

27. A method according to claim 23, in which each substituent is a methyl group.

28. A method according to claim 23, in which the substituted 3-hydroxy-4-pyrone has a single substituent at the 2- or 6-position.

29. A method according to claim 23, in which the compound is a 2:1 iron complex of 3-hydroxy-4-pyrone, 3-hydroxy-3-methyl-4-pyrone, 3-hydroxy-6-methyl-4-pyrone or 2-ethyl-3-hydroxy-4-pyrone.

30. A method according to claim 23, in which the compound is a 2:1 iron complex of 3-hydroxy-2-methyl-4-pyrone.

31. A method according to claim 23, in which the compound is a 2:1 iron complex of 2-ethyl-3-hydroxy-4-pyrone.

32. A method for the treatment of a patient to effect an increase in the level of iron in the patient's bloodstream, which comprises administering to said patient in need thereof a compound being a 1:1 hydroxypyrrone:iron (III) complex of 3-hydroxy-4-pyrone or of a 3-hydroxy-4-pyrone in which at least one of the hydrogen atoms attached to ring carbon atoms is replaced by an aliphatic hydrocarbon group of 1 to 6 carbon atoms, in an amount which is effective to achieve said increase.

33. A method according to claim 32 in which each aliphatic hydrocarbon group is an acyclic group of 1 to 4 carbon atoms.

34. A method according to claim 32, in which each aliphatic hydrocarbon group is an alkyl group.

35. A method according to claim 32, in which the compound is an iron complex of a 3-hydroxy-4-pyrone in which at least one of the hydrogen atoms attached to ring carbon atoms is replaced by a substituent selected from the group consisting of methyl, ethyl, n-propyl and isopropyl.

36. A method according to claim 32, in which each substituent is a methyl group.

37. A method according to claim 32, in which the substituted 3-hydroxy-4-pyrone has a single substituent at the 2- or 6-position.

38. A method according to claim 32, in which the compound is a 1:1 iron complex of 3-hydroxy-4-pyrone, 3-hydroxy-2-methyl-4-pyrone, 3-hydroxy-6-methyl-4-pyrone or 2-ethyl-3-hydroxy-4-pyrone.

39. A method according to claim 32, in which the compound is a 1:1 iron complex of 3-hydroxy-2-methyl-4-pyrone.

40. A method according to claim 32, in which the compound is 1:1 iron complex of 2-ethyl-3-hydroxy-4-pyrone.