

CHREV. 91

ISOELECTRIC POINTS AND MOLECULAR WEIGHTS OF PROTEINS

A TABLE

PIER GIORGIO RIGHETTI and TIZIANA CARAVAGGIO

Department of Biochemistry, University of Milan, Via Celoria 2, Milan 20133 (Italy)

(Received December 2nd, 1975)

CONTENTS

1. Introduction	1
A. Measurement of pH in isoelectric focusing	2
B. pH measurements after IEF in density gradients	3
C. pH measurements after IEF in gel media	4
D. Use of pH markers	5
2. Acknowledgement	22
3. Summary	22
References	22

1. INTRODUCTION

The aim of this article is to provide a list of proteins having known macromolecular parameters, with particular emphasis on isoelectric points (pI) and molecular weights (M.W.). The new updated revision of the classical tables of Darnall and Klotz¹ now has more than 500 entries, each complete with the native molecular weight of the macromolecule, as well as its subunit number and M.W. For each entry, these workers also list the source and, when applicable, the organ of origin. This is very important, since lack of this information in previous tables has generated confusion.

The main aim of Darnall and Klotz¹ was to tabulate proteins having a known quaternary structure. Similar tables have been published by Kleine². However, there is no literature compilation of information on the charge (pI) as well as on the size (M.W.) of proteins. Yet, the pI of a protein is a very important parameter, and its knowledge is very important for the proper use of several techniques, such as disc electrophoresis, isotachopheresis, isoelectric focusing, ion-exchange chromatography and even ammonium sulphate fractionation. Until recently, the literature data on pI values of proteins were scanty and even contradictory. With the advent of isoelectric focusing³⁻⁶, pI data for proteins have been rapidly expanding, to an extent which justifies their collection in a table. Thus, the main emphasis here is on the availability and collection of protein pI values. When available, we also report their M.W. values and quaternary structure. The combined knowledge of these two parameters makes possible the macromolecular mapping of proteins⁷. Most proteins, in fact, are unequivocally determined by values of their pI and M.W. (except, perhaps,

for very close genetic variants of a protein, such as human haemoglobin). Many of the entries in our tables, which lack data on M.W. and subunit structure, can be found in the tables of Darnall and Klotz¹. However, for the same protein the source and/or the organ may be different; therefore we decided not to report such data.

Since most of the proteins reported display microheterogeneity, we have marked the main components with an asterisk when this information was available or could be deduced from graphs or tables. In addition, we also report not only the source and the organ, but also the intracellular location, in the case of isozymes displaying different properties in different subcellular compartments. Unfortunately, there is quite a degree of uncertainty as to the temperature of pH measurement. This is quite an important parameter when defining the pI of a protein, since acid-base equilibria are temperature dependent and a pI of a protein, if quoted without the temperature of measurement, is *per se* ambiguous and lacks an exact physicochemical significance. Some workers never report the temperature of pH measurement. Most workers simply refer to measurements performed at room temperature, which, unfortunately, is not unequivocally determined. Only a few workers have made pH measurements at a precise temperature, in a thermostatted vessel. Since the way in which a pH measurement is made is of utmost importance in defining pI, we summarize here the different techniques used, either in liquid or in gel media.

(A) Measurement of pH in isoelectric focusing

The pI of a protein determined by isoelectric focusing (IEF) also represents its isoionic point in the absence of complex-forming ions⁸. By definition, the isoionic point is a measure of the intrinsic acidity of a pure protein, as it is defined as that pH which does not change on addition of a small amount of pure protein⁹. This definition is also applicable to a protein analyzed by IEF, as the pH of the isoelectric zone does not alter on addition of more protein. It should be remembered that pI values estimated by IEF are temperature dependent and usually decrease with increasing temperature⁶. The difference in pI for the same protein, measured at 25 and at 4°, could be as high as 0.5 pH units, the higher value being obtained at the lower temperature. This difference is usually more pronounced in alkaline regions, and when a protein has a pI value close to the pK of some of its functional groups.

Unfortunately, when given a pI value for a protein at a certain temperature it is difficult to extrapolate this value to another temperature, since the temperature coefficient, dpI/dT , can vary from protein to protein. Thus Bours¹⁰, by measuring the pI values of β -lactoglobulin A and B at 4 and 25°, found a value of dpI/dT of $-(0.9 \pm 0.2) \cdot 10^{-2}$ pH unit per degree. On the other hand, measurements of pI at 4 and 25° for carbonic anhydrase have given a dpI/dT value of $-(1.3 \pm 0.2) \cdot 10^{-2}$ pH unit per degree. In the case of myoglobin¹⁰, the temperature coefficient is as high as $-1.7 \cdot 10^{-2}$ pH unit per degree.

Ideally, pH measurements should be made at the same temperature used during the IEF separation, since in IEF the temperature coefficient, dpI/dT , refers to the corresponding carrier ampholytes rather than to the protein contained in a given fraction. In fact, at the usual concentration of Ampholine (1%), the buffering capacity of the carrier ampholytes in the isoelectric state will normally be sufficient to permit them to dictate the pH even in the presence of as much as 1% of protein^{11,12}. This

means that the pH value assigned to the concentration maximum of a focused protein at a temperature different from the IEF temperature will refer to the corresponding ampholyte fraction rather than to the protein¹³. Furthermore, different protolytic groups are known to display widely different degrees of temperature dependence in their dissociation constants (as a consequence of large differences in their standard heats of ionization)¹⁴. Thus, once a protein has been focused at 4°, a pI measurement made at 20 or at 25° may not represent the true pI of the protein or the pI of the Ampholine molecules surrounding it.

In addition, pH measurements can also be affected by the presence of solutes such as glycerol and sucrose, which are commonly used in IEF. As sucrose lowers the dielectric constant of the solvent and this, in turn, causes the pK values of amino acids to increase¹⁵, it can be expected that the pI value of a given protein is increased on increasing the concentration of sucrose. When IEF is performed in the presence of urea, it should be noted that urea appreciably decreases the activity coefficient of hydrogen ions, resulting in apparently higher pK and pI values of the carrier ampholytes. Correction factors should therefore be used for pI determinations of proteins in urea¹⁶.

(B) pH measurements after IEF in density gradients

In preparative IEF in liquid support media, combined UV and pH readings of all of the collected fractions can be a lengthy and cumbersome procedure. To overcome this, Jonsson *et al.*¹⁷ automated the analysis of column contents by pumping them at a constant rate through series-coupled flow cells, one cell for the recording of pH and the other cell for the determination of UV absorbance. This set-up, however, presented some difficulties in pH determination, which are discussed in detail by these workers¹⁷. More recently, Secchi¹⁸ described a modification of this technique, which appears to overcome many of the problems. A combined glass electrode (Ingold Type 401-M7, provided with a flow vessel and water cooling jacket) is coupled in series with the UV flow cell of an LKB Uvicord II. In this case, the UV signal and pH readings are fed, via a control unit, to the same galvanometric recorder used for the UV readings (LKB 6500). Since the volume lag between the pH and UV cells is only 0.35 ml, accurate and simultaneous pH and UV readings are obtained on the same recorder chart. Moreover, since the pH cell is thermostatted, pH readings are made at the temperature of the focusing column. Another advantage of flow systems is that absorption of carbon dioxide from the air, which might affect pI readings by lowering the pH of the solution, especially in alkaline regions, is avoided.

A similar set-up was reported by Strongin *et al.*¹⁹, who built a pH flow cell in Perspex with Type K-401 and Type GK 2320c pH electrodes from Radiometer A/S. When using flow cells, the peristaltic pump should be attached after the cells, in order to prevent leakage of potassium chloride from the electrodes and to keep the flow-rate as low as possible (to avoid pressure on the glass membrane of the electrode, which could alter the pH readings). Alternatively, for measurement of the pH course in a micro-column, where fractions of only 60 μ l are collected, Fredriksson²⁰ described the use of a Radiometer micro glass electrode, in which the pH-sensitive membrane is shaped as a horizontal capillary. The capillary can be filled, via a vertical polyethylene tube, simply by suction. A volume as small as 20 μ l is sufficient for pH reading. Values correct to within 0.01 pH unit have been reported.

(C) pH measurements after IEF in gel media

When performing IEF in solid support media, it is convenient to be able to measure directly the pH gradient along the gel length. For flat gels, a surface electrode may be used with a coplanar glass reference electrode having either a flat face or a spear tip. In either case, it is advisable to have the probes of both electrodes as close as possible to minimize errors due to differences in conductivity throughout the gel.

One of the first electrodes developed for pH measurements on gel surfaces is the antimony micro-electrode, in conjunction with a calomel reference electrode, reported by Beeley *et al.*²¹. This electrode, which was originally developed by Kleinberg²² to measure the pH of dental plaques, has a spear tip of 1 mm in diameter. Measurements are made by placing the gel over a sheet of graph paper, pressing the reference electrode at any position along the gel length and scanning at regular intervals with the antimony micro-electrode. Where the pH reading has been made, the gel is marked by serration with a scalpel blade. This allows a much greater accuracy in the determination of pI by interpolation on a graph of pH *versus* protein position in the gel. With the antimony electrode, readings of electromotive force are obtained on the millivolt scale and are converted into units of pH by means of an appropriate calibration graph. This type of electrode might be better standardized against Ampholine solutions, which in turn are calibrated against standard buffers²³. One important property of the antimony micro-electrode, in contrast to very small glass electrodes, is its rapid equilibration time (less than 10 sec), even at low temperatures, which makes it very attractive for use at 4°. However, one disadvantage of this electrode is its low reproducibility, given by Beeley *et al.*²¹ as < 0.25 pH units, in comparison with the resolution of 0.02 pH units, and even less, afforded by IEF.

Alternatively, on gel slabs, the pH can be measured directly on the gel surface with a flat membrane electrode, such as LOT Type 403-30-M8 from Ingold. However, since this electrode has a high surface area (membrane diameter, 8 mm), Drysdale²⁴ (in collaboration with Ingold Inc., Lexington, Mass., U.S.A.) has developed a flat membrane micro-electrode. This h-shaped electrode contains the reference unit in one arm, and the measuring membrane in a parallel arm. The two electrodes are coplanar, and are only 3 mm apart in order to offset variations in conductivity along the gel. The pH membrane has a diameter of 2.5 mm. This allows accurate pH measurements over a small surface; however, due to the high impedance of the small glass membrane, this electrode is practically of no use at low temperatures, since the response times are extremely long.

In order to overcome the problems connected with the use of surface electrodes, we have devised a simple method for accurate pH measurements in gel slabs, using a standard combination micro-electrode. The gel slab is focused over plastic graph paper placed on the cooling block of an LKB Multiphor 2117 apparatus, thermostatted at 4°. At the end of the experiment, 22 or 23 gel segments are cut from the side of the gel, at 0.5-cm intervals. Simultaneously, a 1-2-mm diameter hole is bored in the gel (with the help of a gel puncher for immunodiffusion and of a suction pump) in a zone corresponding to the middle of the 0.5-cm gel segment removed for pH measurement. Since this operation is made on the cooling block, and requires only a few minutes, no appreciable diffusion of the protein zones takes place. The rectangular gel segments removed are eluted in small test-tubes with 0.3 ml of 10 mM

KCl. Since the combination micro-electrode, with a saturated solution of KCl and buffers, is kept in the cold room (connected to the electronic control unit outside by a cable passing through the wall), accurate and reproducible pH measurements are obtained from the gel eluates at the focusing temperature. The pH profile can be easily superimposed on the stained gel slab with the aid of the 22 or 23 holes punched in it.

(D) Use of pH markers

An alternative to pH measurements along the gel length, especially when working with thin-layer equipment, is the use of a calibrated mixture of pH markers,

TABLE 1
pH MARKERS FOR ISOELECTRIC FOCUSING

<i>pH marker</i>	<i>pI at 25°</i>	<i>pI at 4°[‡]</i>
<i>Proteins</i>		
Cytochrome <i>c</i>	9.28 ± 0.02*	
Ribonuclease	8.88 ± 0.03*	
Myoglobin (sperm whale):		
major component	8.18 ± 0.02* [‡] ; 8.18 ± 0.04**	
minor component	7.68 ± 0.02*	
Myoglobin (horse):		
major component	7.33 ± 0.01* [‡] ; 7.45 ± 0.04**	7.58 ± 0.02
minor component	6.88 ± 0.02* [‡] ; 7.15 ± 0.04**	7.22 ± 0.05
Bovine haemoglobin A	6.80***	
Carbonic anhydrase (bovine)		6.18 ± 0.02
Conalbumin	5.88 ± 0.02*	
β-Lactoglobulin B	5.31*	5.45 ± 0.02
β-Lactoglobulin A	5.14 ± 0.01*	5.35
Bovine insulin	5.32 ± 0.02***	
Albumin (bovine) Cohn:		
fraction 5	4.90***	4.95 ± 0.02
Ovalbumin	4.70***	
Horse spleen ferritin I	4.50 ± 0.02*	
Horse spleen ferritin II	4.38 ± 0.02*	
Horse spleen ferritin III	4.23 ± 0.03*	
<i>Dyes</i>		
Tris(5-hydroxy-1,10-phenanthroline)iron(II)	7.15**	
Tris(5-hydroxy-1,10-phenanthroline)iron(II) -tris(4-hydroxy-1,10-phenanthroline)- iron(II)	6.82**	
Tris(4-hydroxy-1,10-phenanthroline)iron(II) -tris(5-hydroxy-1,10-phenanthroline)- iron(II)	6.24**	
Congo Red	5.80***	
Tris(4-hydroxy-1,10-phenanthroline)iron(II)	5.45**	
Evans blue	5.35***	
Methyl Blue	3.60***	
Fast green FCF (major component)	3.05***	
Patent blue V	3.00***	

* Ref. 27.

** Ref. 26.

*** Ref. 25.

[‡] Ref. 10.

pI AND M.W. VALUES OF PROTEINS

pI = Acclimated, chloro. = chloroplast, cyto. = cytoplasm, lyso. = lysosome, micro. = microsome, mitoch. = mitochondria, n.g. = not given, nucl. = nucleus, perox. = peroxisome, r.t. = room temperature, s.p.c. = single polypeptide chain and str. = strain.

Protein	Source	Organ and/or subcellular location	M.W.	Subunit		pI	No. of iso-enzymes	Temperature (°C)
				No.	M.W.			
Acetoacetyl-CoA thiolase ^{28,29}	yeast	cytosol mitoch.	1.4 · 10 ⁵ 0.65 · 10 ⁵			5.3 7.8	1 1	n.g. n.g.
Acetoacetyl-CoA thiolase ³⁰	rat	liver cyto. liver mitoch.				4.8 8.4	1 1	n.g. n.g.
Acetoacetyl-CoA thiolase ³⁰	ox	liver cyto. liver mitoch.				5.2 7.8	1 1	n.g. n.g.
Acetoacetyl-CoA thiolase ³¹	rat	brain cyto. brain mitoch.				5.1 8.3	1 1	n.g. n.g.
β -N-Acetyl glucosaminidase ³²	<i>Sclerotinia fructigena</i>		141000			3.76	1	n.g.
β -N-Acetyl glucosaminidase ^{33,34}	jack bean meal		100000			4.8	1	r.t.
β -N-Acetyl glucosaminidase ³⁵	sheep	testis				6.0, 6.3	2	5
N-Acetyl hexosaminidase ³⁶	ram	testis	140000			6.45, 5.85*	3	5
	ram	epididymis	140000			7.15, 7.05*, 6.7, 6.1	4	5
N-Acetyl muramyl-L-alanine amidase ³⁷	<i>Staphylococcus aureus</i>		30000			9.7	1	4
Acetyl transferase A ³⁸	rat	thymus (nucl.)	99000			5.90	1	n.g.
Acetyl transferase B ₁ ³⁸	rat	thymus (nucl.)	110000			4.75	1	n.g.
Acetyl transferase B ₂ ³⁸	rat	thymus (nucl.)	92000			4.75	1	n.g.
Acid β -galactosidase ^{39,40}	rat	intestine	95000			4.2*, 4.6, 5.4*, 6.1, 8.0	5	1.5
Acid phosphatase ⁴¹	rat	liver	100000			7.7, 4.5	2	4
Acid phosphatase ⁴²	<i>S. aureus</i>	str. 408				9.58*, 8.40	2	n.g.
	<i>S. aureus</i>	str. 8325				9.47*, 8.42	2	n.g.
Acid protease ⁴³	<i>Rhizopus chitrensis</i>		35000			5.2*, 5.9	2	n.g.
Acid ribonuclease ⁴⁴	Lymphosarcoma P1798		19000			5.3	1	n.g.
Aconitase ⁴⁵	pig	heart	66000			8.5*, 8.1, 7.9	3	n.g.

ISOELECTRIC POINTS AND MOLECULAR WEIGHTS OF PROTEINS

Aconitate hydratase ⁶⁶	pig	heart cyto.				5.4	1	0
	pig	heart mitoch.				7.4	1	0
Adenine phosphoribosyl trans-ferase ^{67,68}	mouse	fibroblast				5.5	1	r.t.
	human	fibroblast	34000	3	11000	4.48	1	r.t.
	human	erythrocyte	32000			4.70, 4.83*, 4.94, 5.06	4	n.g.
Adenosine deaminase ⁶⁹								
Adenylyate kinase ^{50,51}	rat	liver	46000	2	23000	7.5, 8.0	2	n.g.
Adenylyate kinase ⁵²	<i>Escherichia coli</i>		27000		s.p.c.	4.95	1	n.g.
Albumin ^{53,54}	ox	plasma	69000		s.p.c.	5.18, 5.07, 4.98	3	25
Albumin ^{54,55}	human	serum	69000		s.p.c.	5.85*	3	n.g.
Alcohol dehydrogenase ^{56,57}	horse	liver	80000	2	41000	8.08, 8.28, 8.51, 8.7*, 9.29*	12	n.g.
Alcohol dehydrogenase ⁵⁸	rat	liver	68000	2	39000	9.7	1	0
Aldehyde dehydrogenase ⁵⁹	horse	liver	245000	4	57000	5.05*, 4.80	2	n.g.
Aldolase ^{60,61}	yeast		80000	2	40000	5.1, 5.2*, 5.3	3	n.g.
Aldolase ⁶²	human	muscle	160000	4	40000	9.13, 9.30*, 9.4*, 9.59*, 9.73	5	4
Aldolase ⁶²	hen	muscle	160000	4	40000	9.87, 9.96*, 10.0*, 10.06*, 10.11	5	4
Aldolase ⁶²	pigeon	muscle	160000	4	40000	9.9, 9.97*, 10.0*, 10.09*, 10.15	5	4
Aldolase ⁶²	rabbit	muscle	160000	4	40000	9.1, 9.27, 9.42*, 9.57*, 9.71	5	4
Aldolase ⁶²	rat	muscle	160000	4	40000	9.1, 9.25, 9.39*, 9.53*, 9.70*	5	4
Aldolase ⁶²	yeast		61000	1	23000	5.05	1	n.g.
Alkaline phosphatase ⁶⁵	human	placenta	116000	2	58000	4.6	1	n.g.
Alkaline phosphatase ⁶⁶	human	kidney	140000	2	69000	4.3, 4.8, 5.9	3	n.g.
Alkaline phosphatase ^{66,67}	calf	intestine	80000	2	40000	4.4	1	n.g.
Alkaline phosphatase ⁶⁸	human	liver	80000	2	40000	3.9	1	n.g.
Alkaline phosphatase ^{69,70}	<i>E. coli</i>		135000	2	70000	5.06, 5.17, 5.20, 5.26*, 5.38	5	25
L-Amino acid oxidase ^{71,72}	<i>Crotalus adamanteus</i>	venom	135000	2	70000	(5.2-8.4) 5.60*, 5.71*, 5.87*	18	4

(Continued on p. 8)

TABLE 2 (continued)

Protein	Source	Organ and/or subcellular location	M.W.	Subunit		pI	No. of iso-enzymes	Temperature (°C)
				No.	M.W.			
α -Amylase ^{3,74}	human	parotid	55000		5.95	1	4	
α -Amylase ^{4,75}	human	saliva	55000		6.4, 6.1, 5.6, 5.3	4	22	
α -Amylase ⁶	rat	pancreas	56000	s.p.c.	8.77*, 8.95*	2	n.g.	
α -Amylase ⁷⁷	guinea pig	pancreas	150000	2	8.4	1	5-10	
Anthraniolate synthetase ^{76,79}	<i>Serratia marcescens</i>		210000	2	4.6	1	n.g.	
Arginine esterase ⁸⁰	<i>Contortrix latichctus</i>	venom	30000		9.1	1	n.g.	
Aryl β -glucosidase ^{81,82}	<i>Aspergillus niger</i>		50000		4.21, 4.43*	2	4	
Aryl β -glucosidase ⁸²	<i>Stereum sanguinolentum</i>		50000		4.00, 4.08	2	4	
Aryl β -glucosidase ⁸¹	<i>Fomes annosus</i>		50000		4.08, 4.17	2	4	
Aryl β -glucosidase ⁸²	<i>Chrysosporium lignorum</i>		50000		4.22, 4.38, 4.44	3	4	
Arylesterase ⁸³	guinea pig	cerebral cortex	180000, 78000		5.1*, 5.8*	2	n.g.	
Arylsulphatase ⁸⁴	<i>Pseudomonas aeruginosa</i>			60000	4.85*, 4.9*	2	n.g.	
Arylsulphatase ⁸⁵	rat	kidney			4.4, 6.2	2	n.g.	
Arylsulphatase A ⁸⁶	red kangaroo	liver	100000		5.1, 5.4	2	n.g.	
Arylsulphatase B ⁸⁶	red kangaroo	liver	45000		7.0	1	n.g.	
L-Asparaginase ^{87,88}	<i>Erwinia carotovora</i>							
L-Asparaginase ^{89,90}	yeast		135000	4	8.00	1	r.t.	
Asparagine synthetase ⁹¹	<i>E. coli</i>		133000	4	5.35	1	5	
Aspartate aminotransferase ^{92,93}	<i>E. coli</i>		80000		5.5	1	n.g.	
Aspartate aminotransferase ⁹⁴	pig	heart	92500		5.35, 5.41, 5.43, 5.53, 5.69*	5	25	
Aspartate aminotransferase ⁹⁴	sheep	liver	89000		5.35, 5.56, 5.75	3	n.g.	
Aspartic β -semialdehyde dehydrogenase ⁹⁵	yeast		156000	4	6.17	1	n.g.	
Aspartokinase L-homoserine dehydrogenase ⁹⁶	<i>E. coli</i>	str. K12	360000	4	6.1	1	n.g.	
Biotin carboxyl-carrier protein ⁹⁷	<i>E. coli</i>		45000	2	4.5	1	n.g.	
Bromelain ⁹⁸	<i>Ananas comosus</i>	stem	22000		9.7	1	25	

Butyrylcholinesterase ⁹⁹	human	brain	300000- 400000	(5.6-7.0)	4-5	22
Carbonic anhydrase ¹⁰⁰	bull shark	erythrocyte	38000	4.5	1	n.g.
Carbonic anhydrase ¹⁰¹ (A, B, C, F, D, G, H, M, N, O, P, T, U, X)	human	erythrocyte	25400- 29700	(4.85-8.12)	14	n.g.
Carbonic anhydrase ¹⁰²	guinea pig	colonic mucosa	30000	5.2, 7.4	2	n.g.
Carbonic anhydrase ¹⁰³	pig	erythrocyte	30375	7.3	1	1
Carbonic anhydrase B ^{103,104}	ox		30000	5.89	1	22
Carboxylesterase E ₁ ¹⁰⁵	rat	liver micro.	150000	5.7	1	n.g.
Carboxylesterase E ₂ ¹⁰⁵	rat	liver micro.	70000	6.6	1	n.g.
Carboxylesterase E ₁ ¹⁰⁶	rat	liver micro.	177000	5.65	1	n.g.
Carboxylesterase ¹⁰⁷	beef	liver	55000	5.5	1	n.g.
Carboxypeptidase G ₁ ¹⁰⁸	<i>Pseudomonas</i> <i>sittleri</i>		92000	7.1*	4	n.g.
Carnitine acetyltransferase ¹⁰⁹	rabbit	heart, brain, muscle		6.1, 6.7, 7.9*	3	n.g.
Carnitine acetyltransferase ¹⁰⁹	human	adrenal		6.0, 6.8, 7.4*	3	n.g.
Carnitine acetyltransferase ¹⁰⁹	human	heart		6.1, 6.7, 7.6*	3	n.g.
Carnitine acetyltransferase ¹⁰⁹	human	brain		6.1, 6.6, 7.5*	3	n.g.
Carnitine acetyltransferase ¹⁰⁹	pigeon	muscle		7.0, 5.85*, 7.8*	3	n.g.
Catalase ¹¹⁰	mouse	liver perox., liver cyto.		6.7	1	6
				8.0, 8.2*, 8.5	3	6
				4.7*, 4.9*, 5.1	3	n.g.
Catechol oxidase ¹¹¹ (tyrosinase)	<i>Vitis vinifera</i>			(4.5-5.5) 5.0*	6	n.g.
Cathepsin B ₁ ¹¹²	human	liver	25500	4.27, 4.43*, 4.7*	5	4
Cellulase ¹¹³	<i>Chrysosporium</i> <i>lignarum</i>			5.05*, 5.53	3	4
	<i>Aspergillus niger</i>			3.9, 4.2, 4.52*		
	<i>St. sanguinolentii</i>			3.58	1	4
	<i>Fomes annosus</i>			3.9, 4.2*	2	4
Cellulase ¹¹⁴	<i>Trichoderma</i> <i>koningii</i>			3.8, 3.95*	2	5
Cellulase (C ₁ component) ¹¹⁴	human	plasma	95000	3.0	1	n.g.
Ceramide trihexosidase ¹¹⁵	human	brain	35700	4.74	1	n.g.
Cerebropuprein ¹¹⁶	human					

(Continued on p. 10)

TABLE 2 (continued)

Protein	Source	Organ and/or subcellular location	M.W.	Submit No.	M.W.	pI	No. of iso-enzymes	Temperature (°C)
Cereolysin ¹⁷	<i>Bacillus cereus</i>					6.5	1	n.g.
Choline acetyltransferase ¹¹⁸	pigeon	brain				6.6	1	4-6
Choline acetyltransferase ¹¹⁹	guinea pig	brain				6.8	1	4-6
Choline acetyltransferase ¹¹⁸	rat	brain				7.5, 7.8, 8.3*	3	4-6
Choline acetyltransferase ¹¹⁸	cat	brain				7.0, 7.6, 8.4*	3	4-6
Cholinesterase ¹¹⁹	<i>Pleuronectes platessa</i>							
	ox	muscle				5.3, 6.9, 7.5	3	n.g.
α -Chymotrypsin ^{120,121}		pancreas	21600			8.38, 8.76*	2	n.g.
δ -Chymotrypsin ¹²²	<i>Pseudomonas aeruginosa</i>							
	beef	pancreas	25000			8.5	1	4
Chymotrypsinogen A ¹²³	ox	pancreas	23600	s.p.c.		8.8, 9.2, 9.6*	3	n.g.
Chymotrypsinogen A ^{73,124}	guinea pig	pancreas	25850	s.p.c.		8.97	1	4
Chymotrypsinogen ⁷⁷	trout	pancreas				4.8, 8.7*	2	5-10
Citrate synthase ¹²⁵		liver (cold accl.)				5.76	1	20
		liver (warm accl.)				5.05	1	20
Clostridio peptidase B ¹²⁶	<i>Clostridium perfringens</i>							
	human	serum	44700			4.85	1	18
Coagulation factor VIII ¹²⁷	human					5.6	1	n.g.
Cobalophillin (Vitamin B ₁₂ -binding protein) ¹²⁸	human	saliva	69000	s.p.c.		(4.5-5)	6	r.t.
Cobalophillin (Vitamin B ₁₂ -binding protein) ¹²⁹	human	amniotic fluid	59300	s.p.c.		3.0, 3.3, 3.7*, 3.9	4	r.t.
Colicin E ₂ ¹³⁰	<i>E. coli</i>		62000			7.41, 7.63*	2	22
Colicin E ₃ ¹³⁰	<i>E. coli</i>		62000			6.64	1	22
Co-lipase ¹³¹	pig	pancreas	9650	s.p.c.		5.0	1	n.g.
Concanavalin A ^{132,133}	jack bean		71000	4		4.5, 4.7, 5.05*, 5.5	4	n.g.
Creatine phosphokinase ^{67,134}	rabbit		81000	2		6.6, 6.7, 6.9	3	r.t.
Crotoxin ¹³⁵	rattle snake	venom	9000	s.p.c.		3.7	1	n.g.
	snake		12000	s.p.c.		8.6	1	n.g.
	cow	lens	2.7-10 ⁶ (α_1)	21500		4.5, 4.85*	1	n.g.
α -Crystallin ¹³⁶⁻¹³⁸			620000 (α_2)				2	4

β -Crystallin ¹²⁶⁻¹³⁸	cow	lens	220000 (β_H) 80000 (β_L)	27500	(5.95-7.0) 6.55* 6.80*, 7.0*	10	4
γ -Crystallin ^{137,138}	cow	lens		20500	7.25, 7.35, 7.6*, 7.80	4	4
Cyclic AMP dependent protein kinase ¹³⁹	rat	liver			7.6, 8.2*, 8.5*	3	n.g.
Cyclic nucleotide phosphodiesterase ¹⁴⁰	pea	seedlings	350000		4.3, 4.6, 4.8	3	n.g.
Cystathione synthase ^{141,142}	rat	liver	250000	2	5.5	1	n.g.
				2			
	chicken	liver	125000		6.0	1	n.g.
Cytochrome <i>b</i> -555 ¹⁴³	house fly	larvae	11300	s.p.c.	4.24*, 4.28*, 4.32	3	0-2
Cytochrome <i>b</i> ₅ reductase ¹⁴⁴	cat	liver	43000	s.p.c.	6.8	1	n.g.
Cytochrome <i>c</i> ¹⁴⁵	horse	heart			9.0, 9.4*	2	n.g.
Cytochrome <i>c</i> ¹⁴⁶	beef	heart	12200	s.p.c.	10.37, 10.57, 10.80*	4	4
Cytochrome <i>c</i> ¹⁴⁷	<i>Ustilago</i>						
	<i>sphaerogena</i>		12500	s.p.c.	9.4	1	n.g.
	<i>Spirillum</i>						
Cytochrome <i>c</i> ₅₀ ¹⁴⁸	<i>iterosnii</i>		10800	s.p.c.	9.86	1	4
	<i>Azotobacter vinelandii</i>		12000	s.p.c.	4.63 (reduced) 5.17 (oxidized)	1	15
Cytochrome <i>c</i> ₅₅ ¹⁴⁹	<i>Azotobacter vinelandii</i>		12000	s.p.c.	4.07 (reduced) 4.26 (oxidized)	1	15
	<i>Pseudomonas putida</i>		45000	s.p.c.	4.55	1	20
Cytochrome <i>c</i> peroxidase ¹⁵¹	<i>Pseudomonas fluorescens</i>				6.5, 6.7*	2	9
	house fly		120000	4	6.3, 6.9, 7.4*	3	n.g.
DDT dehydrochlorinase ^{152,153}	<i>S. aureus</i>				2.5*, 5.0, 6.2, 8.5, 10.1*	5	4
Deoxyribonuclease ¹⁵⁴							
	<i>Aspergillus carneus</i>		71000		4.12*, 4.35	2	n.g.
Dextranase ¹⁵⁵	<i>Cytophaga johnsonii</i>		60000		5.1, 5.35*, 5.76*	3	4
Dextranase ¹⁵⁶							

(Continued on p. 12)

TABLE 2 (continued)

Protein	Source	Organ and/or subcellular location	M.W.	Submit		pI	No. of iso-enzymes	Temper-ature (°C)
				No.	M.W.			
N,O-Diacetyl muraminidase ¹⁵⁷	<i>Chalaropsis Aerobacter aerogenes</i>		19000		s.p.c.	7.53	1	20
Diacetyl reductase ¹⁵⁸	<i>Drosophila melanogaster</i>	embryos	100000	4	25000	5.75, 5.9, 6.55*, 6.8*	4	0
DNA polymerase ¹⁵⁹	<i>E. coli</i>					5.3*, 5.8*, 6.3*, 7.5	4	n.g.
DNA polymerase I ¹⁵⁹	ox	plasma	38000			5.2	1	n.g.
Endonuclease ⁶⁰	<i>E. coli</i>					7.3	1	0
Endonuclease ⁶¹	<i>Mitrococcus lysodeikticus</i>		13500			4.3*, 6.1*, 8.5	3	15
Endonuclease ⁶²	yeast		88000	2	44000	4.8*, 9.0*	2	25
Enolase ^{63,163}	<i>S. aureus</i>		27800		s.p.c.	5.4, 5.6*, 6.0, 6.6, 6.7*, 6.8, 7.1	7	n.g.
Enterotoxin A ¹⁶⁴	<i>S. aureus</i>		24500			6.64, 7.26*, 7.68*, 8.14	4	4
Enterotoxin B ¹⁶⁵	<i>S. aureus</i>		24500			7.85, 7.93, 8.25*, 8.55*	4	24
Enterotoxin B ¹⁶⁶	<i>S. aureus</i>		25000		s.p.c.	8.5, 9.05*, 9.4*, 9.56	4	4
Epidermolytic toxin ¹⁶⁷	<i>Laticauda semifasciata</i>	venom	7000		s.p.c.	6.2, 7.05*	2	n.g.
Erbutoxin C ¹⁶⁸	red kidney bean		150000	4	37000	9.24	1	n.g.
Erythroagglutinin ¹⁶⁹	<i>Linnibicus human</i>	blood	3450000	148	23230	6.5	1	n.g.
Erythrocytin ¹⁷⁰	beef	erythrocyte	33600			5.05	1	20
Esterase ¹⁷⁰	horse	liver	55000			4.75	1	n.g.
Ferritin ^{171,173}	human	spleen	440000	24	18500	5.5	1	n.g.
Ferritin ¹⁷⁴	rabbit	liver				(4.27-4.57) 4.27*, 4.35*, 4.40*	9	20
Ferritin ¹⁷⁵	human	liver				5.29, 5.35, 5.45*	5	n.g.
Ferritin ¹⁷⁶	rabbit	liver				5.50, 5.55	5	n.g.
α-Fetoprotein ¹⁷⁷	human	foetus	72000			4.98, 5.00*, 5.15, 5.18	4	n.g.
Fibrinogen ¹⁷⁸	rabbit	plasma				5.16, 5.24*, 5.3	5	20
						4.78*, 5.2	2	25
						5.45, 5.50, 5.55	3	n.g.

Follicle stimulating hormone (FSH) ¹⁷⁹	rat	hypophysis				2.8	1	n.g.
Fructose-1,6-diphosphatase ¹⁸⁰	chicken	breast				8.6	1	n.g.
Fructose-1,6-diphosphatase I and II ¹⁸¹	chicken	liver				8.1	1	n.g.
Fructose diphosphate aldolase ¹⁸²	<i>Paralithodes cambratica</i>	gill				7.5, 5.3*	2	n.g.
	<i>Euglena gracilis</i>	hypodermis				7.2*, 5.4	2	n.g.
		chloro. cyto.				4.6	1	n.g.
		heart				5.7	1	n.g.
Fumarate ¹⁸³	pig		194000	4	48500	(5.9-8.2) 7.58*, 7.73*, 7.90*	11	n.g.
α -D-Galactosidase ¹⁸⁴	<i>Aspergillus niger</i>					4.1, 4.5*	2	n.g.
β -D-Galactosidase ¹⁸⁴	<i>Aspergillus niger</i>					4.3	1	n.g.
β -D-Galactosidase ¹⁸⁵	<i>Neurospora crassa</i>		96000			4.78*, 5.82	2	n.g.
β -1,3-Glucan hydrolase ¹⁸⁶	<i>Nicotiana glutinosa</i>		45000			4.87	1	n.g.
Glucose-6-phosphate dehydrogenase ¹⁸⁷	<i>Candida utilis</i>		110000	2	55000	5.50, 5.87, 6.54*	3	n.g.
Glucose-6-phosphate dehydrogenase ¹⁸⁸	human	erythrocyte				5.79, 5.94, 6.11, 6.6*, 7.04*	6	i.t.
β -D-Glucosidase ¹⁸⁹	pig	kidney	50000			4.9*, 5.1	2	n.g.
β -Glucosidase ¹⁹¹	<i>Aspergillus niger</i>					4.46	1	4
β -Glucosidase ¹⁹²	<i>Chrysosporium lignorum</i>					4.38	1	4
β -Glucosidase A ₃ ¹⁹⁰	<i>Aspergillus wentii</i>		170000		s.p.c.	3.8	1	n.g.
Glucosinolate ¹⁹¹	<i>Sinapis alba</i>					5.0, 5.55	2	20
Glucosyl transferase ¹⁹²	parsley	cell culture	50000			5.0	1	n.g.
β -Glucuronidase ¹⁹³	rat	kidney				4.6, 6.7	2	n.g.
β -Glucuronidase ¹⁹³	rat	preputials	280000	4	75000	6.15	1	20
L-Glutamate-phenylpyruvate aminotransferase ¹⁹⁴	beef	kidney (mitoch.)				9.8	1	n.g.
L-Glutaminase ¹⁹⁵	<i>E. coli</i>					4.2	1	n.g.
Glutaminase-asparaginase ¹⁹⁶	Achromobacteraceae		138000	4	35000	8.3	1	n.g.
L-Glutamylcyclotransferase ¹⁹⁷	sheep	brain				4.65, 5.25, 5.45*, 5.75*, 6.20	5	n.g.

(Continued on p. 14)

TABLE 2 (continued)

Protein	Source	Organ and/or subcellular location	M.W.	Subunit		pI	No. of iso-enzymes	Temperature (°C)
				No	M.W.			
L-Glutamylcyclotransferase ¹⁹⁷	human	brain				4.06*, 4.25	2	n.g.
Glutathione epoxide transferase ¹⁹⁸	rat	liver	40000	2	24000	7.14	1	4
Glutathione-S-transferase ¹⁹⁹	rat	liver	40000			6.9*, 8.1*	2	n.g.
Glyceraldehyde-3-phosphate dehydrogenase ^{95,200}	yeast		144700	4	35000	6.25	1	n.g.
Glyceralddehyde-3-phosphate dehydrogenase ²⁰¹	rabbit	liver, muscle	140000	4	36000	8.32, 8.52*	2	n.g.
L-Glycerol-3-phosphate dehydrogenase ⁶⁰	rabbit	adipose tissue	66000			6.35	1	n.g.
	rabbit	skeletal muscle	78000	2	37500	6.45	1	n.g.
α_1 -Glycoprotein ²⁰²	bull frog	serum	206000			4.5	1	9
GSH transferase ²⁰³	sheep	liver				7.1, 9.5	2	n.g.
	rat	liver				7.6, 8.0*, 8.6*, 9.4	4	n.g.
	mouse	liver				8.2*, 9.8	2	n.g.
	house fly					5.15*, 5.9, 6.8	3	n.g.
	cockroach					4.5, 6.6*, 8.5	3	n.g.
	grass grub					4.5, 5.2*, 8.8	3	n.g.
Guanine phosphorybosyl transferase ²⁰⁴	<i>E. coli</i>					5.5	1	4
Gulonolactonase ²⁰⁵	rat	kidney	42000		s.p.c.	5.20	1	4
Haemoglobin A ^{206,207}	human	erythrocytes	64500	4	16000	6.95	1	r.t.
Haemoglobin C ^{206,207}	human	erythrocytes	64500	4	16000	7.40	1	r.t.
Haemoglobin S ^{206,207}	human	erythrocytes	64500	4	16000	7.25	1	r.t.
Haemoglobin F ₁ ^{206,207}	human	foetus	64500	4	16000	6.90	1	r.t.
Haemoglobin F _{II} ^{206,207}	human	foetus	64500	4	16000	7.15	1	r.t.
Haemoglobin A (α -chain) ²⁰⁸	human	erythrocytes			16000	8.12	1	4
Haemoglobin S (β -chain) ²⁰⁸	human	erythrocytes			16000	7.08	1	4
Haemoglobin ²⁰⁹	harp seal	blood				7.06*, 8.04	2	n.g.
α -Haemolysin (α -toxin) ²¹⁰	<i>S. aureus</i>	str. V8	44000			5.0, 7.0, 8.5*	3-4	4
β -Haemolysin ²¹¹	<i>S. aureus</i>	str. R1				3.5, 9.5*	2	4
δ -Haemolysin (δ -lysin) ^{211,212}	<i>S. aureus</i>	str. R1	74000			9.6	1	4
Hemerythrin ²¹³	<i>Dendrostromum pyroides</i>		107000	8	13000	6.71	1	n.g.

Hepatocuprein ¹¹⁶	human	liver	35600		4.76	1	n.g.
Hexokinase ^{114,215}	yeast		102000	2	4.7	1	n.g.
Hyaluronate lyase ¹¹⁶	<i>S. aureus</i>	str. M18			7.4, 7.9*	2	4
Hyaluronate lyase ¹¹⁷	<i>S. aureus</i>	str. V8			6.4*, 8.2*	4	4
L-3-Hydroxyacyl coenzyme A dehydrogenase ^{219,219}	pig	heart	65000	2	8.95	1	4
Hypoxanthine-guanine phosphoribosyltransferase ²²⁰	human	erythrocytes	68000	2	5.66, 5.82, 6.00	3	5
Insulin ^{73,221}	beef		11466	2	5.72	1	4
Insulin ⁸⁷	pig				6.0	1	r.t.
Invertase ²²²	yeast				(3.4-4.4) 4.02*, 4.24*	10	4
Invertase ²²³	<i>Neurospora crassa</i>		210000	4	4.78	1	n.g.
Isocitrate dehydrogenase ²²⁴	baker's yeast		375000	10	5.5	1	n.g.
Isocitrate lyase ²²⁵	<i>Neurospora crassa</i>				5.5	1	n.g.
2-Keto-3-deoxy-6-phosphogluconate aldolase ²²⁶	<i>Pseudomonas putida</i>		73300	3	4.7	1	n.g.
α -Ketoglutarate-glyoxylate carboligase ²²⁷	<i>Mycobacterium phlei</i>				5.6	1	n.g.
Laccase A ^{228,229}	<i>Polyporus versicolor</i>		62000		3.07, 3.27	2	4
Laccase B ²²⁸	<i>Polyporus versicolor</i>				5.70*, 5.9, 6.13*, 6.26, 6.36, 6.56	6	4
β -Lactamase (penicillinase) ^{230,231}	<i>S. aureus</i>	str. 408	56000		8.38, 9.25*	2	n.g.
Lactase ²²²	rat	str. 8325			8.48, 9.55*	2	n.g.
Lactate dehydrogenase ²²⁵	chinese hamster	intestine			4.4	1	n.g.
Lactate dehydrogenase ⁶⁰	rabbit	skeletal muscle			5.7	1	n.g.
Lactate dehydrogenase ²³³	<i>Bacillus stearothermophilus</i>	muscle			8.3, 8.4, 8.55*	3	n.g.
β -Lactoglobulin ^{67,234}	cow	milk	135000	4	5.5	1	n.g.
β -Lactoglobulin A ²³⁵	cow	milk	35000	2	5.1, 5.26*, 5.34*	3	r.t.
β -Lactoglobulin B ²³⁵	cow	milk			5.13	1	25
					5.23	1	25

(Continued on p. 16)

TABLE 2 (continued)

Protein	Source	Organ and/or subcellular location	M.W.	Subunit		pI	No. of iso-enzymes	Temperature (°C)
				No.	M.W.			
Lactoperoxidase ²³⁶	cow	milk	85000			9.16, 9.32, 9.49, 9.68*, 9.7*, 9.80*	6	n.g.
Leghaemoglobin ²³⁷	<i>Lupin serradella</i>	root nodules	20000		s.p.c.	5.08, 5.13	2	n.g.
Leucoagglutinin ²³⁸	kidney bean	root nodules	18000		s.p.c.	5.00, 5.05	2	n.g.
Leucoagglutinin ⁶⁹	kidney bean		126000	4	31000	5.1	1	n.g.
Leucyl-t-RNA synthetase ²³⁹	kidney bean		140000	4	35000	5.0	1	n.g.
Lipase ²⁴⁰	yeast		120000	2	60000	4.9	1	n.g.
	<i>Corynebacterium acnes</i>				80000	3.8	1	n.g.
Lipase 177	guinea pig	pancreas			66000	5.1	1	5-10
Lipase 277	guinea pig	pancreas			49700	8.1	1	5-10
Liponamide dehydrogenase ²⁴¹	pig	heart	100000			5.6, 6.0, 6.5*, 6.8*	4	n.g.
Lipoxydase (lipoxygenase) ^{242, 243}	soybean		108000	2	54000	5.65	1	25
Lipoxygenase ^{243, 244}	soybean		108000	2	54000	5.68, 6.15, 6.26	3	n.g.
Lipoxygenase ²⁴⁵	pea		70000			5.80*, 5.82*	2	n.g.
Lipoxygenase ²⁴⁶	pea		70000			5.85, 5.92, 5.97, 6.01, 6.06, 6.17	6	25
Lipoyldehydrogenase ²⁴⁷	pig	heart				5.9, 6.1, 6.3, 6.6, 6.8*, 7.0, 7.2	7	n.g.
Luciferase ²⁴⁸	firefly		100000	2	50000	5.7, 6.4*	2	n.g.
Luteinizing hormone (LH) ^{249, 250}	rat	hypophysis	31000	2	15500	9.0*	4	n.g.
Lysin ²⁵¹	<i>Tegula pfeifferi</i>	egg membrane	8800			7.9	1	n.g.
Lysozyme ²⁵²	human	serum				10.0*, 11.0	2	4
Malic enzyme ²⁵³	<i>Ascaris suum</i>	muscle	248000	4	66400	6.63	1	5
Malonyl CoA-ACP transacylase ²⁵⁴	<i>E. coli</i>		36600		s.p.c.	4.65	1	n.g.
Mannanase ²⁵⁵	<i>Aspergillus niger</i>					3.95	1	4
	<i>St. sanguinolentum</i>					3.58	1	4
	<i>Fomes annosus</i>					3.9*, 4.2	2	4
	<i>Clryosporium lignorum</i>					4.11	1	4
	<i>Aspergillus niger</i>					4.68	1	4
Mannosidase ²⁵⁶	soybean		175000			4.5, 4.75*	2	n.g.

ISOELECTRIC POINTS AND MOLECULAR WEIGHTS OF PROTEINS

Methionyl t-RNA synthetase ²⁵⁵	<i>Bacillus brevis</i>					4.4	1	n.g.
Methylsterase ²⁵⁷	tomato	100000				5.0*	1	n.g.
Mevalonic kinase ²⁵⁸	hog	200000		pectin		8.4	1	20
Myoglobin ²⁵⁹	horse	26300		liver		4.7	1	n.g.
		98000		muscle		6.47, 6.53, 6.60,		
		17500	s.p.c.			6.79, 6.83, 6.92,		
						7.17*	7	20
Myoglobin ⁷³	horse	17500	s.p.c.	muscle		6.76, 7.16	2	4
Myoglobin ⁶	horse	17500	s.p.c.	muscle		6.8, 6.86, 7.27, 7.76	4	4
Myoglobin ⁶⁷	horse	17500	s.p.c.	muscle		6.8, 7.0, 7.1*, 7.5*	4	r.t.
Myoglobin ²⁵⁹	ox	17500	s.p.c.	muscle		6.56, 6.74, 7.01*	3	20
Myoglobin ⁷³	whale			sperm		7.52, 7.79	2	4
Myoglobin ⁶⁷	whale			sperm		7.1, 7.4, 7.6, 7.68*		
						8.01*	5	r.t.
Myosin ²⁶⁰	Acanthamoeba	180000	1			8.0*	3	n.g.
			1					
			1					
Myrosinase C ²⁶¹	rapeseed	135000	2			4.96, 4.99, 5.06*	3	n.g.
	white mustard	151000	2			5.08	1	n.g.
NADH dehydrogenase ²⁶²	<i>Rhodospirillum rubrum</i>	33000				4.80*, 4.94	2	20
		26000				4.50	1	20
NADP isocitrate dehydrogenase ²⁶³	rainbow trout			liver (2°)		5.75	1	n.g.
				liver (17°)		5.72*, 6.9	2	n.g.
				salivary gland		4.50, 9.30*	2	n.g.
Nerve growth factor ^{264, 265}	mouse	26500	2			4.80	1	n.g.
Neuraminidase ²⁶⁶	<i>Vibrio cholerae</i>					4.95	1	n.g.
Neuraminidase ²⁶⁶	<i>Cl. perfringens</i>	200000	4			5.2, 5.35, 5.5*	6	20
Neuraminidase ²⁶⁷	Influenza virus	69800		A ₂ /1957		5.8*, 6.2, 6.5	1	r.t.
						4.9	1	r.t.
Neuraminidase ²⁶⁸	Pneumococcus			polymorphonuclear leucocytes (lyso.)		4.2*, 4.8, 5.2*, 6.3,	7	n.g.
Neutral protease ²⁶⁹	rabbit					7.0, 8.2, 9.1		
Nitrate reductase ²⁷⁰	spinach	240000				3.5, 4.9*	2	n.g.
Nitrogenase ²⁷¹	<i>Klebsiella pneumoniae</i>	218000	2			5.0	1	n.g.
		66800	2			4.0*, 5.9	3	n.g.

(Continued on p. 18)

TABLE 2 (continued)

Protein	Source	Organ and/or subcellular location	M.W.	Subunit		pI	No. of iso-enzymes	Temperature (°C)
				No.	M.W.			
Nucleoside diphosphokinase ²⁷²	human	erythrocyte	85000			5.4, 5.8*, 6.3 6.8, 7.3, 8.3	6	n.g.
Nucleotide phosphotransferase ²⁷³	<i>E. coli</i>		29000			6.8, 7.1*	2	n.g.
Ornithine aminotransferase ²⁷⁴	rat	liver	132000	4	33000	5.38	1	6
Ornithine transcarbamylase ²⁷⁵	human	liver				3.2*, 4.4	2	0
Ornithine transcarbamylase ²⁷⁶	ox	liver	108000	3	37800	6.19, 6.36, 6.44, 6.49, 6.59, 6.77, 6.95*	7	22
Ovotransferrin ²⁷⁷	hen	egg	80000			5.62, 5.78*, 6.05, 6.25*, 6.50, 6.05, 6.25*, 6.50, 6.73*, 5.7*, 6.3, 6.7*	6	20
Oxoacyl CoA thiolase ²⁷⁸	ox	liver (mitoch.)				6.95, 8.15*	3	n.g.
Palmitoyl CoA synthetase ²⁷⁹	rat	liver (mitoch.)	168000	6	27000	5.5	1	n.g.
Parvalbumins ²⁸⁰	rat	liver	11900			4.45	1	r.t.
	<i>Raja clavata</i>	muscle	12300			4.65	1	r.t.
	<i>Raja montagui</i>	muscle	33000			2.86*, 2.94	2	4
Pepsin ^{281, 282}	hog	gastric juice	45900			4.0, 8.8, >11	4	n.g.
Peroxidase ²⁸³	horseradish	root	40000			(3.9-9) 6.1*, 6.6*, 7.2*	20	20
Peroxidase ^{284, 285}	horseradish							
Phosphonacetyl glucosamine mutase ²⁸⁶	pig	submaxillary gland	110000			4.8	1	r.t.
Phosphodiesterase ²⁸⁷	sugar beet	leaves				3.85	1	n.g.
Phosphoenolpyruvate carboxylase ^{288, 289}	<i>E. coli</i>	muscle	402000	4	99600	4.92	1	21
Phosphoglucosmutase ²⁹⁰	rabbit		64900		s.p.c.	6.8*	4	n.g.
6-Phosphogluconate dehydrogenase ²⁹¹	<i>Neurospora crassa</i>		115000	2	57000	4.93, 5.50*	2	10
Phosphoglucose isomerase ²⁹²	human	erythrocyte	125000	2	62000	9.0, 9.1, 9.2*	3	n.g.
Phospholypase A ²⁹³	honey bee	venom	40000	2	19500	10.5	1	n.g.
Phospholypase A ²⁹⁴	<i>Naja naja</i>	venom	8500-					
			20200			(4.60-5.66) 4.95*	11	n.g.
Phospholypase A ²⁹⁴	<i>Vipera russellii</i>	venom	15000-			(4.62-9.90) 9.52*		
			23800			9.90*	7	n.g.

ISOELECTRIC POINTS AND MOLECULAR WEIGHTS OF PROTEINS

Phospholypase A ²⁹⁵	<i>Laticauda semi-fasciata</i>	venom	11000	6.6	1	n.g.
Phospholypase A ²⁹⁶	<i>Agkistrodon halys blomhoffii</i>		13800	4.0*, 10.0*	2	20
Phospholypase C ¹⁷	<i>Cl. perfringens</i>		47000	5.2*, 5.5*	2	n.g.
Phosphomannanase ²⁹⁷	<i>Bacillus circulans</i>	heart		5.9, 6.8*	2	n.g.
Phosphorylase a ²⁹⁸	pig	heart		5.45, 5.9, 6.3	3	n.g.
Phosphorylase b ²⁹⁸	pig	heart		5.5, 5.9, 6.25, 6.35*	4	n.g.
C-Phycocyanin ²⁹⁹	<i>Oscillatoria agardhii</i>		26300	4.65	1	3
Plasminogen ³⁰⁰	human	plasma	81000	(6.4-8.5) 7.5*, 7.8*, 8.1*	8	r.t.
Procarboxypeptidase A ³⁰¹	human	pancreatic juice		6.22, 6.37, 6.56, 6.72	4	r.t.
Procarboxypeptidase A ⁷⁷	guinea pig	pancreas	45000	4.6, 4.8*	2	5-10
Procarboxypeptidase B ⁷⁷	guinea pig	pancreas	47700	6.6	1	5-10
Progesterone binding globulin ³⁰²	guinea pig	serum	82800	3.5, 4.4	2	4
Progesterone binding plasma protein ³⁰³	guinea pig	plasma	77500	3.6*	3	n.g.
Proelastase ⁷⁷	guinea pig	pancreas	28000	6.9, 7.5	2	5-10
Prolactin ³⁰⁴	human	plasma, amniotic fluid, pituitary	27500	6.53	1	n.g.
Prolamine kinase ³⁰⁵	rat	brain		5.3, 5.8, 7.4, 8.2	4	n.g.
Protense ³⁰⁶	<i>Bacteroides amylophilus</i>		60000	4.25*	2	n.g.
Protense ³⁰⁷	<i>Aspergillus niger</i>					
Prothrombin ³⁰⁸	ox	blood			3	0
Protocollagen proline hydroxylase ³⁰⁹	chick	embryo	248000	4.4	1	n.g.
Purine nucleoside phosphorylase ³¹⁰	human	erythrocyte	84000	6.0*, 6.5, 7.0, 7.5, 7.8	5	n.g.
Pyridine nucleotide dehydrogenase ³¹¹	human	erythrocyte	28000	6.2*, 7.45	2	n.g.

(Continued on p. 20)

Protein	Source	Organ and/or subcellular location	M.W.	Subunit		pI	No. of iso-enzymes	Temperature (°C)
				No.	M.W.			
IMP: Pyrophosphate phospho-ribosyl transferase ¹⁰ Pyruvate kinase ³¹²	human rat	erythrocyte liver	68000			(5.5-7.5) 6.0*, 6.5* 5.50*, 5.75*, 6.0, 6.75	9	n.g.
	rat	muscle				6.75	4	n.g.
Pyruvate kinase ³¹⁵	oyster	mantle				6.75	1	n.g.
	oyster	adductor muscle				6.35*	3	n.g.
Ribonuclease ⁷⁷	guinea pig	pancreas	13500			5.6, 6.5*	3	n.g.
Ribose 5-phosphate isomerase ³¹⁴	pea	leaf (chloro.) leaf (cyto.)		s.p.c.		8.7, 8.8 4.95*	2	5-10
	human	HeLa cells	500000			4.75*	3	0
RNA polymerase II ³¹⁵	human	HeLa cells	500000			4.75	2	0
				2200000, 1400000, 35000, (25000), (170000)			1	n.g.
Serine sulphhydrase ¹⁴¹	chicken	liver	125000			6.0	1	n.g.
Seryl t-RNA synthetase ³¹⁶	<i>E. coli</i>	str. B	103000	2	53000	6.15	1	n.g.
Seryl t-RNA synthetase ³¹⁷	<i>E. coli</i>	str. K12	100000	2	50000	3.9	1	n.g.
Staphylokinase ³¹⁷	<i>S. aureus</i>					6.5	1	4
Subtilisin ³¹⁶						8.4*	4	4
Subtilopeptidase ¹²²	<i>Pseudomonas aeruginosa</i>							
Succinate thiokinase ³¹⁹	pig	heart	78000			7	1	4
						5.8, 5.9, 6.0, 6.12*, 6.36	5	n.g.
Sulphatase A ³²⁰	human	liver (lyso.)	400000			4.7, 4.83, 4.98	3	n.g.
Sulphatase A ^{321,322}	ox	liver	107000			3.6	1	n.g.
Sulphatase B ³²¹	ox	liver	25000	2	50000	8.3	1	n.g.
Superoxide dismutase ³²³	ox	liver	73000			7.9	1	n.g.
Thromboplastin ³⁰⁸	ox	serum				4.1	3	0
Thymidine kinase ³²⁴	monkey	kidney (cyto.)	85000			9.8	1	n.g.
	monkey	kidney (mitoch.)	75000			6.0, 9.0, 9.8*	3	n.g.
Thymidine kinase ³¹³	mouse	LM cells (cyto.)	86000			8.8, 9.0*	2	4
	human	LM cells (mitoch.) HeLa cells (cyto.)	69000 86000			5.0*, 8.1 9.6	2 1	4 4
Thyroglobulin ³²⁶	hog	HeLa cells (mitoch.) thyroid gland	69500			5.6, 8.4, 9.7* 4.5*	3 3-4	4 22

Thyroid stimulating hormone (TSH) ³²⁷	sea whale	pituitary				8.17, 8.38*, 8.51*, 8.70	4	n.g.
Thyrotropin (TSH) ^{326, 329}	ox	pituitary	2	32000	15000	8.28, 8.58*, 8.68*, 8.8	4	25
Thyroxine binding globulin ³³⁰	human	serum				(4.2-5.2) 4.3*, 4.4*, 4.6*	9	n.g.
α -Toxin A ³³¹	<i>S. aureus</i>			28000	s.p.c.	7.2	1	n.g.
α -Toxin B ³³¹	<i>S. aureus</i>			28000	s.p.c.	8.4	1	n.g.
Transcobalamin II ³³²	human	serum		36000		6.25, 6.45	2	r.t.
Transferrin (apo) ³³³	rabbit	serum				6.0	1	n.g.
Transferrin (1 Fe ³⁺) ³³³	rabbit	serum				5.5	1	n.g.
Transferrin (2 Fe ³⁺) ³³³	rabbit	serum				5.0	1	n.g.
Triosephosphate isomerase ³³⁴	human	erythrocyte		56000		5.5, 6.0*, 6.4	3	3
Triosephosphate isomerase ^{334, 335}	rabbit	muscle	2	53000	26500	5.8, 6.2, 7.0*, 7.7, 8.0	5	3
Triosephosphate isomerase ³³⁶	pea	leaf (chloro.)				4.75*	3	0
Triosephosphate isomerase ⁶²	<i>Euglena gracilis</i>	leaf (cyto.)				4.8	1	0
		cytoplasm		58000		4.8	1	n.g.
		chloroplast		58000		4.45	1	n.g.
Trypsin and chymotrypsin inhibitor ³³⁷	groundnut	pancreas		7500	s.p.c.	8.5	1	n.g.
Trypsinogen ⁷⁷	guinea pig				24500	8.7*, >9.3	2	5-10
Tryptophanyl t-RNA synthetase ^{338, 339}	<i>E. coli</i> B			74000		6.2	3	0
Tryptophanyl t-RNA synthetase ³⁴⁰	human	placenta	2	120000		5.2	1	0
Tyrosine α , β -amino mutase ³⁴¹	<i>Bacillus brevis</i>	str. Vm4		75000	s.p.c.	4.6	1	n.g.
L-Tyrosine methyl ester sulphotransferase ³⁴²	rat	liver				5.3*, 5.8, 6.6	3	n.g.
Urease ³⁴³	jack bean			485000		4.88	1	n.g.
UDP-glucosyl transferase ³⁴⁴	parsley			50000		5.0	1	n.g.
Vitamin B ₁₂ -binding protein ³⁴⁵	human	amniotic fluid		59300	s.p.c.	3.3, 3.7*, 3.9	3	n.g.
Xylanase ⁶²	<i>Aspergillus niger</i>					3.9*, 4.5*	2	4
	<i>St. saugitrolentum</i>					3.62, 4.30*	2	4
	<i>Fomes annosus</i>					4.1, 4.6, 7	3	4
	<i>Chrysosporium lignarium</i>					4.44, 6.0*	2	4

* The pI value of the major component. When seven or more isozymes are given for a single entry, the range of pI is given in parentheses, followed by the pI values of only the major component(s). Thus, in the case of L-amino acid oxidase, which is resolved into 18 isozymes, the values (5.2-8.4) represent the pI range of the 18 components, while the values 5.60, 55.71 and 5.87 represent the pI values of the three major components.

covering the pH range of interest. Ideally, these pH markers should be coloured proteins or dyes. In the case of protein markers, they should not be heterogeneous, or at least have a major, easily identifiable, band in order to avoid confusion in pH assessments. Studies on the use of pH markers have been reported by Conway-Jacobs and Lewin²⁵, Nakhleh *et al.*²⁶, Bours¹⁰ and Radola²⁷. We have listed the most common protein and dye markers in Table 1. This table will be followed by the compilation of data on pI and M.W. values of proteins.

In these tables, the most accessible references are given for each entry; they do not necessarily indicate the source most deserving of credit for the macromolecular parameters given. Some references might have been overlooked. We would appreciate comments from other laboratories on missing references and data or on possible errors in these tables. We hope that this compilation will prove useful for teaching and research purposes.

2. ACKNOWLEDGEMENT

This project was supported by research grants CT74.00255.04 and No. 75.00686.115.4756 from Consiglio Nazionale delle Ricerche (C.N.R.), Italy.

3. SUMMARY

Proteins with known isoelectric points (pI), as determined by isoelectric focusing, have been tabulated. When available, the native molecular weight as well as subunit molecular weight and stoichiometry have been reported. For each entry, the source and, when applicable, the organ of origin and/or subcellular location are given.

The pI values are listed together with the respective temperature of pH measurement. Moreover, in the case of proteins displaying microheterogeneity, the major components have been indicated. The table lists a total of approximately 800 pI values.

REFERENCES

- 1 D. W. Darnall and I. M. Klotz, *Arch. Biochem. Biophys.*, 166 (1975) 651.
- 2 V. R. Kleine, *Fortschr. Arzneimitt. Forsch.*, 16 (1972) 365.
- 3 H. Svensson, *Acta Chem. Scand.*, 15 (1961) 325.
- 4 H. Svensson, *Acta Chem. Scand.*, 16 (1962) 456.
- 5 H. Svensson, *Arch. Biochem. Biophys.*, Suppl. 1 (1962) 132.
- 6 O. Vesterberg and H. Svensson, *Acta Chem. Scand.*, 20 (1966) 820.
- 7 P. G. Righetti and J. W. Drysdale, *J. Chromatogr.*, 98 (1974) 271.
- 8 O. Vesterberg, *Sv. Kem. Tidskr.*, 80 (1968) 213.
- 9 S. P. L. Sorensen, K. Linderstrom-Lang and E. Lund, *C.R. Trav. Lab. Carlsberg*, 16 (1926) 5.
- 10 J. Bours, *Sci. Tools*, 20 (1973) 29.
- 11 O. Vesterberg, *Ann. N.Y. Acad. Sci.*, 209 (1973) 23.
- 12 H. Davies, *Prot. Biol. Fluids*, 17 (1970) 389.
- 13 S. Fredriksson, *Ph. D. Thesis*, Chalmers Institute of Technology, University of Gothenburg, Gothenburg, Sweden, 1975.
- 14 E. J. King, *Acid-Base Equilibria*, Pergamon Press, Oxford, 1965.
- 15 J. T. Edsall and J. Wyman, *Biophysical Chemistry*, Vol. I, Academic Press, New York, 1958, p. 471.
- 16 N. Ui, *Biochim. Biophys. Acta*, 229 (1971) 567.
- 17 M. Jonsson, E. Pettersson and H. Rilbe, *Acta Chem. Scand.*, 23 (1969) 1553.
- 18 C. Secchi, *Anal. Biochem.*, 51 (1973) 448.
- 19 A. J. A. Strongin, A. P. Balduiev and E. D. Levin, *Sci. Tools*, 20 (1973) 34.

- 20 S. Fredriksson, *Anal. Biochem.*, 50 (1972) 575.
- 21 J. A. Beeley, S. M. Stevenson and J. G. Beeley, *Biochim. Biophys. Acta*, 285 (1972) 293.
- 22 I. Kleinberg, *Brit. Dent. J.*, 104 (1958) 197.
- 23 P. G. Righetti, *Separ. Purif. Methods*, 4 (1975) 23.
- 24 J. W. Drysdale, personal communication.
- 25 A. Conway-Jacobs and L. M. Lewin, *Anal. Biochem.*, 43 (1971) 394.
- 26 E. T. Nakhleh, S. A. Samra and Z. L. Awdeh, *Anal. Biochem.*, 49 (1972) 218.
- 27 B. J. Radola, *Biochim. Biophys. Acta*, 295 (1973) 412.
- 28 J. A. Kornblatt and H. Rudney, *J. Biol. Chem.*, 246 (1971) 4417.
- 29 J. A. Kornblatt and H. Rudney, *J. Biol. Chem.*, 246 (1971) 4424.
- 30 B. Middleton, *Biochem. J.*, 125 (1971) 69p.
- 31 B. Middleton, *Biochem. J.*, 125 (1971) 70p.
- 32 F. Reyes and R. J. W. Byrde, *Biochem. J.*, 131 (1973) 381.
- 33 Y. T. Li and S. C. Li, *J. Biol. Chem.*, 243 (1968) 3994.
- 34 Y. T. Li and S. C. Li, *J. Biol. Chem.*, 245 (1970) 5153.
- 35 B. G. Winchester, M. Caffrey and D. Robinson, *Biochem. J.*, 121 (1971) 161.
- 36 S. Bullock and B. G. Winchester, *Biochem. J.*, 133 (1973) 593.
- 37 T. Wadström and O. Vesterberg, *Acta Pathol. Microbiol. Scand.*, 79 (1971) 248.
- 38 D. Gallwitz and I. Sures, *Biochim. Biophys. Acta*, 263 (1972) 315.
- 39 N. G. Asp, *Biochem. J.*, 117 (1970) 369.
- 40 J. Krome, O. Koldowsky, A. Heringova, V. Jirsova, K. Kacel, M. Ledrina and H. Pelichova, *Biochem. J.*, 114 (1969) 621.
- 41 M. Igarashi and V. P. Hollander, *J. Biol. Chem.*, 243 (1968) 6084.
- 42 W. Schaeg, R. Bingöl and H. Blobel, *Biochim. Biophys. Acta*, 268 (1972) 542.
- 43 D. Tsuru, A. Hattori, H. Tsuji and J. Fukumoto, *J. Biochem. (Tokyo)*, 67 (1970) 415.
- 44 S. Biswas and V. P. Hollander, *J. Biol. Chem.*, 244 (1969) 4185.
- 45 C. Kennedy, R. Rauner and O. Gawron, *Biochem. Biophys. Res. Commun.*, 47 (1972) 740.
- 46 R. Z. Zanes and E. Kun, *Biochim. Biophys. Acta*, 227 (1971) 204.
- 47 T. Kusano, C. Long and H. Green, *Proc. Nat. Acad. Sci. U.S.A.*, 68 (1971) 82.
- 48 C. B. Thomas, W. J. Arnold and W. N. Kelley, *J. Biol. Chem.*, 248 (1973) 2529
- 49 R. A. Qsborne and N. Spencer, *Biochem. J.*, 133 (1973) 117.
- 50 V. Sapić, G. Litwack and W. E. Criss, *Biochim. Biophys. Acta*, 258 (1972) 436.
- 51 W. E. Criss, V. Sapić and G. Litwack, *J. Biol. Chem.*, 245 (1970) 6346.
- 52 R. K. Holmes and M. F. Singer, *J. Biol. Chem.*, 248 (1973) 2014.
- 53 A. R. Williamson, M. R. Salaman and H. R. Kreth, *Ann. N.Y. Acad. Sci.*, 209 (1973) 210.
- 54 J. T. Edsall, in H. Neurath and F. Bailey (Editors), *The Proteins*, 1st ed., Academic Press, New York, 1953, p. 549.
- 55 E. Valmet, *Prot. Biol. Fluids*, 17 (1970) 401.
- 56 U. M. Lutstorf, P. M. Schuerch and J. P. van Wartburg, *Eur. J. Biochem.*, 17 (1970) 497.
- 57 H. Theorell and D. Winer, *Arch. Biochem. Biophys.*, 83 (1959) 291.
- 58 Y. Tokuma and H. Terayama, *Biochem. Biophys. Res. Commun.*, 54 (1973) 341.
- 59 R. I. Feldman and H. Weiner, *J. Biol. Chem.*, 247 (1972) 260.
- 60 W. A. Susor, M. Kochman and W. J. Rutter, *Science*, 165 (1969) 1260.
- 61 C. E. Harris, R. D. Kobes, D. C. Teller and W. J. Rutter, *Biochemistry*, 8 (1969) 2442.
- 62 Y. Ikehara, S. Yanagi and T. Kamiya, *J. Biochem. (Tokyo)*, 72 (1972) 203.
- 63 G. H. Sheys, W. J. Arnold, J. A. Watson, J. A. Hayashi and C. C. Doughty, *J. Biol. Chem.*, 246 (1971) 3824.
- 64 G. H. Sheys and C. C. Doughty, *Biochim. Biophys. Acta*, 235 (1971) 414.
- 65 M. Usategui-Gomez, F. M. Yeager and P. Tarbutton, *Clin. Chim. Acta*, 50 (1974) 405.
- 66 A. L. Latner, M. Parsons and A. W. Skillen, *Enzymologia*, 40 (1970) 1.
- 67 M. Fosset, D. Chapelet-Tordo and M. Lazdunski, *Biochemistry*, 13 (1974) 1783.
- 68 A. L. Latner, M. E. Parson and A. W. Skillen, *Biochem. J.*, 118 (1970) 299.
- 69 H. Csopak, M. Jonsson and B. Hallberg, *Acta Chem. Scand.*, 26 (1972) 2412.
- 70 A. Garen and C. Levinthal, *Biochim. Biophys. Acta*, 38 (1960) 470.
- 71 M. B. Hayes and D. Wellner, *J. Biol. Chem.*, 244 (1969) 6636.
- 72 A. Dekok and A. B. Rawitch, *Biochemistry*, 8 (1969) 1405.
- 73 J. A. Beeley, S. M. Stevenson and J. G. Beeley, *Biochim. Biophys. Acta*, 285 (1972) 293.
- 74 H. Mutzbauer and G. V. Schulz, *Biochim. Biophys. Acta*, 102 (1965) 526.
- 75 S. Scharpè, A. Lauwers and W. Coreman, *Biochimie*, 55 (1973) 1003.

- 76 T. G. Sanders and W. J. Rutter, *Biochemistry*, 11 (1972) 130.
- 77 G. A. Scheele, *J. Biol. Chem.*, 250 (1975) 5375.
- 78 F. Robb, M. A. Hutchinson and W. L. Belser, *J. Biol. Chem.*, 246 (1971) 6908.
- 79 H. Zalkin and L. H. Hwang, *J. Biol. Chem.*, 246 (1971) 6899.
- 80 P. M. Toom, T. N. Solie and A. T. Tu, *J. Biol. Chem.*, 245 (1970) 2549.
- 81 E. Ahlgren, K. E. Eriksson and O. Vesterberg, *Acta Chem. Scand.*, 21 (1967) 937.
- 82 E. Ahlgren and K. E. Eriksson, *Acta Chem. Scand.*, 21 (1967) 1193.
- 83 H. B. Bosman, *Biochim. Biophys. Acta*, 276 (1972) 180.
- 84 G. Delisle and F. H. Milazzo, *Biochim. Biophys. Acta*, 212 (1970) 505.
- 85 A. Goldstone and H. Koning, *Biochem. J.*, 141 (1974) 527.
- 86 A. B. Roy, *Biochim. Biophys. Acta*, 227 (1971) 129.
- 87 M. R. Salaman and A. R. Williamson, *Biochem. J.*, 122 (1971) 93.
- 88 K. A. Cammack, D. I. Mariborough and D. S. Miller, *Biochem. J.*, 126 (1972) 361.
- 89 H. K. Robinson, *Anal. Biochem.*, 49 (1972) 353.
- 90 B. H. Frank, A. H. Pekar, A. J. Veros and P. P. K. Ho, *J. Biol. Chem.*, 245 (1970) 3716.
- 91 H. Cedar and J. H. Schwartz, *J. Biol. Chem.*, 244 (1969) 4112.
- 92 G. Marino, M. DeRosa, V. Buonocore and V. Scardi, *FEBS Lett.*, 5 (1969) 347.
- 93 M. Martinez-Carrion, C. Turano, E. Chiancone, F. Bossa, A. Giartosio, F. Riva and P. Fasella, *J. Biol. Chem.*, 242 (1967) 2397.
- 94 M. Campos-Carrires and E. A. Munn, *Biochem. J.*, 135 (1973) 683.
- 95 M. J. Holland and E. W. Westhead, *Biochemistry*, 12 (1973) 2276.
- 96 F. Falcoz-Kelley, J. Janin, J. C. Saari, M. Veron, P. Truffa-Bachi and G. N. Cohen, *Eur. J. Biochem.*, 28 (1972) 507.
- 97 R. R. Fall and P. R. Vagelos, *J. Biol. Chem.*, 247 (1972) 8005.
- 98 W. Berndt, U. Hoffmann and K. Mueller-Wieland, *Z. Gastroenterol.*, 6 (1968) 185.
- 99 J. B. Carlsen and O. Svensmark, *Biochim. Biophys. Acta*, 207 (1970) 477.
- 100 J. R. Maynard and J. E. Coleman, *J. Biol. Chem.*, 246 (1971) 4455.
- 101 S. Funakoshi and H. F. Deutsch, *J. Biol. Chem.*, 244 (1969) 3438.
- 102 M. J. Carter and D. S. Parsons, *Biochim. Biophys. Acta*, 206 (1970) 190.
- 103 R. B. Ashworth and T. E. Spencer, *Arch. Biochem. Biophys.*, 142 (1971) 122.
- 104 M. Jonsson and E. Pettersson, *Acta Chem. Scand.*, 22 (1968) 712.
- 105 A. Ljungquist and K. B. Augustinsson, *Eur. J. Biochem.*, 23 (1971) 303.
- 106 R. Arndt, E. Heymann, W. Junge, K. Krisch and H. Hollandt, *Eur. J. Biochem.*, 36 (1973) 120.
- 107 D. Wynne and Y. Shalitin, *Arch. Biochem. Biophys.*, 154 (1973) 199.
- 108 J. L. McCullough, B. A. Chabner and J. R. Bertino, *J. Biol. Chem.*, 246 (1971) 7207.
- 109 H. L. White and J. C. Wu, *Comp. Biochem. Physiol.*, 48B (1974) 575.
- 110 R. Feinstein and C. Peraino, *Biochim. Biophys. Acta*, 214 (1970) 230.
- 111 P. Coggon, G. A. Moss and G. W. Sanderson, *Phytochemistry*, 12 (1973) 1947.
- 112 A. J. Barret, *Biochem. J.*, 131 (1973) 809.
- 113 K. E. Eriksson and W. Rzedowski, *Arch. Biochem. Biophys.*, 129 (1969) 683.
- 114 T. M. Wood and S. I. McCrae, *Biochem. J.*, 128 (1972) 1183.
- 115 C. A. Mapes, C. H. Suelter and C. C. Sweely, *J. Biol. Chem.*, 248 (1973) 2471.
- 116 R. J. Carrico and H. F. Deutsch, *J. Biol. Chem.*, 244 (1969) 6087.
- 117 A. W. Bernheimer, P. Grushoff and L. S. Avigad, *J. Bacteriol.*, 95 (1968) 2439.
- 118 D. Malthe-Sorensen and F. Fonnum, *Biochem. J.*, 127 (1972) 229.
- 119 U. Brodbeck, R. Gentinetta and S. J. Lundin, *Acta Chem. Scand.*, 27 (1973) 561.
- 120 N. Ui, *Biochim. Biophys. Acta*, 229 (1971) 582.
- 121 G. W. Schwert and S. Kaufman, *J. Biol. Chem.*, 190 (1951) 807.
- 122 S. Arvidson and T. Wadström, *Biochim. Biophys. Acta*, 310 (1973) 418.
- 123 D. Bobb, *Ann. N.Y. Acad. Sci.*, 209 (1973) 225.
- 124 J. Wilcox, K. Kraut, T. Wade and H. Neurath, *Biochim. Biophys. Acta*, 24 (1957) 72.
- 125 P. W. Hochachka and J. K. Lewis, *J. Biol. Chem.*, 245 (1970) 6567.
- 126 W. M. Mitchell, *Biochim. Biophys. Acta*, 178 (1969) 194.
- 127 A. Gladhaug and H. Prydz, *Biochim. Biophys. Acta*, 215 (1970) 105.
- 128 R. Gräsbeck, K. Visuri and U. Hakan Stenman, *Biochim. Biophys. Acta*, 263 (1972) 721.
- 129 U. Hakan Stenman and R. Gräsbeck, *Biochim. Biophys. Acta*, 286 (1972) 243.
- 130 H. R. Herschman and D. R. Helinski, *J. Biol. Chem.*, 242 (1967) 5360.
- 131 C. Erlanson and B. Borgström, *Biochim. Biophys. Acta*, 271 (1972) 400.
- 132 G. Entlicher, J. V. Kostir and J. Kocourek, *Biochim. Biophys. Acta*, 236 (1971) 795.

- 133 M. O. J. Olson and I. E. Liener, *Biochemistry*, 6 (1967) 3801.
- 134 P. M. Bayley and A. R. Thomson, *Biochem. J.*, 104 (1967) 33p.
- 135 J. Horst, R. A. Hendon and H. Fraenkel-Conrat, *Biochem. Biophys. Res. Commun.*, 46 (1972) 1042.
- 136 H. Bloemendal and P. Herbrink, *Ophthalm. Res.*, 6 (1974) 81.
- 137 J. Bours, *Exp. Eye Res.*, 16 (1973) 501.
- 138 J. Bours, *J. Chromatogr.*, 60 (1971) 225.
- 139 L. J. Chen and D. A. Walsh, *Biochemistry*, 10 (1971) 3614.
- 140 B. P. Lin and J. E. Varner, *Biochim. Biophys. Acta*, 276 (1972) 454.
- 141 A. E. Braunstein, E. V. Goryachenkova, E. A. Tolosa and I. H. Willhardt, *Biochim. Biophys. Acta*, 242 (1971) 247.
- 142 S. Kashiwamata, Y. Kotake and D. M. Greenburg, *Biochim. Biophys. Acta*, 212 (1970) 501.
- 143 Y. Okada and K. Okunuki, *J. Biochem. (Tokyo)*, 67 (1970) 487.
- 144 L. Spatz and P. Strittmatter, *J. Biol. Chem.*, 248 (1973) 793.
- 145 A. Heaney and D. L. Weller, *J. Chem. Educ.*, 47 (1970) 724.
- 146 T. Flatmark and O. Vesterberg, *Acta Chem. Scand.*, 20 (1966) 497.
- 147 S. N. Vinogradov, K. G. Bitar and K. W. Junk, *Biochim. Biophys. Acta*, 257 (1972) 273.
- 148 G. D. Clark-Walker and J. Lascelles, *Arch. Biochem. Biophys.*, 136 (1970) 153.
- 149 W. H. Campbell, W. H. Orme-Johnson and R. H. Burris, *Biochem. J.*, 135 (1973) 617.
- 150 K. Dus, M. Katagiri, C. A. Yu, D. L. Erbes and I. C. Gunsalus, *Biochem. Biophys. Res. Commun.*, 40 (1970) 423.
- 151 N. Ellfolk and R. Soininen, *Acta Chem. Scand.*, 24 (1970) 2126.
- 152 B. Goodchild and J. N. Smith, *Biochem. J.*, 117 (1970) 1005.
- 153 M. L. Dinamarca, L. Levenbook and E. Valdes, *Arch. Biochem. Biophys.*, 147 (1971) 374.
- 154 T. Wadström, *Biochim. Biophys. Acta*, 147 (1967) 441.
- 155 N. Hiraoka, J. Fukumoto and D. Tsuru, *J. Biochem. (Tokyo)*, 71 (1972) 57.
- 156 J. C. Janson, *Ph. D. Thesis*, University of Uppsala, Uppsala, Sweden, 1972.
- 157 W. M. Mitchell and J. H. Hash, *J. Biol. Chem.*, 244 (1969) 17.
- 158 Ø. Hetland, K. Bryn and F. C. Stormer, *Eur. J. Biochem.*, 21 (1971) 206.
- 159 L. H. Cohen, P. E. Penner and L. A. Loeb, *Ann. N.Y. Acad. Sci.*, 209 (1973) 354.
- 160 M. Anai, H. Haraguchi and Y. Takagi, *J. Biol. Chem.*, 247 (1972) 193.
- 161 S. M. Hadi, D. Kirtikar and D. A. Goldthwait, *Biochemistry*, 12 (1973) 2747.
- 162 H. Nakayama, S. Okubo and Y. Takagi, *Biochim. Biophys. Acta*, 228 (1971) 67.
- 163 K. G. Mann, F. J. Catellino and P. A. Hargrave, *Biochemistry*, 9 (1970) 4002.
- 164 E. J. Schantz, W. G. Roessler, M. J. Woodburn, J. M. Lynch, H. M. Jacoby, S. M. Silverman and J. C. Gorman and L. Spero, *Biochemistry*, 11 (1972) 360.
- 165 P. C. Chang and N. Dickie, *Biochim. Biophys. Acta*, 236 (1971) 367.
- 166 J. F. Metzger, A. D. Johnson and W. S. Collins, *Biochim. Biophys. Acta*, 257 (1972) 183.
- 167 J. P. Arbuthnot, B. Billcliffe and W. D. Thompson, *FEBS Lett.*, 46 (1974) 92.
- 168 N. Tamiya and H. Abe, *Biochem. J.*, 130 (1972) 547.
- 169 T. H. Weber, H. Aro and C. T. Nordman, *Biochim. Biophys. Acta*, 263 (1972) 94.
- 170 M. R. Rossi-Fanelli, E. Chiancone, P. Vecchini and E. Antonini, *Arch. Biochem. Biophys.*, 141 (1970) 278.
- 171 S. Hartz and H. F. Deutsch, *J. Biol. Chem.*, 244 (1969) 4565.
- 172 I. Urushizaki, Y. Niitsu, K. Ishitani, M. Matsuda and M. Fukuda, *Biochim. Biophys. Acta*, 243 (1971) 187.
- 173 R. R. Crichton, *New England J. Med.*, 284 (1971) 1413.
- 174 E. Alpert, R. L. Coston and J. W. Drysdale, *Nature (London)*, 242 (1973) 194.
- 175 B. K. van Kreel, H. G. van Eijk and B. Leijnse, *Acta Haematol.*, 47 (1972) 59.
- 176 I. Urushizaki, K. Ishitani and Y. Niitsu, *Biochim. Biophys. Acta*, 328 (1973) 95.
- 177 E. Alpert, J. W. Drysdale and K. J. Isselbacher, *Ann. N.Y. Acad. Sci.*, 209 (1973) 387.
- 178 S. Krantz, M. Lober and H. Fiedler, *FEBS Lett.*, 11 (1970) 100.
- 179 H. Kercret and J. Duval, *Biochimie*, 57 (1975) 85.
- 180 J. P. Olson and R. R. Marquardt, *Biochim. Biophys. Acta*, 268 (1972) 453.
- 181 P. W. Hochachka, *Biochem. J.*, 127 (1972) 781.
- 182 Y. Mo, B. G. Harris and R. W. Gracy, *Arch. Biochem. Biophys.*, 157 (1973) 580.
- 183 P. E. Penner and L. H. Cohen, *J. Biol. Chem.*, 246 (1971) 4261.
- 184 Y. C. Lee and V. Wacek, *Arch. Biochem. Biophys.*, 138 (1970) 264.
- 185 H. N. Johnson and A. G. DeBusk, *Arch. Biochem. Biophys.*, 138 (1970) 412.

- 186 A. E. Moore and B. A. Stone, *Biochim. Biophys. Acta*, 258 (1972) 238.
- 187 R. Chillas, K. M. Doering, G. F. Domagk and M. Rippa, *Arch. Biochem. Biophys.*, 159 (1973) 235.
- 188 V. M. Der Kaloustian, S. H. Idriss-Daouk, R. T. Hallal and Z. L. Awdeh, *Biochem. Genet.*, 12 (1974) 51.
- 189 H. E. Abrahams and D. Robinson, *Biochem. J.*, 111 (1969) 749.
- 190 G. Legler, M. von Radloff and M. Kempfle, *Biochim. Biophys. Acta*, 257 (1972) 40.
- 191 J. R. Vose, *Phytochemistry*, 11 (1972) 1649.
- 192 A. Sutter, R. Ortmann and H. Grisebach, *Biochim. Biophys. Acta*, 258 (1972) 71.
- 193 K. Ohtsuka and M. Wakabayashi, *Enzymologia*, 39 (1970) 109.
- 194 R. Scandurra, C. Cannella and R. Elli, *Sci. Tools*, 16 (1969) 17.
- 195 H. A. Campbell and L. T. Mashburn, *Biochemistry*, 8 (1969) 3768.
- 196 J. Roberts, J. S. Holcenberg and W. C. Dolowy, *J. Biol. Chem.*, 247 (1972) 84.
- 197 M. Orlowski, P. G. Richman and A. Meister, *Biochemistry*, 8 (1969) 1048.
- 198 T. A. Fjellsted, R. C. Allen, B. K. Duncan and W. B. Jakoby, *J. Biol. Chem.*, 248 (1973) 3702.
- 199 B. Gillham, *Biochem. J.*, 135 (1973) 797.
- 200 R. Jaenicke, D. Schmid and S. Knof, *Biochemistry*, 7 (1968) 919.
- 201 C. M. Smith and S. F. Velick, *J. Biol. Chem.*, 247 (1972) 273.
- 202 H. Nagano, T. Shimada and R. Shukuya, *J. Biol. Chem.*, 248 (1973) 3052.
- 203 A. G. Clark, J. N. Smith and T. W. Speiz, *Biochem. J.*, 135 (1973) 385.
- 204 R. L. Miller, G. A. Ramsey, T. A. Krenitsky and G. B. Elion, *Biochemistry*, 11 (1972) 4723.
- 205 S. H. Grossman and B. Axelrod, *J. Biol. Chem.*, 248 (1973) 4846.
- 206 J. W. Drysdale, P. G. Righetti and H. F. Bunn, *Biochim. Biophys. Acta*, 229 (1971) 42.
- 207 G. Braunitzer, K. Hilse, V. Rudloff and N. Hilschmann, *Advan. Protein Chem.*, 19 (1964) 1.
- 208 M. Perrella, D. Bresciani, G. Guglielmo, V. Russo, F. Rossi, L. Rossi-Bernardi, P. G. Righetti and J. V. Kilmartin, *Progr. Clin. Biol. Res.*, 1 (1975) 329.
- 209 H. J. Suderman, C. Yeh-Ku and K. Ronald, *Comp. Biochem. Physiol.*, 458 (1973) 515.
- 210 T. Wadström, *Biochim. Biophys. Acta*, 168 (1968) 228.
- 211 R. Möllby and T. Wadström, *Prot. Biol. Fluids*, 17 (1970) 465.
- 212 T. Yoshida, *Biochim. Biophys. Acta*, 71 (1963) 544.
- 213 R. E. Ferrell and G. B. Kitto, *Biochemistry*, 9 (1970) 3053.
- 214 B. Wurster and B. Hess, *Eur. J. Biochem.*, 36 (1973) 68.
- 215 J. R. Pringle, *Biochem. Biophys. Res. Commun.*, 39 (1970) 46.
- 216 O. Vesterberg, *Biochim. Biophys. Acta*, 168 (1968) 218.
- 217 O. Vesterberg, T. Wadström, K. Vesterberg, H. Svensson and B. Malmgren, *Biochim. Biophys. Acta*, 133 (1967) 435.
- 218 B. E. Noyes and R. A. Bradshaw, *J. Biol. Chem.*, 248 (1973) 3052.
- 219 B. E. Noyes, B. E. Glatthaar, J. S. Garavelli and R. A. Bradshaw, *Proc. Nat. Acad. Sci. U.S.*, 71 (1974) 1334.
- 220 W. J. Arnold and W. N. Kelley, *J. Biol. Chem.*, 246 (1971) 7398.
- 221 D. F. Waugh, *Advan. Protein Chem.*, 9 (1954) 325.
- 222 O. Vesterberg and B. Berggren, *Ark. Kemi*, 27 (1966) 119.
- 223 Z. D. Meachum, Jr., H. J. Calvin, Jr. and H. D. Braymer, *Biochemistry*, 10 (1971) 326.
- 224 J. A. Illingworth, *Biochem. J.*, 129 (1972) 1119.
- 225 D. L. Weller, A. Heaney and R. E. Sjogren, *Biochim. Biophys. Acta*, 168 (1968) 576.
- 226 R. H. Ammerstedt, H. Möhler, K. A. Decker and W. A. Wood, *J. Biol. Chem.*, 246 (1971) 2069.
- 227 H. Yamasaki and T. Moriyama, *Biochim. Biophys. Acta*, 242 (1971) 637.
- 228 M. Jonsson, E. Pettersson and B. Reinhammar, *Acta Chem. Scand.*, 22 (1968) 2135.
- 229 H. Mosbach, *Biochim. Biophys. Acta*, 73 (1963) 204.
- 230 W. Schnaeg, R. Bingöl and H. Blobel, *Biochim. Biophys. Acta*, 268 (1972) 542.
- 231 J. Pedersen, *J. Phys. Chem.*, 51 (1947) 164.
- 232 S. Schlegel-Haueter, P. Hore, K. R. Kerry and G. Semenza, *Biochim. Biophys. Acta*, 258 (1972) 506.
- 233 E. Kolb and J. I. Harris, *Biochem. J.*, 124 (1971) 76p.
- 234 R. Townsend and S. N. Timasheff, *J. Amer. Chem. Soc.*, 79 (1957) 3613.
- 235 S. Fredriksson, *Anal. Biochem.*, 50 (1972) 575.
- 236 A. Carlström and O. Vesterberg, *Acta Chem. Scand.*, 21 (1967) 271.
- 237 W. J. Broughton, M. J. Dilworth and C. A. Godfrey, *Biochem. J.*, 127 (1972) 309.
- 238 V. Räsänen, Th. H. Weber and R. Gräsbeck, *Eur. J. Biochem.*, 38 (1973) 193.

- 239 J. G. Chirikjian, K. Kanagalingam, E. Lau and J. R. Fresco, *J. Biol. Chem.*, 248 (1973) 1074.
240 G. S. Hassing, *Biochim. Biophys. Acta*, 242 (1971) 381.
241 M. L. Cohn and I. R. McManus, *Biochim. Biophys. Acta*, 276 (1972) 70.
242 N. Catsimpoolas, *Arch. Biochem. Biophys.*, 131 (1969) 185.
243 F. C. Stevens, D. M. Brown and L. E. Smith, *Arch. Biochem. Biophys.*, 136 (1970) 413.
244 J. P. Christopher, E. K. Pistorius and B. Axelrod, *Biochim. Biophys. Acta*, 284 (1972) 54.
245 C. E. Eriksson and S. G. Svensson, *Biochim. Biophys. Acta*, 198 (1970) 449.
246 S. Fredriksson, *Anal. Biochem.*, 57 (1974) 452.
247 J. E. Wilson, *Arch. Biochem. Biophys.*, 144 (1971) 216.
248 J. L. Denburg and W. D. McElroy, *Biochemistry*, 9 (1970) 4619.
249 H. Kercret and J. Duval, *Biochimie*, 57 (1975) 85.
250 D. N. Ward, L. E. Reichart, Jr., B. A. Fitak, H. S. Nahm, C. M. Sweeney and J. D. Neill, *Biochemistry*, 10 (1971) 1796.
251 K. Haino, *Biochim. Biophys. Acta*, 229 (1971) 459.
252 G. Lundblad, O. Vesterberg, R. Zimmerman and J. Lind, *Acta Chem. Scand.*, 26 (1972) 1711.
253 D. W. Fodge, R. W. Gracy and B. G. Harris, *Biochim. Biophys. Acta*, 268 (1972) 271.
254 F. E. Ruch and P. R. Vagelos, *J. Biol. Chem.*, 248 (1973) 8086.
255 M. Saita, T. Ikenaka and Y. Matsushima, *J. Biochem. (Tokyo)*, 70 (1971) 827.
256 A. P. Surguchev, I. G. Surgucheva, T. M. Ermokhima and G. N. Zaitseva, *Biochim. Biophys. Acta*, 224 (1970) 623.
257 H. Delincée and B. J. Radola, *Biochim. Biophys. Acta*, 214 (1970) 178.
258 E. Beytia, J. K. Dorsey, J. Marr, W. W. Cleland and J. W. Porter, *J. Biol. Chem.*, 245 (1970) 5450.
259 A. H. A. van den Oord, J. J. Wesdorp, A. F. van Dam and J. A. Verheij, *Eur. J. Biochem.*, 10 (1969) 140.
260 T. D. Pollard and E. D. Korn, *J. Biol. Chem.*, 248 (1973) 4682.
261 B. Lönnnerdal and J. Ch. Janson, *Biochim. Biophys. Acta*, 278 (1972) 175.
262 T. Horio, R. G. Bartsch, T. Kakuno and M. D. Kamen, *J. Biol. Chem.*, 244 (1969) 5899.
263 T. W. Moon and P. W. Hochachka, *Biochem. J.*, 123 (1971) 695.
264 V. Bocchini, *Biochim. Biophys. Acta*, 243 (1971) 420.
265 R. H. Angeletti and R. A. Bradshaw, *Proc. Nat. Acad. Sci. U.S.A.*, 68 (1971) 2417.
266 A. R. Neurath, R. W. Hartzell and B. A. Rubin, *Experientia*, 26 (1970) 1210.
267 A. P. Kendal, M. P. Kiley and E. A. Eckert, *Biochim. Biophys. Acta*, 317 (1973) 28.
268 W. L. Stahl and R. D. O'Toole, *Biochim. Biophys. Acta*, 268 (1972) 480.
269 P. Davies, G. A. Rita, K. Krakauer and G. Weissman, *Biochem. J.*, 123 (1971) 559.
270 B. A. Notton, E. J. Hewitt and A. H. Fielding, *Phytochemistry*, 11 (1972) 2447.
271 R. R. Eady, B. E. Smith, K. A. Cook and J. R. Postgate, *Biochem. J.*, 128 (1972) 655.
272 Y. C. Cheng, R. P. Agarwal and R. E. Parks, Jr., *Biochemistry*, 10 (1971) 2139.
273 E. F. Brunngraber and E. Chargaff, *Biochemistry*, 12 (1973) 3005.
274 C. Peraino, L. G. Bunville and T. N. Tahmisian, *J. Biol. Chem.*, 244 (1969) 2241.
275 S. Arashima and I. Matsuda, *Biochem. Biophys. Res. Commun.*, 45 (1971) 145.
276 M. Marshall and P. P. Cohen, *J. Biol. Chem.*, 247 (1972) 1641.
277 R. V. Wenn and J. Williams, *Biochem. J.*, 108 (1968) 69.
278 B. Middleton, *Biochem. J.*, 132 (1973) 717.
279 J. Bar-Tana and G. Rose, *Biochem. J.*, 131 (1973) 443.
280 C. Gerday and J. C. Tewis, *Biochim. Biophys. Acta*, 271 (1972) 320.
281 O. Vesterberg, *Acta Chem. Scand.*, 27 (1973) 2415.
282 R. C. Williams, Jr. and T. G. Rajagopalan, *J. Biol. Chem.*, 241 (1966) 4951.
283 K. G. Paul and T. Stigbrand, *Acta Chem. Scand.*, 24 (1970) 3607.
284 H. Delincée and B. J. Radola, *Biochim. Biophys. Acta*, 200 (1970) 404.
285 A. Cecil and J. Ogston, *Biochem. J.*, 49 (1951) 105.
286 A. Fernandez-Sörensen and D. M. Carlson, *J. Biol. Chem.*, 246 (1971) 3485.
287 B. Lerch and G. Wolf, *Biochim. Biophys. Acta*, 258 (1972) 206.
288 T. E. Smith, *Arch. Biochem. Biophys.*, 128 (1968) 611.
289 T. E. Smith, *J. Biol. Chem.*, 246 (1971) 4234.
290 S. Harshman and H. R. Six, *Biochemistry*, 8 (1969) 3423.
291 W. A. Scott and T. Abramsky, *J. Biol. Chem.*, 248 (1973) 3535.
292 K. K. Tsuboi and K. Fukunaga, *J. Biol. Chem.*, 246 (1971) 7586.

- 293 R. A. Shipolini, G. L. Callewaert, R. C. Cottrell, S. Doonan, C. A. Vernon and B. E. C. Banks, *Eur. J. Biochem.*, 20 (1971) 459.
- 294 J. I. Salach, P. Turini, R. Seng, J. Hauber and T. P. Singer, *J. Biol. Chem.*, 246 (1971) 331.
- 295 A. T. Tu, R. B. Passey and F. M. Toom, *Arch. Biochem. Biophys.*, 140 (1970) 96.
- 296 S. Kawachi, S. Iwanaga, Y. Samejima and T. Suzuki, *Biochim. Biophys. Acta*, 236 (1971) 142.
- 297 W. L. McLellan, Jr., L. E. McDaniel and J. O. Lampen, *J. Bacteriol.*, 102 (1970) 261.
- 298 K. Hanabusa, H. Kohno and S. Matsuyama, *J. Biochem.*, 72 (1972) 1261.
- 299 P. A. Torjesen and K. Sletten, *Biochim. Biophys. Acta*, 263 (1972) 258.
- 300 L. Summaria, L. Azzadon, P. Bernabe and K. C. Robbins, *J. Biol. Chem.*, 247 (1972) 4691.
- 301 W. J. Kim and T. T. White, *Biochim. Biophys. Acta*, 242 (1971) 441.
- 302 O. A. Lea, *Biochim. Biophys. Acta*, 317 (1973) 351.
- 303 E. Milgrom, P. Allouch, M. Atger and E. E. Baulieu, *J. Biol. Chem.*, 248 (1973) 1106.
- 304 M. Ben-David, D. Rodbard, R. W. Bates, W. E. Bridson and A. Chrambach, *J. Clin. Endocrinol. Metab.*, 36 (1973) 951.
- 305 Y. Inoue, H. Yamamura and U. Nishizuka, *Biochem. Biophys. Res. Commun.*, 50 (1973) 228.
- 306 E. M. Lesk and T. H. Blackburn, *J. Bacteriol.*, 106 (1971) 394.
- 307 H. B. Bosmann, *Biochim. Biophys. Acta*, 293 (1973) 476.
- 308 L. Pechet and J. A. Smith, *Biochim. Biophys. Acta*, 200 (1970) 475.
- 309 M. Pänkäläinen, H. Aro, K. Simons and K. I. Kivirikko, *Biochim. Biophys. Acta*, 221 (1970) 559.
- 310 M. R. Davies and B. M. Dean, *FEBS Lett.*, 18 (1971) 283.
- 311 Y. Sugita, S. Nomura and Y. Yoneyama, *J. Biol. Chem.*, 246 (1971) 6072.
- 312 W. E. Criss, *Biochem. Biophys. Res. Commun.*, 35 (1969) 901.
- 313 T. Mustafa and P. W. Hochachka, *J. Biol. Chem.*, 246 (1971) 3196.
- 314 L. E. Anderson, *Biochim. Biophys. Acta*, 235 (1971) 245.
- 315 B. Sugden and W. Keller, *J. Biol. Chem.*, 248 (1973) 3777.
- 316 E. A. Boeker, A. P. Hays and G. L. Cantoni, *Biochemistry*, 12 (1973) 2379.
- 317 R. M. Waterson, S. J. Clarke, F. Kalousek and W. H. Konigsberg, *J. Biol. Chem.*, 248 (1973) 4181.
- 318 B. Svensson, *FEBS Lett.*, 29 (1973) 167.
- 319 D. P. Baccanari and S. Cha, *J. Biol. Chem.*, 248 (1973) 15.
- 320 E. Neuwelt, D. Stumpf, J. Austin and P. Kohler, *Biochim. Biophys. Acta*, 236 (1971) 333.
- 321 E. Allen and A. B. Roy, *Biochim. Biophys. Acta*, 168 (1968) 243.
- 322 A. B. Roy and A. Jerfy, *Biochim. Biophys. Acta*, 207 (1970) 156.
- 323 S. Marklund, *Acta Chem. Scand.*, 27 (1973) 1458.
- 324 S. Kit, W. Ch. Leung, D. Trkula, D. R. Dubbs and G. Jorgensen, *Intervirology*, 2 (1973/1974) 137.
- 325 S. Kit, W. Ch. Leung, D. Trkula and G. Jorgensen, *Int. J. Cancer*, 13 (1974) 203.
- 326 N. Ui, *Biochim. Biophys. Acta*, 257 (1972) 350.
- 327 H. Tamura and N. Ui, *Biochim. Biophys. Acta*, 214 (1970) 566.
- 328 J. S. Fawcett, M. L. Dedman and C. J. O. R. Morris, *FEBS Lett.*, 3 (1969) 250.
- 329 G. Hennen, G. Maghuin-Rogister and G. Mamoir, *FEBS Lett.*, 9 (1970) 20.
- 330 J. S. Marshall, J. Pensky and S. Williams, *Arch. Biochem. Biophys.*, 156 (1973) 456.
- 331 H. R. Six and S. Harshman, *Biochemistry*, 12 (1973) 2677.
- 332 L. Puutula and R. Gräsbeck, *Biochim. Biophys. Acta*, 263 (1972) 734.
- 333 H. G. Van Eyk, R. J. Vermaat and B. Leijne, *FEBS Lett.*, 3 (1969) 193.
- 334 E. E. Rozacky, T. H. Sawyer, R. A. Barton and R. W. Gracy, *Arch. Biochem. Biophys.*, 146 (1971) 312.
- 335 F. C. Hartman, *Biochemistry*, 10 (1971) 146.
- 336 L. E. Anderson, *Biochim. Biophys. Acta*, 235 (1971) 237.
- 337 A. Tur-Sinai, Y. Birk, A. Gertler and M. Rigbi, *Biochim. Biophys. Acta*, 263 (1972) 666.
- 338 D. R. Joseph and K. H. Muench, *J. Biol. Chem.*, 246 (1971) 7602.
- 339 D. R. Joseph and K. H. Muench, *J. Biol. Chem.*, 246 (1971) 7610.
- 340 N. S. Penneys and K. H. Muench, *Biochemistry*, 13 (1974) 560.
- 341 Z. Kurylo-Borowska and T. Abramsky, *Biochim. Biophys. Acta*, 264 (1972) 1.
- 342 D. J. Barford and J. G. Jones, *Biochem. J.*, 125 (1971) 76p.
- 343 C. C. Contaxis and F. J. Reithel, *Enzymologia*, 41 (1971) 320.
- 344 A. Sutter, R. Ortmann and H. Grisebach, *Biochim. Biophys. Acta*, 258 (1972) 71.
- 345 U. Hakan Stenman, *Biochim. Biophys. Acta*, 342 (1974) 173.