Федеральное государственное автономное образовательное учреждение высшего образования

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Методические материалы по дисциплине:

Основы профессионального перевода

основная профессиональная образовательная программа высшего профессионального образования – программа специалитета

30.05.01 Медицинская биохимия

## Тестовые задания:

Оценочное средство	Эталон ответа	Уровень
7		применения*
Лексико-грамматический тест по дисциплине		$\Pi A$
«Основы профессионального перевода»:		
At the molecular level medicines interact with	A	
in the body.		
A. proteins		
B. enzymes		
C. receptors		
X 4:1 C4:: 4 4: 11		
You can think of this interaction like	C	
·		
A 1 1		
A. a docking process		
B. drug-protein binding		
C. a key fitting a lock		
The word CHONPS stands for	C	
The word CHONPS stands for		
A. the list of common elements in our body		
B. the backbone of our DNA		
C. the list of common elements of drugs organic in nature		
III liature		
Salt bridges between two charged groups contribute	A	
to	А	
A. a drug potency		
B. a drug specificity		
C. a drug stability		
C. a drug stability		
To provide specificity for different protein targets	В	
drug designers use	D .	
A. hydrophobic groups		
B. hydrogen bonds		
C. amino groups		
e. umme groups		
When there is a good fit between the shape of the	A	
drug molecule and the shape of the target protein the		
main role in drug-protein binding play		
·		
A. hydrophobic groups		
B. hydrophilic groups		
C. carboxylic groups		
Sulphanilamide is a metabolite of prontosil	A	
1 r		
A. True		

B. False		
D. Faisc		
IPTD used to treat typhoid fever caused deafness	В	
A. True		
B. False		
Carbutamide has a hypoglycaemic effect but a short	A	
half-life	A	
A. True B. False		
Tolbutamide was much more toxic than carbutamide	B	
A. True		
B. False		
It was sulfonylurea – the central section of a	A	
chlorpropamine molecule – that gave the name to a	A	
general class of antidiabetic agents		
A. True		
B. False		
Gliclazide is a next generation sylfanylurea drug	A	
A. True		
B. False		
adding an aromatic ring to the nitrogen atom	D	
<ul><li>A. increase the elimination half-life</li><li>B. reduce toxicity</li></ul>		
C. stimulate the pancrease to release more		
insulin		
D. improve solubility replacing right hand ring with a short chain of carbon	C	
atoms		
A		
<ul><li>A. increase the elimination half-life</li><li>B. reduce toxicity</li></ul>		
C. stimulate the pancrease to release more		
insulin		
D. improve solubility		
replacing the amino group on the benzene ring with a	В	
methyl group		
A. increase the elimination half-life		
B. reduce toxicity		
C. stimulate the pancrease to release more		
insulin		

D. improve solubility	
replacing the methyl group with a chlorine atom	A
A. increase the elimination half-life	
B. reduce toxicity	
C. stimulate the pancrease to release more	
insulin	
D. improve solubility	
D. Improve solubility	
HCM CoA	B
HGM-CoA	B
A. product	
B. substrate	
C. inhibitor	
D. enzyme	
HGM-CoA reductase	D
A. product	
B. substrate	
C. inhibitor	
D. enzyme Mevalonic acid C6H12O4	A .
Mevalonic acid CoH12O4	A
A. product	
B. substrate	
C. inhibitor	
D. enzyme	
mevastatin	C
A. product	
B. substrate	
C. inhibitor	
D. enzyme	
	A
How many enzymes are involved in the production	A
of cholesterol within body cells?	
A. 30	
B. 13	
C. 10	
What is an important building block in the	В
generation of cholesterol?	
A. HGM-CoA	
B. mevalonic acid	
C. CoA	
C. CUA	
Wil .	l n
What microorganisms can produce statins?	B
A. which constantly battling each other	
for resources and spaces	
	<del>-</del>

D1:-1 124	T T	
B. which don't require HGM-CoA reductase for survival		
C. fungal microorganisms		
How many microorganisms were screened to find	A	
mevastatin?	A	
mevastatiii:		
A. 6,000		
B. 600		
C. 1,600		
C. 1,000		
The first semisynthetic statin was made from	В	
lovastatin by adding		
lovastatin by adding		
A. an oxygen atom		
B. a methyl group		
C. a six-membered ring		
C. a six-inclinered ring		
When the statin is converted to the ring opened form	C	
in the body it resembles		
in the body it resembles		
A. HGM-CoA		
B. Mevolonate		
C. Both a) and b)		
Statins can block HGM-CoA reductase because	A	
Statilis call block Troivi-Coa reductase because	A	
A. they have spatial and chemical match		
needed to fit within the enzyme		
pocket		
B. they are complementary in structure		
with HGM-CoA		
C. because they have hydroxyl and		
carboxyl groups in their structure		
carboxyr groups in their structure		
benzene ring A	В	
ochzene imgri		
A. no functional groups		
B. a hydroxyle group -OH attached		
C. an ether oxygen		
D. an amino group -NH2 with basic N and a		
methyl group attached		
metry group attached		
six-membered ring B	A	
A. no functional groups		
B. a hydroxyle group -OH attached		
C. an ether oxygen		
D. an amino group -NH2 with basic N and a		
methyl group attached		
mondy by orb annoned		
ring D	D	
<i>S</i> -		
A. no functional groups		
	1	

B. a hydroxyle group -OH attached		
C. an ether oxygen		
D. an amino group -NH2 with basic N and a		
methyl group attached		
7 8 m		
five-membered ring E	C	
A C		
A. no functional groups		
B. a hydroxyle group -OH attached		
C. an ether oxygen		
D. an amino group -NH2 with basic N and a		
methyl group attached		
7 8 - 1		
The structure of morphine comprises five rings that	A	
are joined into a specific scaffold		
are joined into a specific scarrold		
A. True		
B. False		
D. Taisc		
The morphine molecule has three hydroxyl groups in	В	
its structure		
its structure		
A. True		
B. False		
The special feature of the molecule is that it is	В	
	Б	
flexible		
A. True		
B. False		
The key functional groups that give morphine its	A	
biological activity are held in place in specific		
position relative to each other		
position relative to each other		
A. True		
B. False		
D. Paisc		
The human body produces its own pain relieving	A	
1		
chemicals, known as enkephalins		
A. True		
B. False		
D. Faist		
A pyridine ring is a five-membered ring with one	В	
_ = -		
carbon atom replaced with a nitrogen atom		
A. True		
B. False		
In pyrrolidine ring nitrogen atom has a methyl group	A	
	**	
attached		

А Т		
A. True		
B. False		
Y 4 11 1		
In the bloodstream the nitrogen atom of the	A	
pyrrolidine ring of nicotine has a charge of plus 1		
A. True		
B. False		
A hashed bond between the two rings of a nicotine	В	
molecule indicates that the pyridine ring is oriented		
towards the viewer		
A. True		
B. False		
Chiral molecules can be superimposed	В	
A. True		
B. False		
D. Tuise		
Morphine is a good example where only one mirror	A	
_ = = = = = = = = = = = = = = = = = = =	А	
image form gives pain relief		
A T		
A. True		
B. False		
The binding site for the drug in the receptor must	A	
have the complimentary shape of the active drug		
A. True		
B. False		
In drug design it is extremely useful to know which	A	
conformation is biologically relevant		
A. True		
B. False		
Cytisine interacts with the nicotine receptor	A	
A. True		
B. False		
Cytisine does not contain a basic nitrogen	В	
A. True		
B. False		
Cytisine molecule is not flat, but rigit	A	
Symbolic molecule is not mat, but light	11	
A. True		
B. False		
D. Paisc		

The left hand ring of cytisine has a carboxyl group	В
A. True	
B. False	
ARR17779 is in the phase I clinical trials	В
-	
A. True	
B. False	
The structure of ARR is made up of a five-membered	A
ring with an oxygen and nitrogen	
A. True	
B. False	
D. Tube	
Common to all three molecules – nicotine, cytisine	A
and ARR – is the presence of a basic nitrogen atom	
A. True	
B. False	
The pyrimidine nitrogen and carbonyl groups make	A
hydrogen bonds with the nicotinic receptor	
A. True	
B. False	
the bearing should together by a sulphur atom	B
– the basic amino group	
A. Phenbenzamine antihistamine	
B. Chlorpromazine antipsychotic	
C. Noradrenaline natural neurotransmitter	
<ul><li>D. Imipramine tricyclic antidepressant</li><li>E. Diphenhydramine antihistamine</li></ul>	
F. Fluoxetine Prozac selective serotonin	
reuptake inhibitor SSRI	
two benzene rings – a three carbon chain – the basic	D
amino group	
A. Phenbenzamine antihistamine	
B. Chlorpromazine antipsychotic	
C. Noradrenaline natural neurotransmitter	
<ul><li>D. Imipramine tricyclic antidepressant</li><li>E. Diphenhydramine antihistamine</li></ul>	
F. Fluoxetine Prozac selective serotonin	
reuptake inhibitor SSRI	
two benzene rings – an oxygen – two carbon chain -	
the basic amino group	<u> </u>

<ul> <li>A. Phenbenzamine antihistamine</li> <li>B. Chlorpromazine antipsychotic</li> <li>C. Noradrenaline natural neurotransmitter</li> <li>D. Imipramine tricyclic antidepressant</li> <li>E. Diphenhydramine antihistamine</li> <li>F. Fluoxetine Prozac selective serotonin reuptake inhibitor SSRI</li> </ul>		
two benzene rings bonded together by a two atom chain – the basic amino group with two methyl groups attached	A	
<ul> <li>A. Phenbenzamine antihistamine</li> <li>B. Chlorpromazine antipsychotic</li> <li>C. Noradrenaline natural neurotransmitter</li> <li>D. Imipramine tricyclic antidepressant</li> <li>E. Diphenhydramine antihistamine</li> <li>F. Fluoxetine Prozac selective serotonin reuptake inhibitor SSRI</li> </ul>		
two benzene rings – an oxygen – three carbon chain - the basic amino group attached to a right hand benzene ring	F	
<ul> <li>A. Phenbenzamine antihistamine</li> <li>B. Chlorpromazine antipsychotic</li> <li>C. Noradrenaline natural neurotransmitter</li> <li>D. Imipramine tricyclic antidepressant</li> <li>E. Diphenhydramine antihistamine</li> <li>F. Fluoxetine Prozac selective serotonin reuptake inhibitor SSRI</li> </ul>		
a benzene ring with two hydroxyl groups – a two carbon chain with a hydroxyl group – the basic amino group	С	
<ul> <li>A. Phenbenzamine antihistamine</li> <li>B. Chlorpromazine antipsychotic</li> <li>C. Noradrenaline natural neurotransmitter</li> <li>D. Imipramine tricyclic antidepressant</li> <li>E. Diphenhydramine antihistamine</li> <li>F. Fluoxetine Prozac selective serotonin reuptake inhibitor SSRI</li> </ul>		
Certain antihistamines can also block serotonine and nonadrenaline reuptake	A	
A. True B. False		
The reuptake blocker nisoxetine was used as a starting point to develop the antihistamine	В	

diphenhydramine	
A T	
A. True B. False	
Nisoxetine was very useful in drug development as it	A
was more selective at inhibitining noradrenaline reuptake with lesser effect on serotonin reuptake	
A. True B. False	
D. raise	
Nisoxetine was marketed in 1987	В
A. True	
B. False	
	2
Prozac is very specific for serotonin reuptake with fewer side effects than antihistamines	B
A. True B. False	
D. raise	
By combining the shapes of different antidepressants	A
molecules we get a picture of the pocket of the target protein	
protein	
A. True	
B. False	
The picture of the pocket of the target protein could	A
be used for the design of new drugs	
A. True	
B. False	

ДОКУМЕНТ ПОДПИСАН ЭЛЕКТРОННОЙ ПОДПИСЬЮ

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