Aminoacids Peptides Proteins

Books

Harper`s Biochemistry Stryer – Biochemistry Lehninger – Biochemistry Moore JT, Langley RH – Biochemistry for Dummies Salway JG – Medical Biochemistry at a Glance Kaneko – Clinical Biochemistry



getting acquinted with biochemical characterisits and metabolic meaning of aminoacids, peptides and proteins

 getting acquinted with biologically important representatives of aminoacids, peptides and proteins

Learning effects

 ability to join together chemical properties of aminoacids, peptides and proteins with their function in living body and the participation in biochemical pathways

 understanding of the meaning of aminoacids, peptides and proteins for appropriate structure and finction of cells

Ehlers Danlos syndrome

Plasma

Liquid which is obtained from full blood collected with anticoagulant after the centrifugation of morphologic elements.

Plasma depending of used anticoagulant can be citrate, heparin or EDTA related. Moreover, it can be platelletrich or platellet-free – it can be distinguished depending on the speed of blood centrifugation.

Plasma

The concentration of total protein in **plasma** 66 – 87 g/dm³

The concentration of total protein in serum $65 - 82 \text{ g/dm}^3$

The role of plasma proteins

- the distrubution of liquids between blood and intercellular space
- transport of hormons, metabolites, metals, drugs etc
- enzymes, regulators
- proteins of immunological system and cloting system
- components of buffers
- hormons and receptors
- components of connective tissue (adhesive)
- nutritional components

Albumins are storage material of body which can be used during starvation and the loss of proteins during the course of different diseases. They contain small content of tryptophan. Albumins maintain constant volume of blood and oncotic pressure. They transport fatty acids, bilirubin, cholesterol and selected ions.

Globulins are characterised by higher molecular weight as albumins. Following fractions are known: α_1 1-4% α_2 4-13% β 7-13% γ<u>8-19%</u>

acute phase proteins – markers of inflammation. They possess inhibitory properties for proteases (the protection of organism from lysosomal enzymes) or are carriers of different substances (haptoglobin). The representatives are: C-reactive **protein** (CRP), α_1 antitrypsin, α_1 acid glikoprotein (transports progesteron), haptoglobin (binds and transports extracellular hemoglobin), ceruloplasmin (cooper binding).

proteins of complement system
transferrin
fibrinogen
hormon binding proteins
enzymes
hormons

immunoglobulins – contain 2 light and 2 heavy chains bound by disulfide bonds Classes: IgG – they are the component of humoral acquired answer (75%) IgA – immunological protection of mucosal membranes, are present in saliva, tears, milk (15%) **IgM** – primary immunological answer (5-10%) IgD – biological function not fully recognised **IgE** – relevant for alergic diseases



Is formated after the centrifugation of blood collected for so called "clot". Blood collected to dry glass clots and is centrifuged to separate serum from clot. Serum differs from plasma the lack of fibrynogen.

Hypoproteinemia

The decrease in protein concentration is the results of the loss of proteins, the inhibition of their synthesis or the dilution of blood. Main reason for hipoproteinemia is the decrease in albumin concentration in blood or more seldom the decrease in immunoglobulin. Critical concentration of total protein is -45g/dm^3 (albumins) below 20 g/dm³) – in this situation edema, transsudation and hypovolemia appear due to the decrease in oncotic pressure and the escape of water from vessels. Often hypoproteinemia is accompaned by dysproteinemia – the alteration in the ratio between concentrations of particular proteins eg. albumin to globulin.

Hyperproteinemia

Is the result of increased production of one or few classes of immunoglobulins. Together with hyperalbuminemia may appear during dehydratation.



Aminoacids

Organic compounds, derivatives of carobxylic acids containing amino group

In accordance to configuration:
1. Protein
2. Rarely present in proteins
3. Non-protein

Consequences: α i L Biosynthesis of proteins



Non-protein aminoacids

β Alanine – component of dipeptides carnosin and anserin as well as pantotein acid which builts coenzyme A

Ornithine, cytruline – intermediates of urea cycle

 γ aminobutyric acid – important for nervous system

Rarely present in proteins

- 5-hydroxylysine
- 4-hydroxyproline
- allysine
- Are the result of posttranslational modifications:
- Adding of **OH** groups to selected prolins and lysins in collagen and gelatin
- Adding of **methyl** groups to selected lysins and histidins in myosin of muscles
- Adding of **carboxyl** groups to selected glutamins in cloth proteins, blood and bone proteins

- Adding of **phosphate** groups to selected serins, treonins and tyrosins

In accordance to the possibility to synthesise branched – isoleucine, 1. Exogenous: leucine, valine aromatic – phenylalanine, tryptophan containing S – methionine basic – arginine, lysine, histidine containing OH group – treonine 2. Endogenous: remaining

Consequences: diet; transamination

In acordance to Karlson With apolar side chain 1. alanine, valine, leucine, isoleucine, proline, phenyloalanine, tryptophan, methionine 2. With polar side chain but without charge glycine, serine, treonine, cysteine, tyrosine, asparagine, glutamine 3. Monoaminodicarboxylic (negatively charged) asparaginic acid, glutaminic acid 4. **Diaminomonocarboxylic** (positively charged) lysine, arginine, histidine

- In accordance to participation in metabolic pathways
- 1. glucogenic:
 - pyruvate pathway (and acetylCoA): threonine, glycine, serine, cystine, cysteine, alanine
 - glutaminic acid pathway (and α -ketoglutarate): arginine, proline, histidine, glutamine
 - succinate pathway: methionine, isoleucine, valine
 - oxalacetate pathway: asparagine, asparaginic acid
- 2. ketogenic:
 - acetoacetyl-CoA pathway: leucine
- 3. mixed: isoleucine, lysine, phenyloalanine, tyrosine

in accordance to the participation in secondary structures of proteins

1. Stabilizing of α helix: alanine, leucine, phenyloalanine, tyrosine, tryptophan, cysteine, methionine, histidine, asparagine, glutamine, valine

2. Destabilizing of α helix: serine, isoleucine, threonine, glutaminic acid, asparaginic acid, lysine, arginine, glycine

3. Interrupting of α helix: proline, hydroxyproline

Reactions of amino group

 with nitrous acid – nitrogen is formatted and the exchange of amino group into hydroxyl group appear (the determination of liberated nitrogen is the basis for quantitative estimation of aminoacids in accordance to the method of van Slyke)

N-acylation with halides or acid anhydrides - N-acyloaminacids are formatted, it can be used for the protection of amino groups during the synthesis of peptides

 N-alkylation, methylation – betains of appropriate aminoacids are formatted, the donor is adenozylmethionin

Ninhydrine reaction – serves for quantitative determination of aminoacids

Reactions of amino group

Reaction of Sanger with fluorodinitrobenzene – serves for labelling and quantitative determination of amino groups in aminoacids and peptides Reaction of Edman with fenyloizotiocyanate – fenylotiokarbamoyl derivatives are formatted, which after treatment with acid undergo cyclisation with the formation of fenylotiohydantoins. Reaction serves for the identification of NH₂-terminal aminoacids in polypeptide chains and the determination of the sequence of aminoacids Reaction with dansyl chloride

Reactions of amino group

deamination –ketoacids are formatedtransamination

Reactions of carboxyl group

Reduction – aminoalkohols are formated
Esterification – serves for the protection of carboxyl groups in the synthesis of peptides
Decarboxylation – amines are formated (biogenic),

the reaction requires pirydoxal phosphate

Biogenic amines

Histidine – histamine – tissue hormon which regulates blood pressure, responsible for alergic reactions Asparaginic acid - β alanine – element of CoA Glutaminic acid – γ aminobutyric acid (GABA) Serine – kolamine – element of conjugated lipids Threonine - **propanolamine** – element of vitamin B12 Cysteine - cysteamine - element of CoA Tyrosine – tyramine – tissue hormon - **dopamine** – substrate for adrenalin synthesis Tryptophan – tryptamine – tissue hormon - serotonine – tissue hormon

Reactions of functional groups

thiol group of cysteine

with ions of heavy metals – mercaptic
 derivatives are formated (inactivation of active
 centers of enzymes)

- Oxidation – cystine is formated

Physico-chemical properties

Acid-base properties depend on environmental pH

In acidic environment aminoacid accepts proton, behaves like cation and in accordance to Bronsted is proton-donor

NH₃⁺ - CHR - COOH
in basic environment aminoacid returns proton, behaves like anion and is proton-acceptor
NH₂ - CHR - COO⁻
in isoelectric point
NH₃⁺ - CHR - COO⁻

Physico-chemical properties

Isoelectric point – pH where aminoacid contains balance between positive and negative charges and is neutrally charged - jon obojnaczy. In this pH value aminoacid is characterised by the lowest solubility and is not moving in electric field.

Consequences – activity of enzymes

- buffers
- isolation

Physical properties of aminoacids

Smell – glutaminic acid – przyprawa

products of reaction between prolin and glucose –
fresh bred

Taste – glutaminic acid – "umami" (5th taste)
Toxicity – pelagra – excess of Leu, insufficiency of Trp

neurotoxicity – excess of Tyr

The determination of aminoacids

- ninhydrin method blue product is formated (except from prolin and hydroxyprolin - yellow)
- cysteine method black sediment of PbS is formated
- Millon's method red product is formated
- Method of Adamkiewicz-Hopkins purple ring on the border of two phases is formated
- xantoprotein method yellow nitro derivatives of aromatic aminoacids are formated
- formol titration in accordance to Sorensen

The separation of aminoacids

electrophoresischromatography

Molecular sieving

Ion exchange chromatography

Affinity chromatography

Laboratory methods for the synthesis of aminoacids

Acidic hydrolysis
Basic hydrolysis
Enzymatic hydrolysis
Microbiologic methods
Synthetic methods
Prebiotic methods

The transportation of aminoacids via membranes

system A – transports majority of neutral aminoacids

- ma cechy wtórnego transportu aktywnego
- zależy od wewnątrzkomórkowego stężenia aminokwasów
- podlega regulacji hormonalnej
- układ ASC transportuje alaninę, serynę i cysteinę
- układ Gly- transportuje glicynę
- układ N transportuje histydynę, glutaminę i asparaginę
- układ L transportuje leucynę, izoleucynę, walinę i fenyloalaninę
 - działa na zasadzie dyfuzji ułatwionej



Definition

Chain molecules containing from 2 up to 100 aminoacids, which are bound by amide bond named peptide bond. These molecules can cross dialysis membranes (molecular weight up to 10 000 D) and are not susceptible to denaturation due to the lack of secondary structure.

from 2 up to 10 aminoacids – OLIGOPEPTIDS
more than 10 aminoacids – POLYPEPTIDS

homeomeric – contain ONLY aminoacids
 heteromeric (peptolides) – contain additional structural elements

homodetic – contain ONLY peptide bonds
heterodetic – other bonds such as ester, disulfide, thioester may occur

The meaning of peptides

biologically active hormons
antybiotics
toxins
in food industry

Peptide bond

Substituted amide bond that binds amino group of one aminoacid with carboxyl group of next aminoacid (water is liberated). It has the character of a covalent bond and shows the following features:

semiunsaturated – single bond of C-N posseses in 40% character of double bond while double bond between C=O posseses in similar range the character of single bond

planar – atoms of C and N, as well as C adjacent to them are in the same plane

cross substituted – C atoms are always in trans position to each other and to peptide bond

polar

The synthesis of polypeptide chain

Aims:

- 1. The confirmation of proposed primary structures by chemical synthesis
- 2. The examination of the relationship between structure and biologicalactivity via synthetic analogs
- 3. The chemical changes of biologically active peptides in order to modify pharmacological effects
- 4. Economic requirements
- 5. The synthesis of model peptides

The synthesis of dipeptide

The stages for the determination of aminoacid sequence

- 1. The identification of terminal NH₂ and COOH aminoacid
- 2. The hydrolysis of polypeptide chain with trypsin
- 3. Electrophoretic or chromatographic separation of obtained fragments as well as the indetification of terminal NH_2 i COOH aminoacids
- 4. Gradual degradation of chain in accordance to Edman
- 5. The hydrolysis of chain with the use of different peptidase

Method of Sanger with 2,4-dinitrofluorobenzen

After hydrolysis newly formated DNP-aminoacids are extracted and determined chromatographically

Method of Edmana with phenyl-isotiocyanate

'labelled" aminoacid and the chain shorter for one aminoacid are obtained

Biologically important oligopeptides

- carnosin and anserin
- aspartam (more sweet as saccharose)
- glutathion
- enkephalins
- angiotensin II
- bradykinin
- vasopresin, oxytocin

Biologically important polypeptids

Glucagon
Insulin
Endorfins
ACTH
Parathormon
Calcitonin

Glycine (1820r.)

characteristic for connective tissue
is able to neutralise toxic substances via connection with their carboxyl groups for example: benzoic acid + glycine → hippuric acid
participates in the synthesis of purine bases
does not contain assymetric carbon atom
binds to bile acids



Alanine (1888 r.)

is present nearly in each protein
 after deamination pyruvate is formated – the substrate for gluconeogenesis



Serine (1865)

after decarboxylation colamine is formated – the compound of conjugated lipids
 easily undergoes esterification with phosphate acid





easily undergoes esterification with phosphate acid



Phenyloalanine (1879) and Tyrosine (1846) precursor of thyroid gland hormons, adrenaline and melanine metabolic blocks in biochemical pathways: Alkaptonuria – the lack of brak homogentisin oxygenase Fenyloketonuria – the lack of phenyloalanin hydrolase

- Albinizm – the lack of o-difenol oxygenase



Tryptophan (1901)

- precursor in the synthesis of vitamin B3 nicotinic acid
- after decarboxylation formates serotonine
- thanks to the presence of aromatic rings may absorb ultraviolet light at 280 nm – it provides with the possibility of spectrophotometric determinations
- formates yellow nitro derivatives with nitric acid (quantitative determinations)
- metabolic blocks in pathways:
- Disease of Hartnup –
 the lack of tryptophan
 oxygenase





after decarboxylation histamine is formated



Cysteine (1884)

• the component of glutathione

- derivative of cysteine taurine creates complexes with bile acids
- derivative cystine (1810)
- oxidation leads to the formation of cystein acid
- heating in basic solutions causes the liberation of sulfur, ammonia and pyruvate



Methionine (1922)

 donor of single carbon fragments do reactions of synthesis (as Sadenosyl-metionine)
 ovstoing reaction is pagative

cysteine reaction is negative

$CH_3 - S - CH_2 - CH_2 - COO^-$

Valine (1901), Leucine (1819), Isoleucine (1904)

exogenous – due to branched chain



Lysine (1889), Arginine (1895)



Proline (1901)

 characteristic compound of collagen – protein of connective tissue

derivative of proline is hydroxyproline



Asparaginic acid (1868) and Asparagine

ability to bind ammonia



Glutaminic acid (1866) and Glutamine

ability to transport ammonia ions

