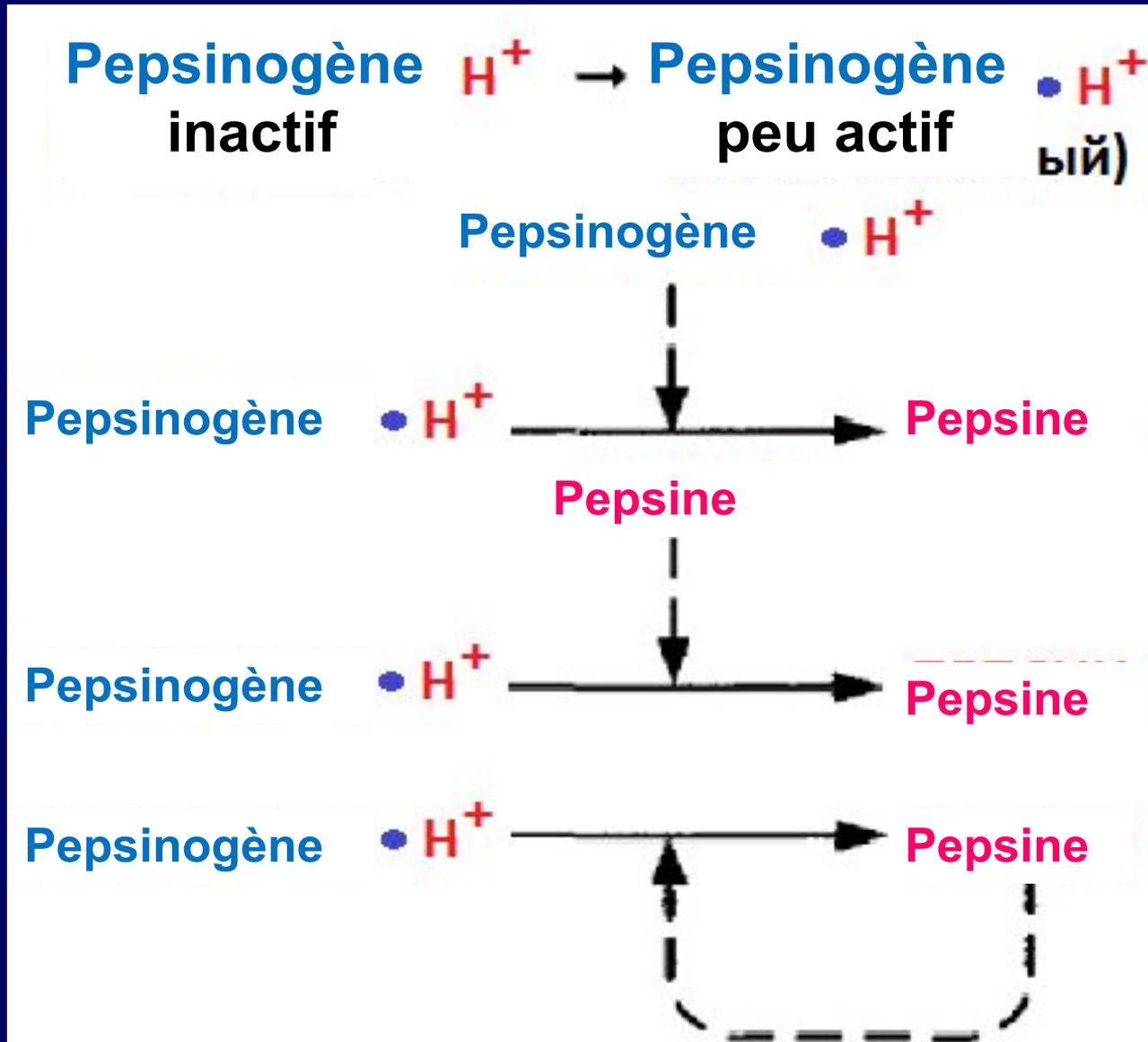
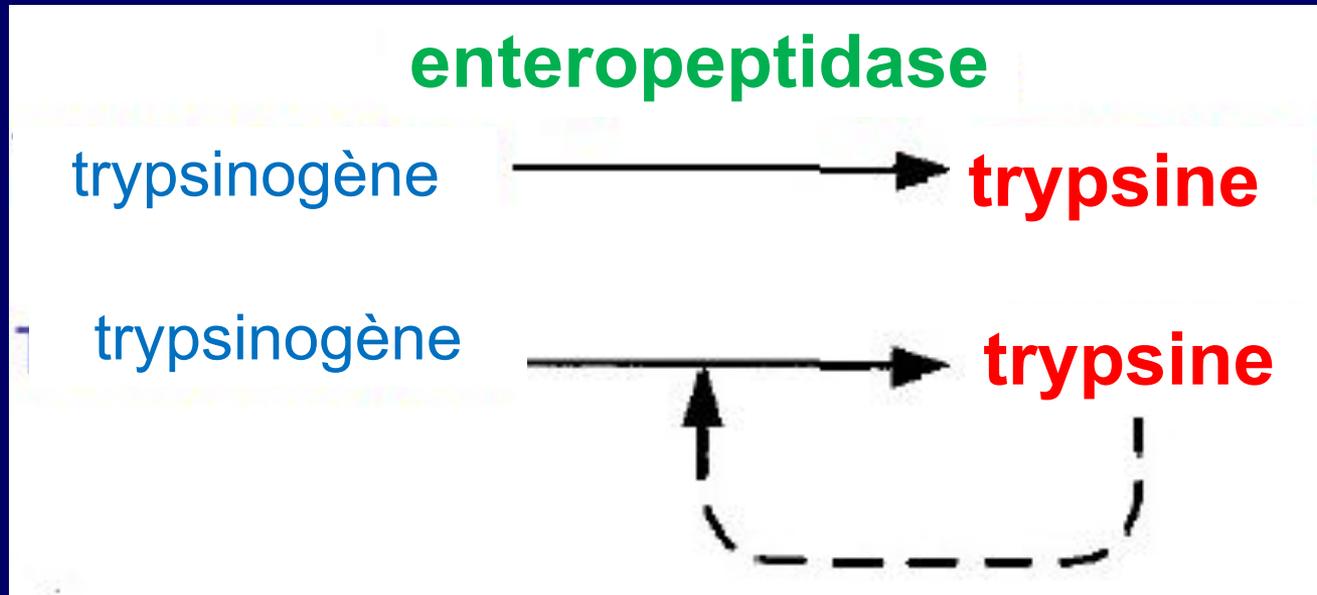


MÉTABOLISME DES PROTÉINES

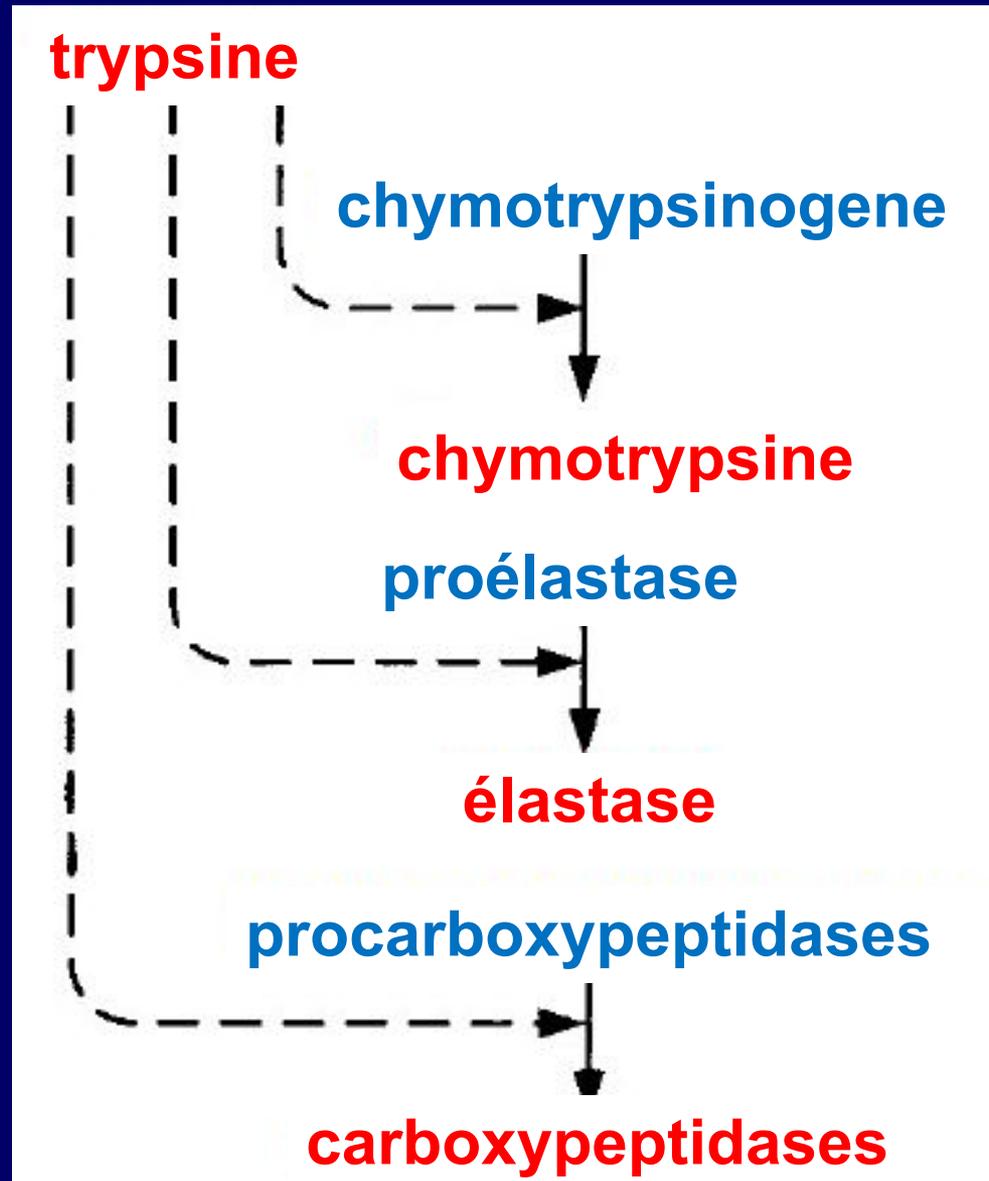
Activation de la pepsine



Activation de la trypsine



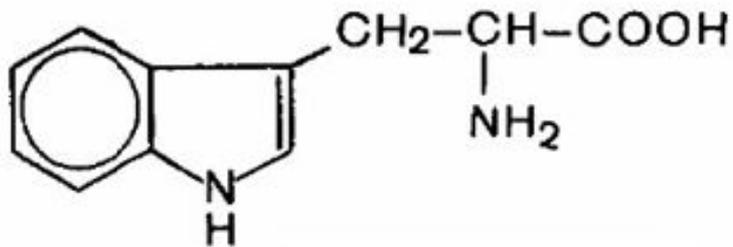
Activation des proteases de l'intestin



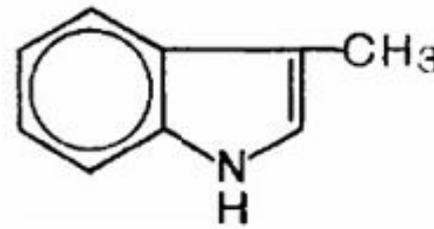
Sélectivité de l'action des peptidases

Enzyme	Liaisons peptidiques rompues
Pepsine	Phénylalanine, tyrosine, acide glutamique
Trypsine	Lysine, arginine
Chymotrypsine	Tryptophane, phénylalanine, tyrosine
Élastase	Glycine, alanine, sérine
Carboxypeptidases A	Tryptophane, phénylalanine, tyrosine
Carboxypeptidases B	Lysine, arginine

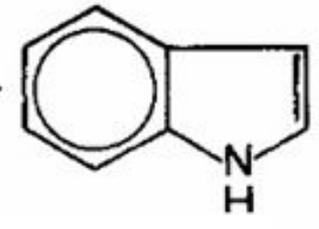
La putréfaction des protéines dans l'intestin



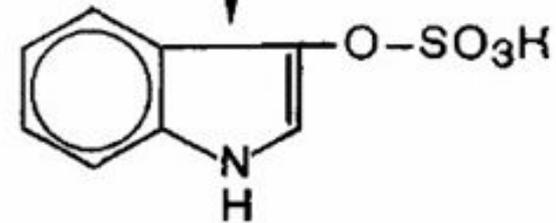
tryptophane



scatole



indole

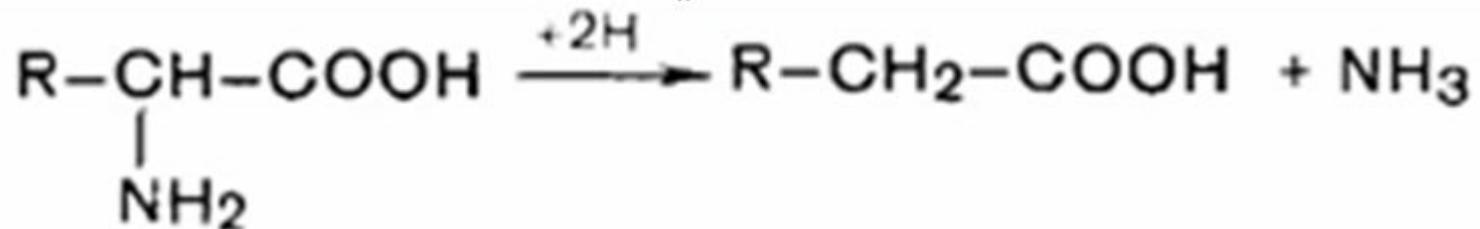


indican animal

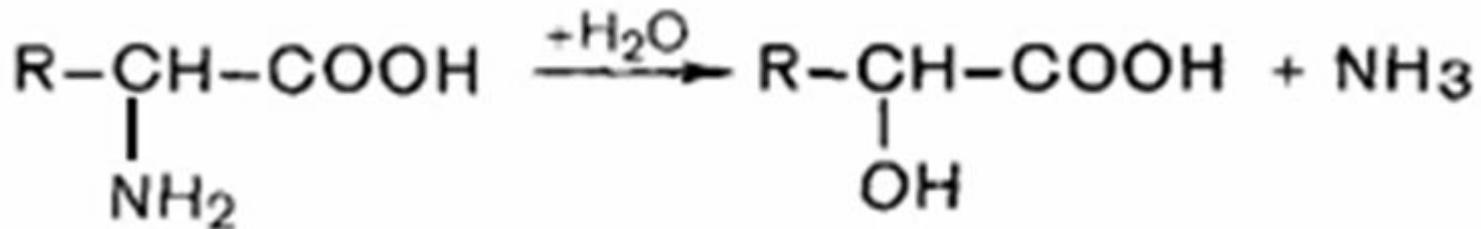
Les réactions des acides aminés

1. Désamination

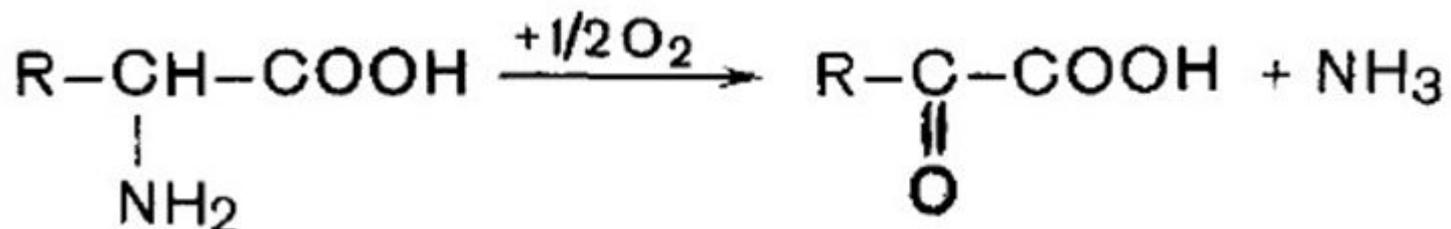
- réductrice :



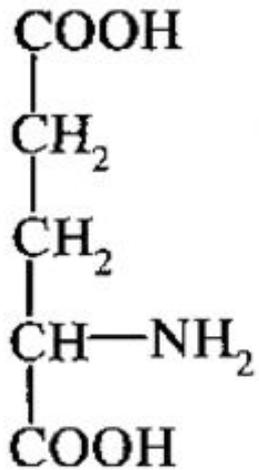
- hydrolytique:



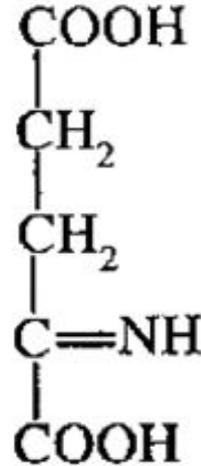
- oxydative:



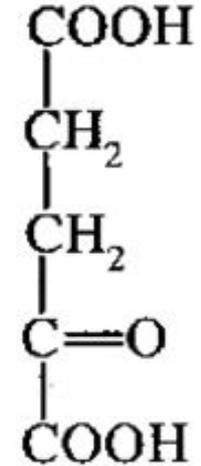
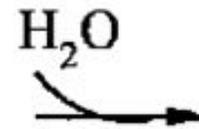
La désamination de l'acide glutamique



Acide glutamique



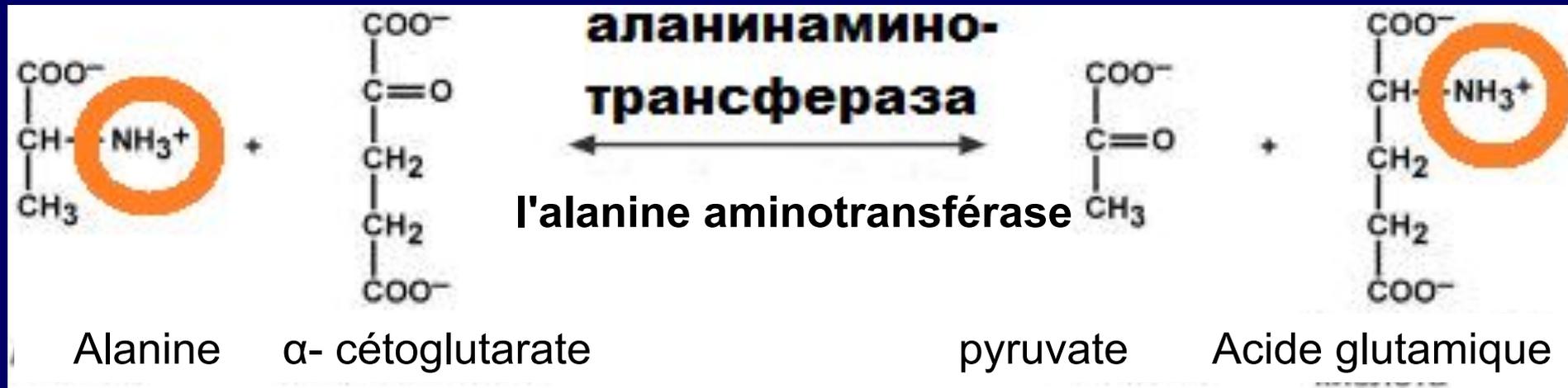
acide α -aminoglutarique



acide α -cétoglutarique



2. Transamination



Transaminases

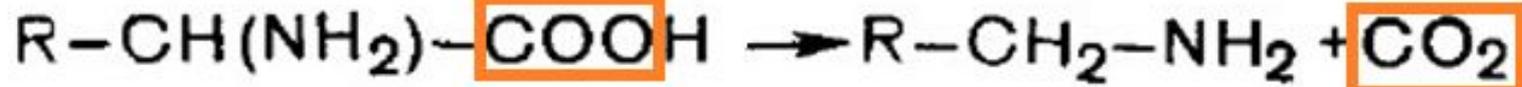
-Aspartate aminotransférase :

aspartate + α -cétoglutarate \rightarrow \rightarrow
oxaloacétate + glutamate

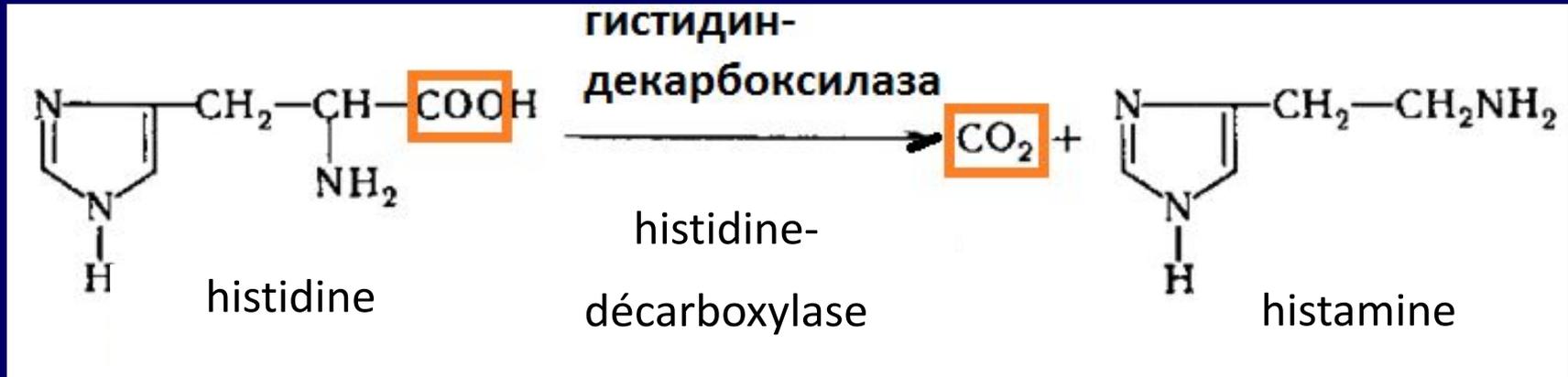
- Alanine aminotransférase:

alanine + α -cétoglutarate \rightarrow
 \rightarrow pyruvate + glutamate

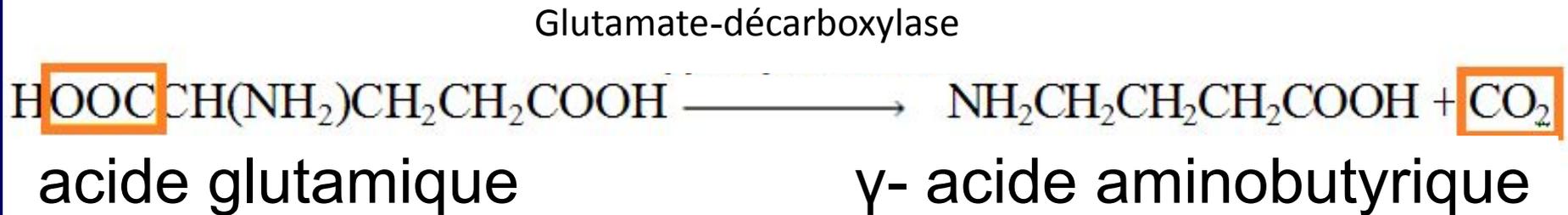
3. Décarboxylation



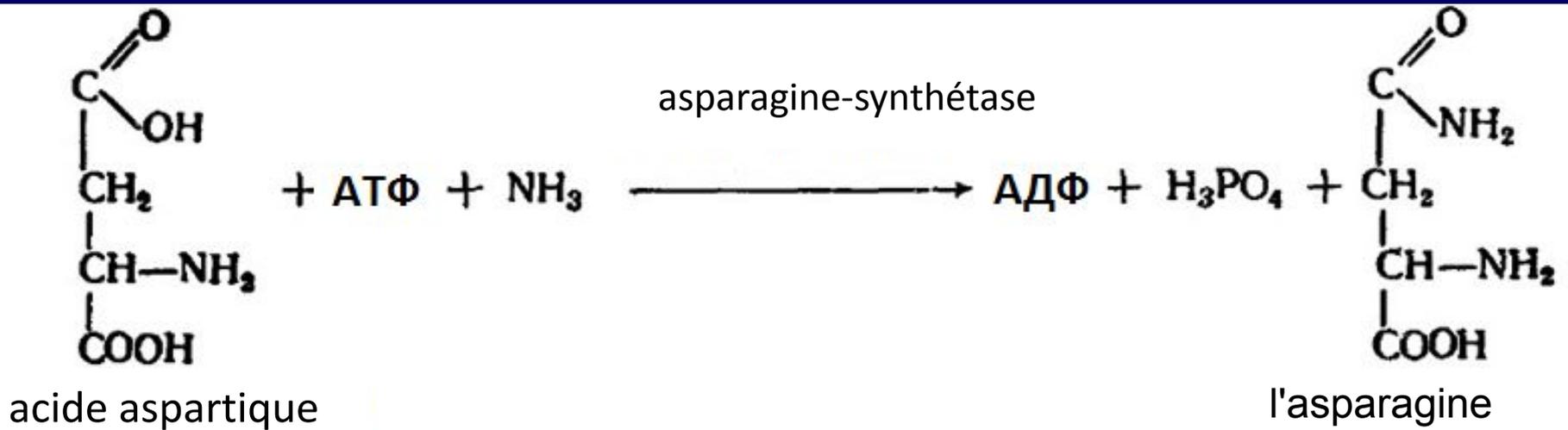
Décarboxylation de l'histidine :



Décarboxylation de l'acide glutamique :



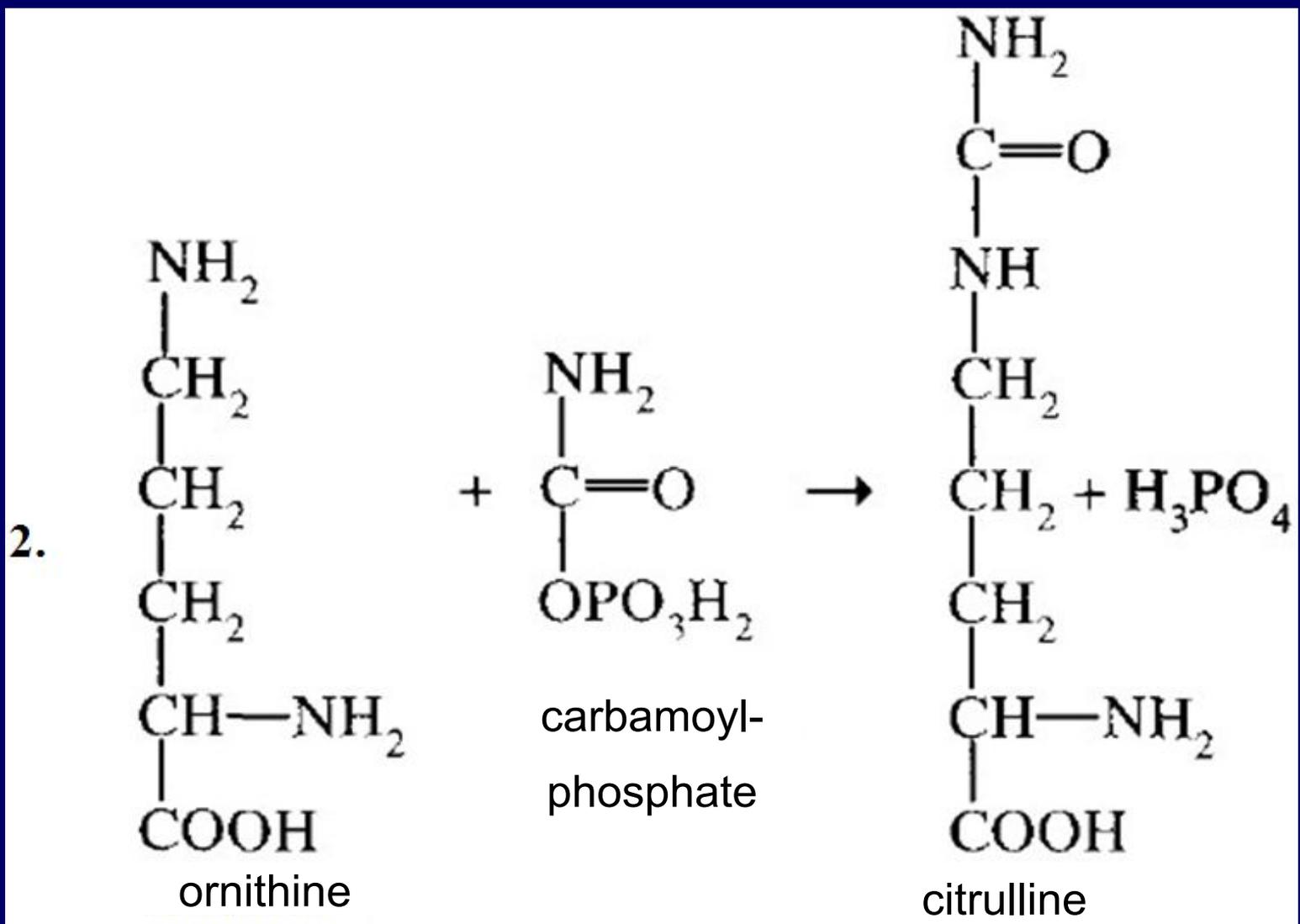
La fixation primaire de l'ammoniac



Le cycle ornithine

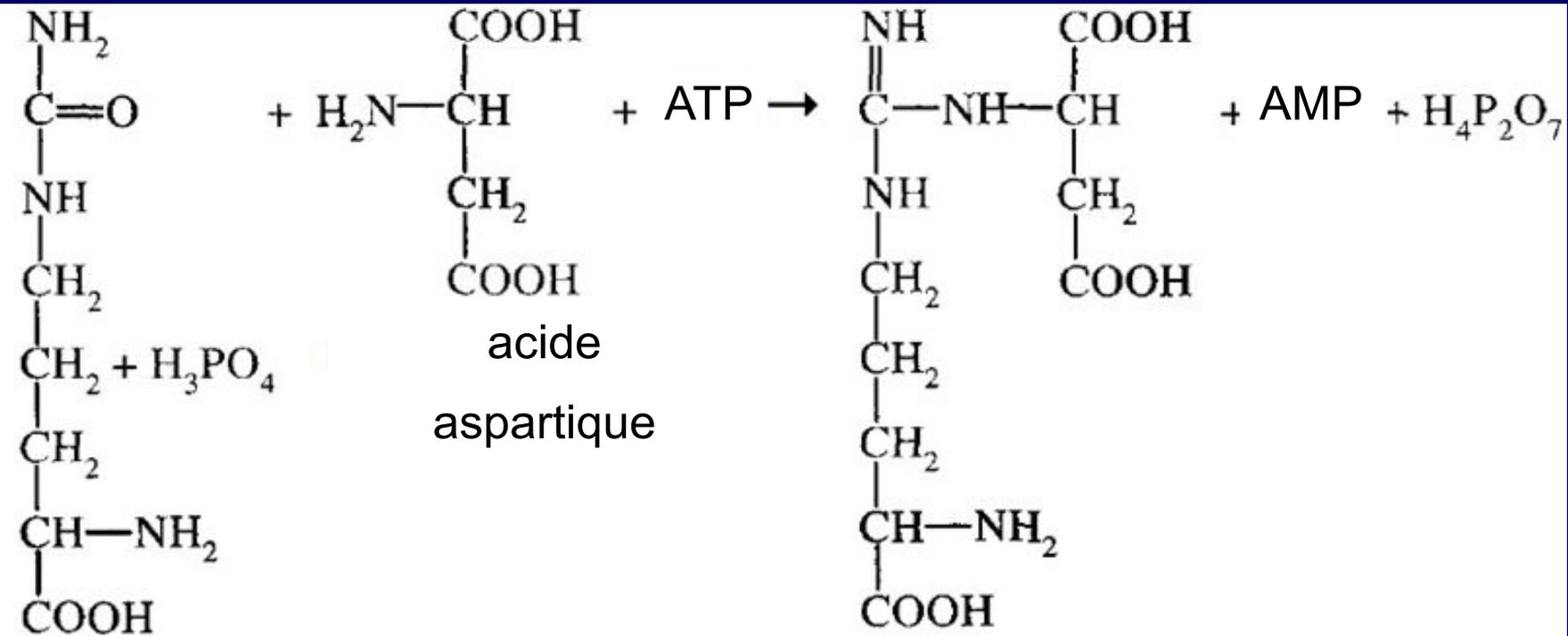


**E – carbamoylphosphatesynthétase
(ammoniaquedépendente)**



E – ornithinecarbamoyltransferase

3.

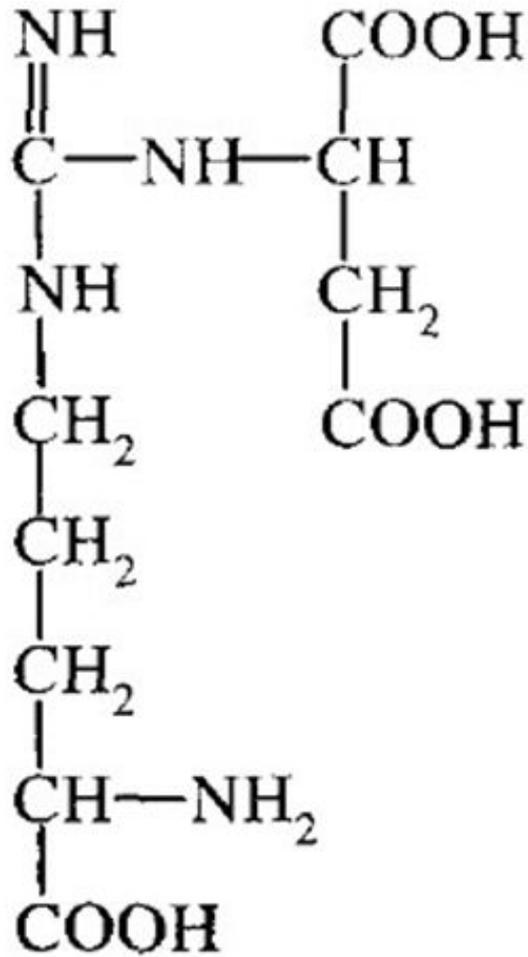


citrulline

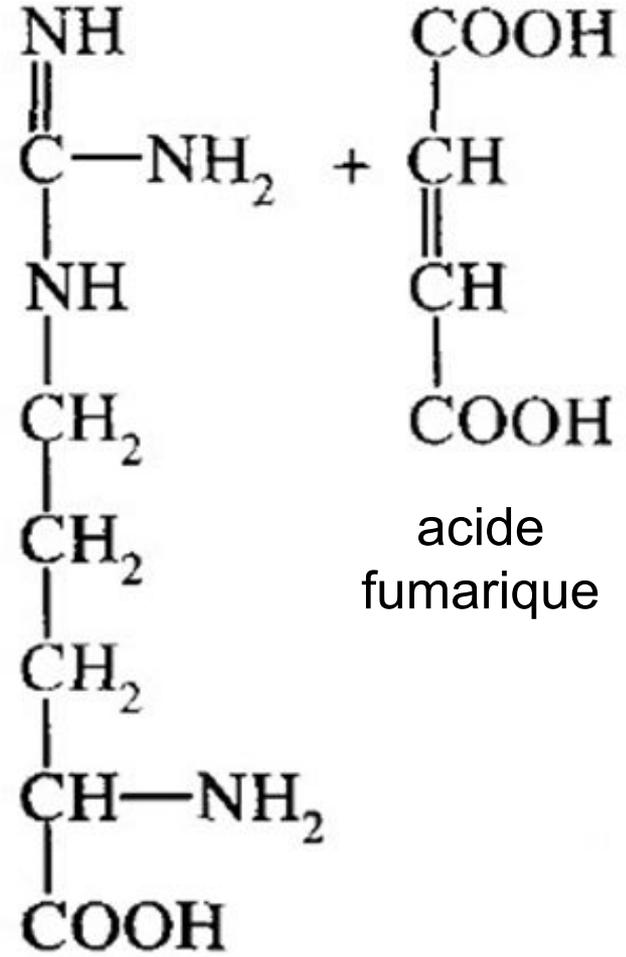
argininesuccinate

E - argininesuccinatesynthétase

4.



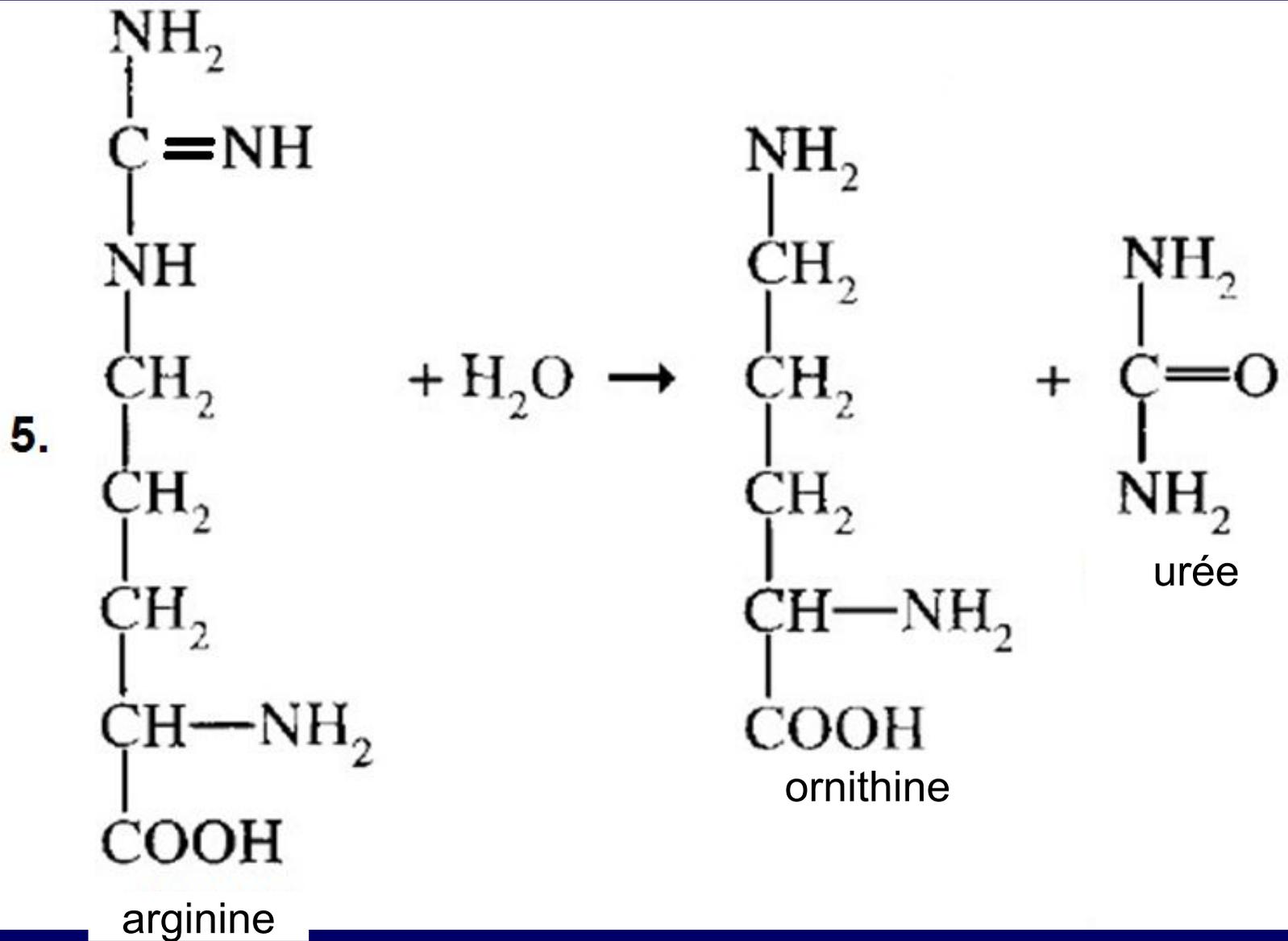
argininesuccinate



arginine

acide
fumarique

E – argininesuccinateliase



E – arginase

Foie



urée



sang



reins



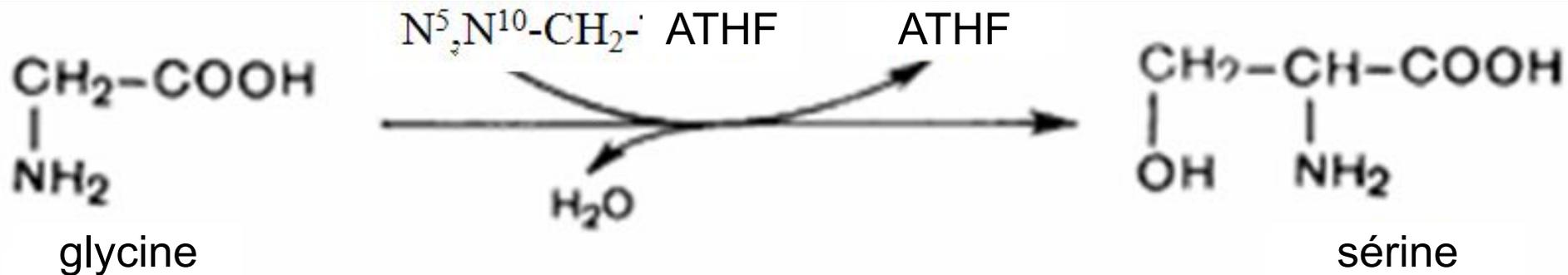
urine
(urée)

La pellagre infantile



métabolisme des acides aminés

- glycine, sérine, thréonine



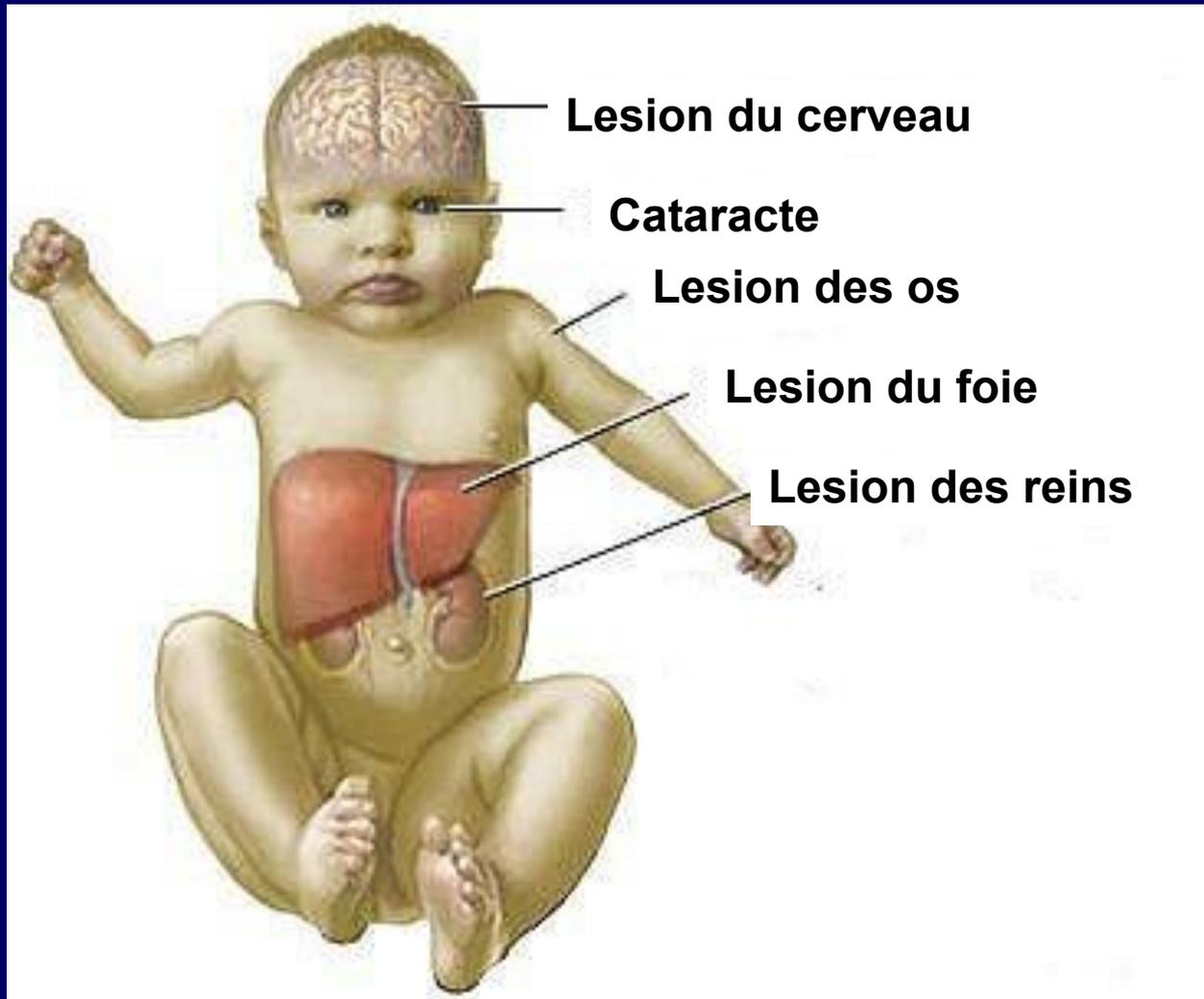
- des acides aminés sulfurés

Méthionine + ATP → S-adénosylméthionine + PP + P

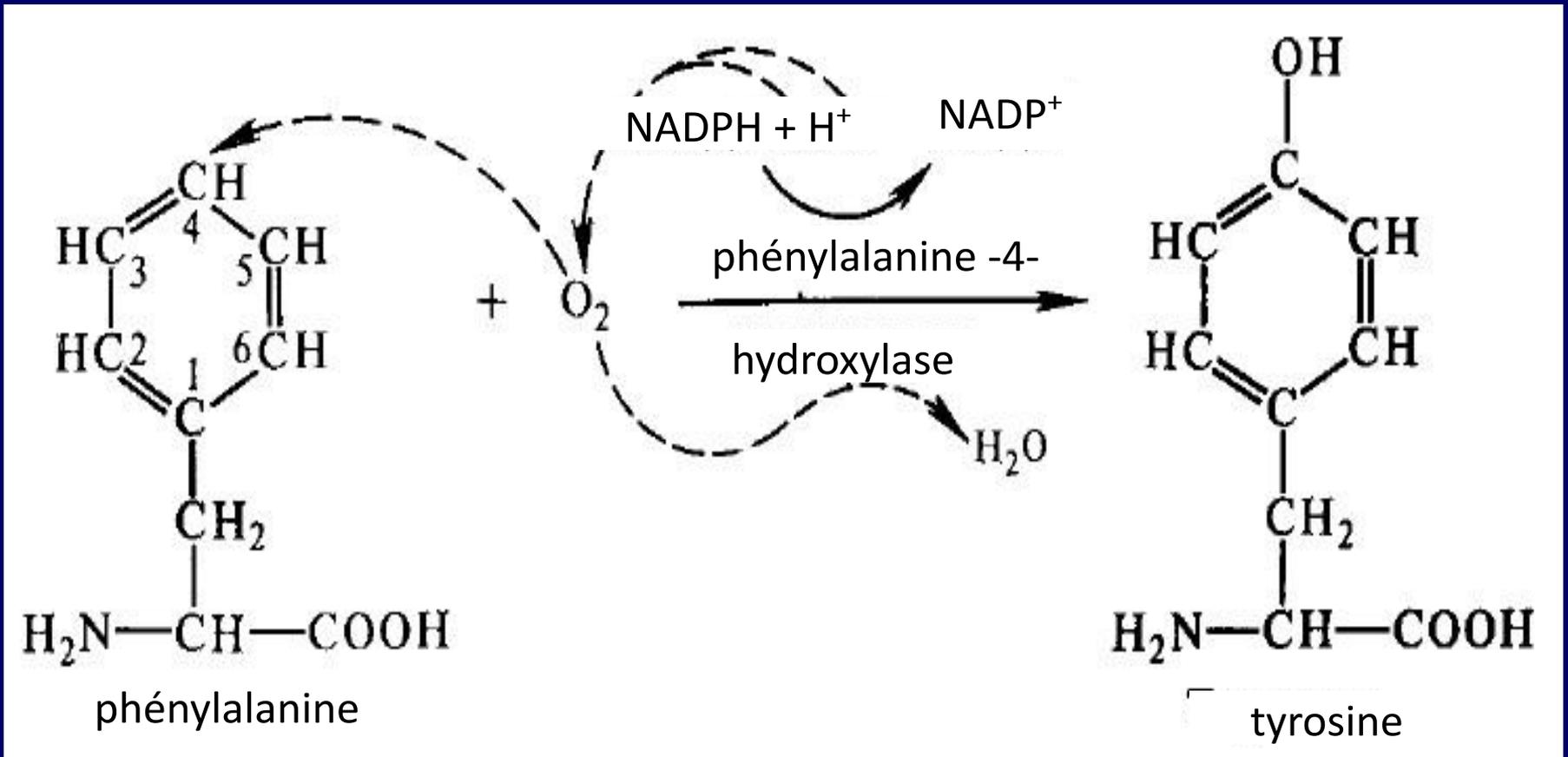
- des acides aminés à la chaîne ramifiée

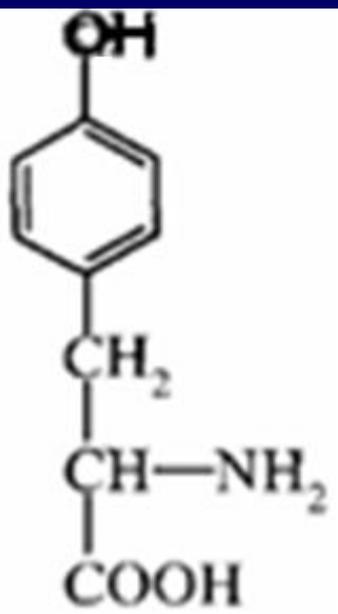
Leu, Ile, Val → α-céto-acides → l'acyl-CoA dérivés

la maladie du sirop d'érable



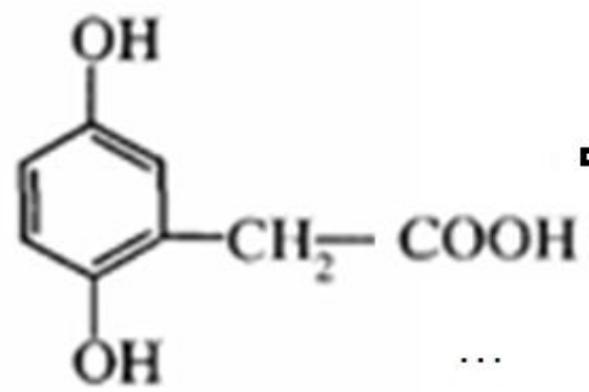
- La phénylalanine et la tyrosine





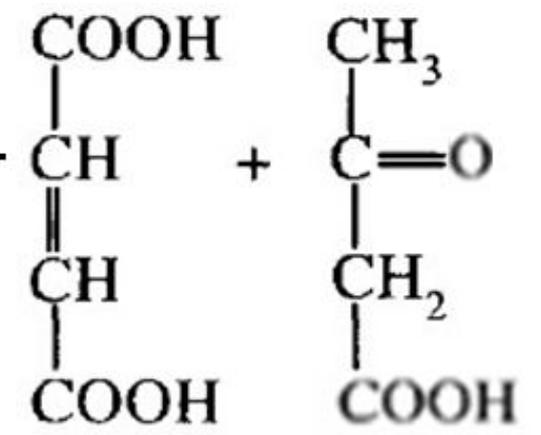
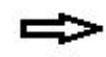
tyrosine

...



acide homogentisique

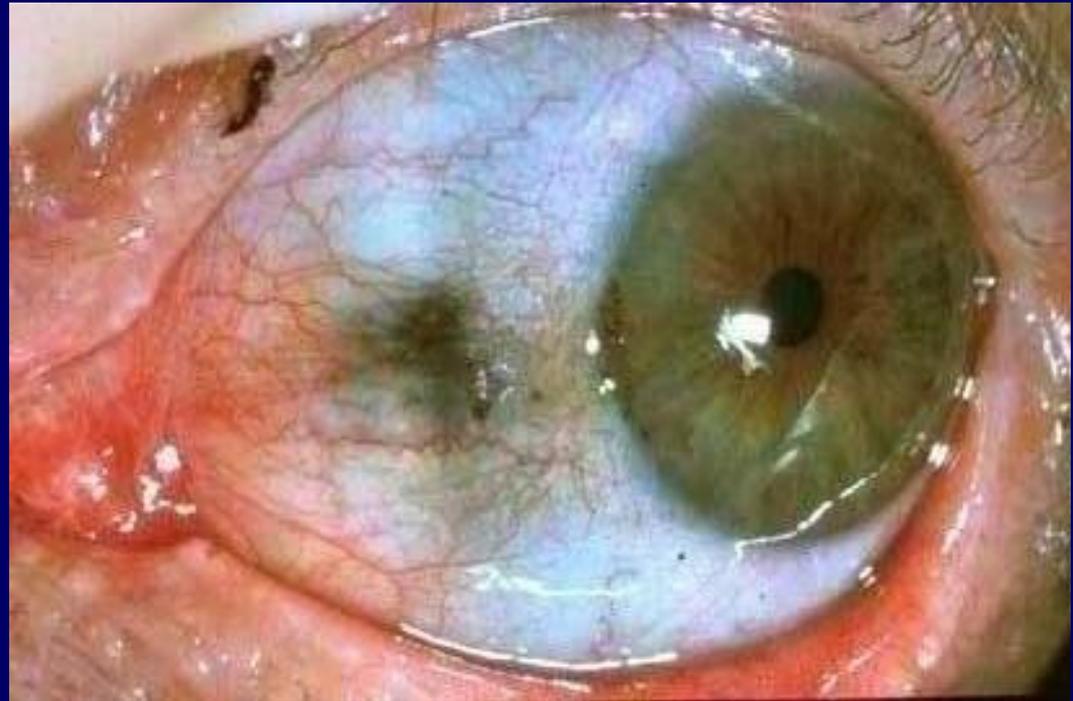
...

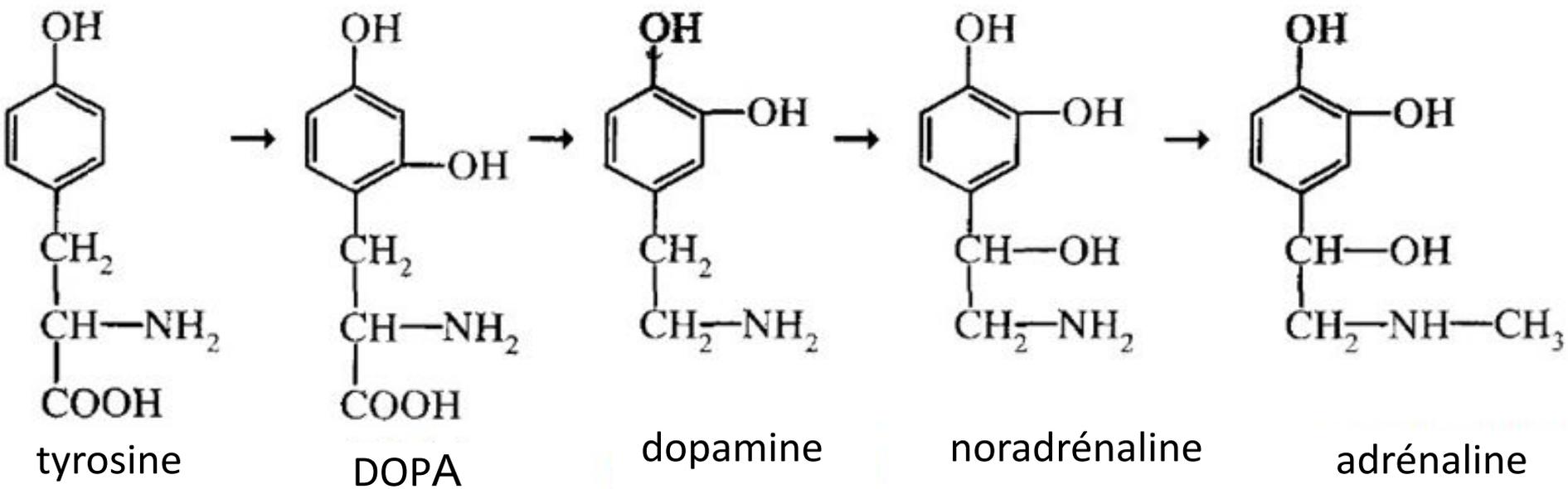


fumarate

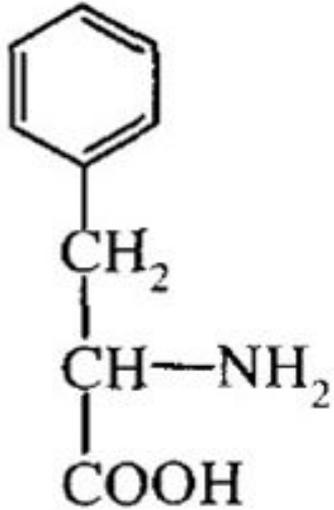
acéto-acétate

L'alcaptonurie

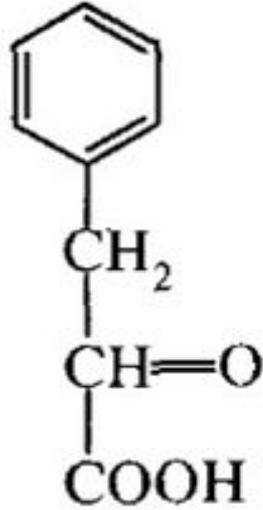




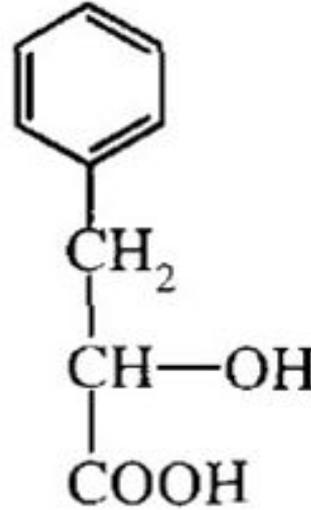
La phénylcétonurie



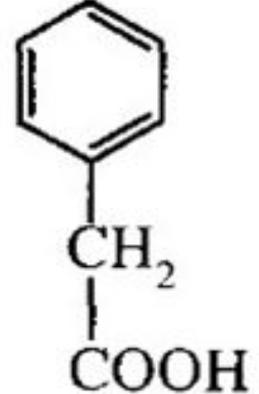
phénylalanine



phénylpyruvate

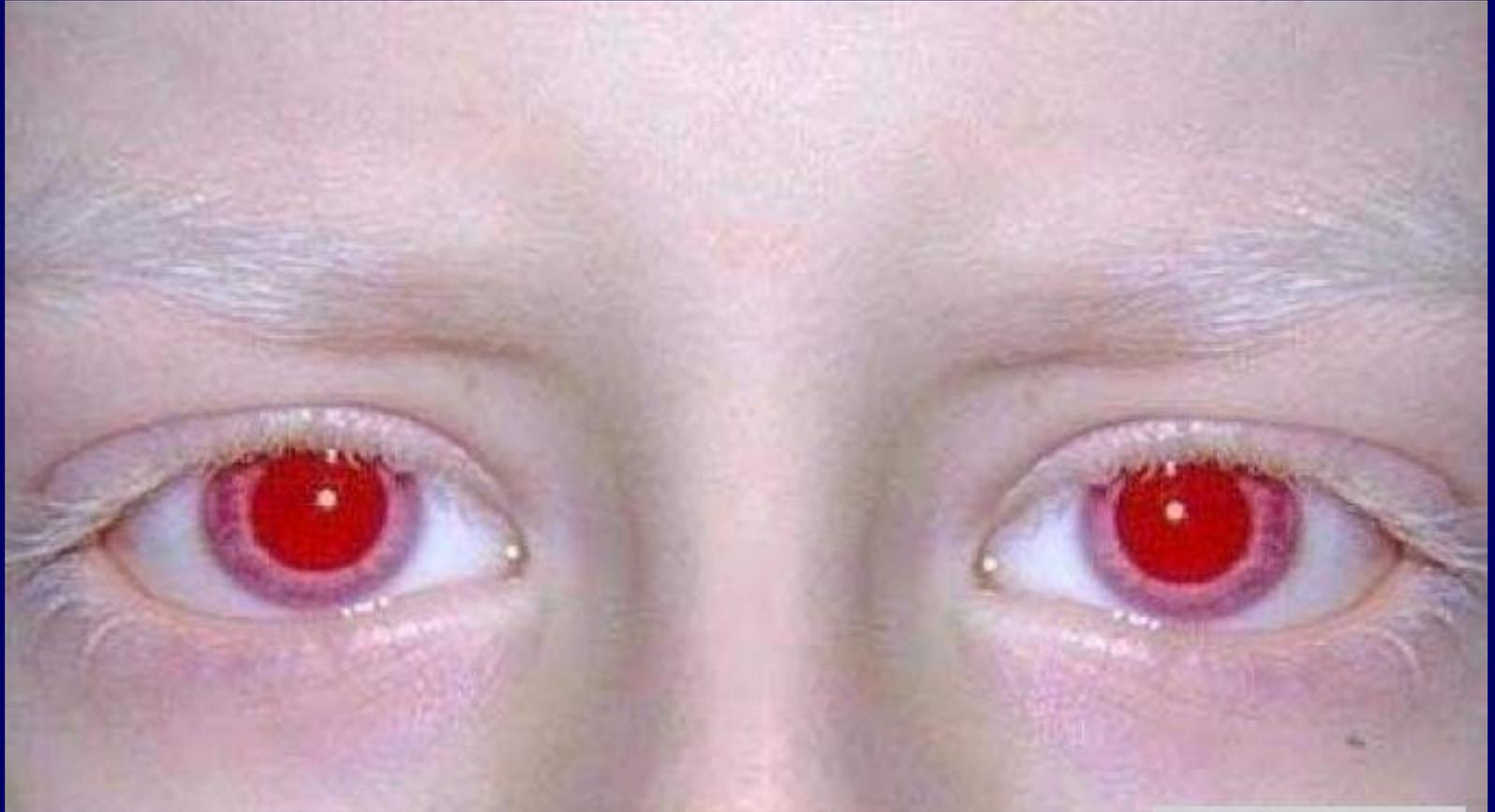


phényllactate



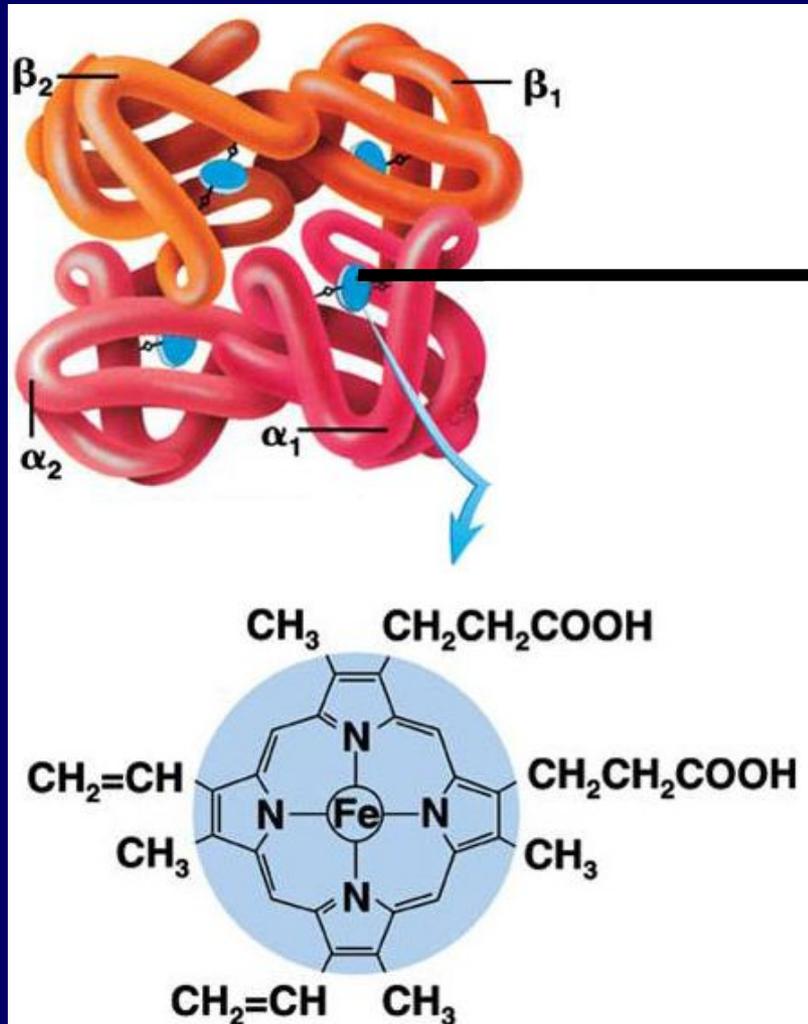
phénylacétate

L'albinisme

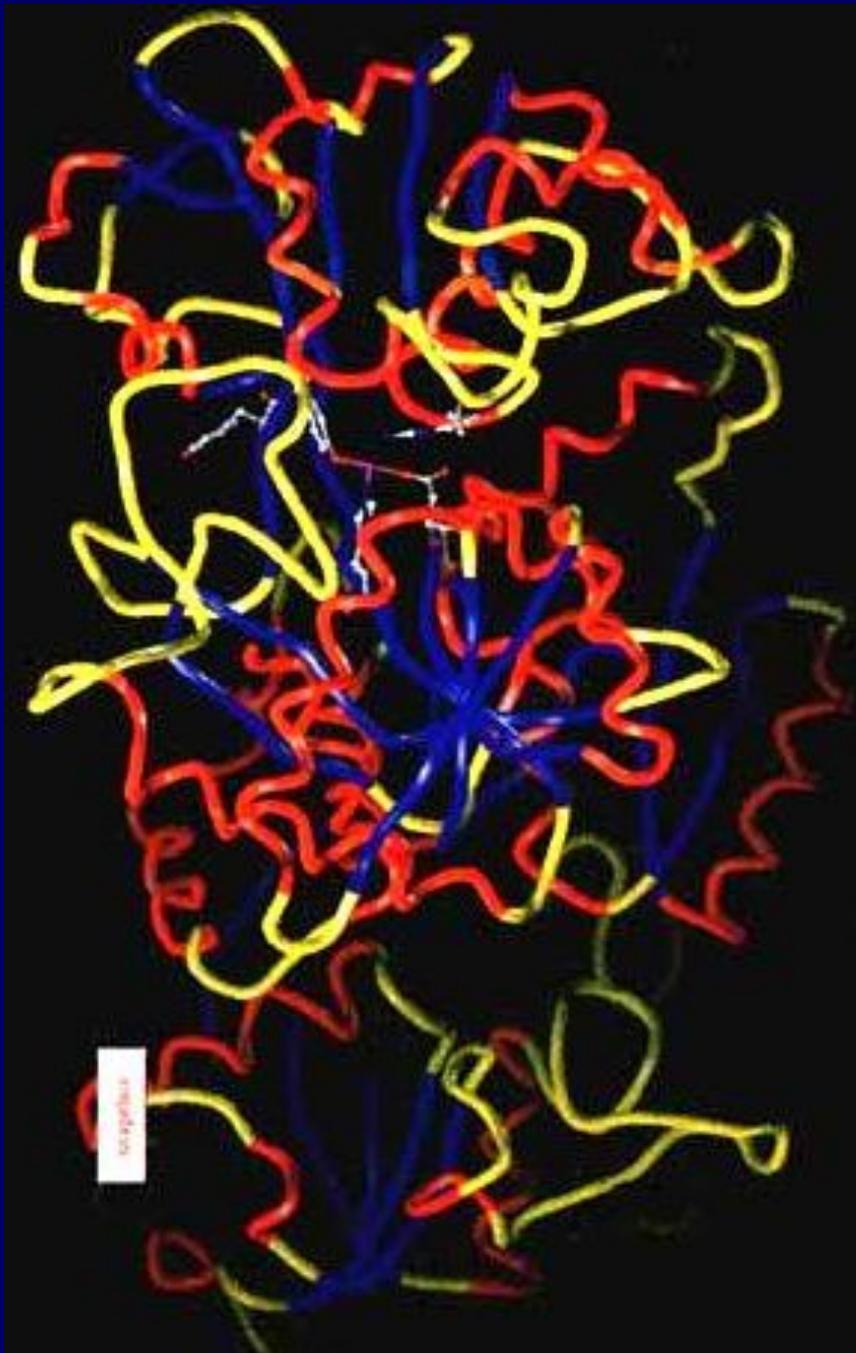


Métabolisme des protéines conjuguées

L'hémoglobine

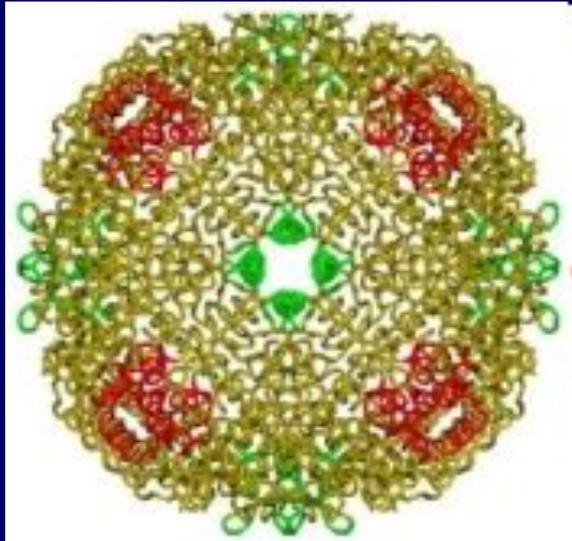


hém

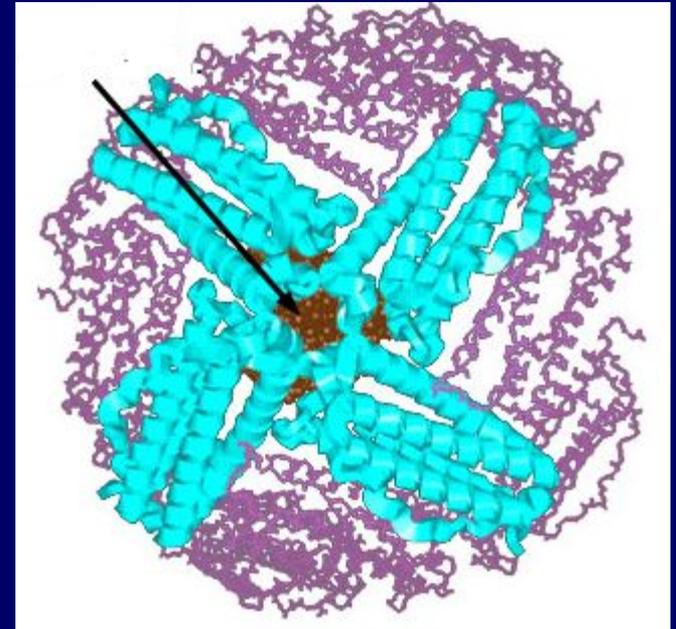


Le transferrine

Formation de la ferritine

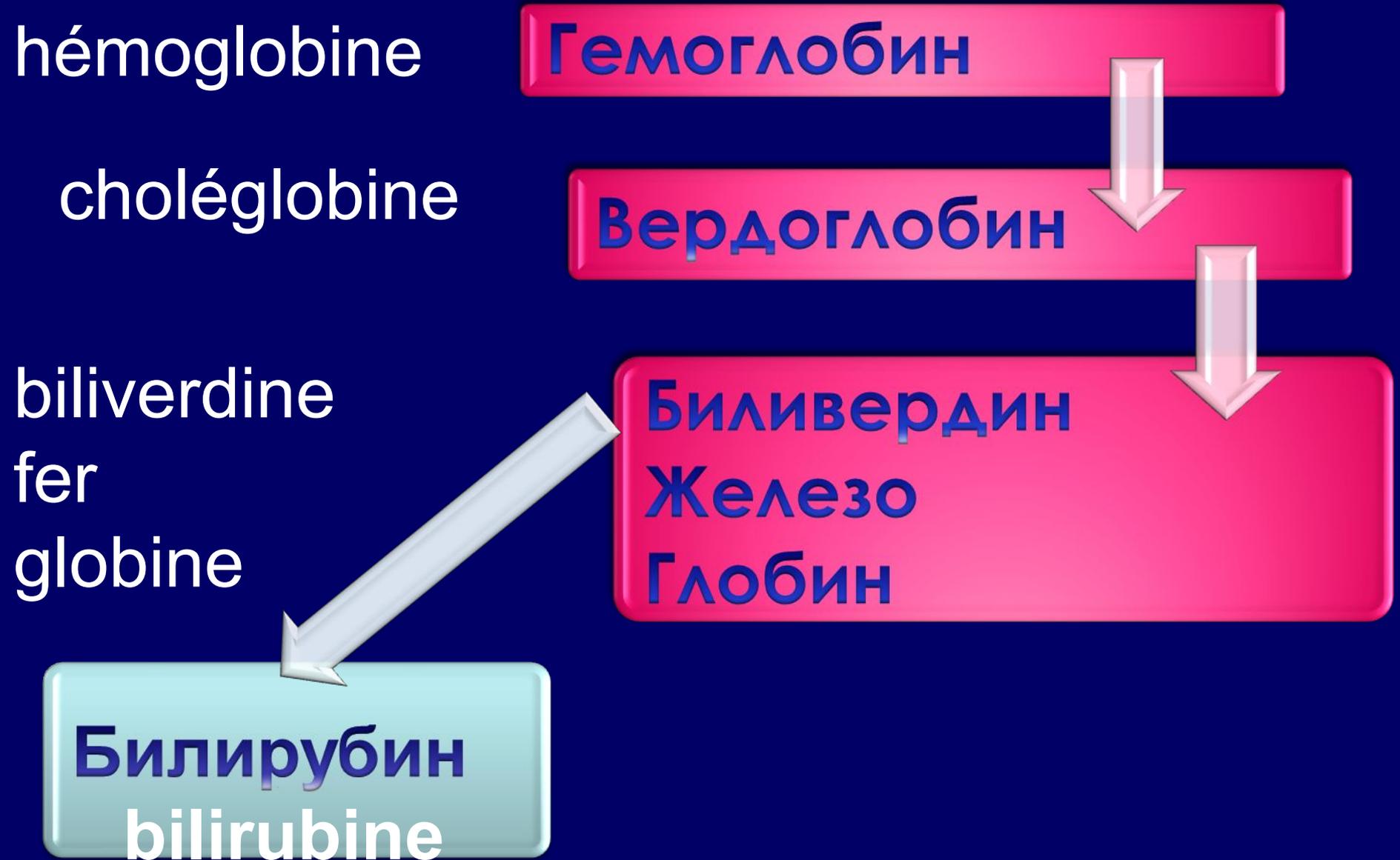


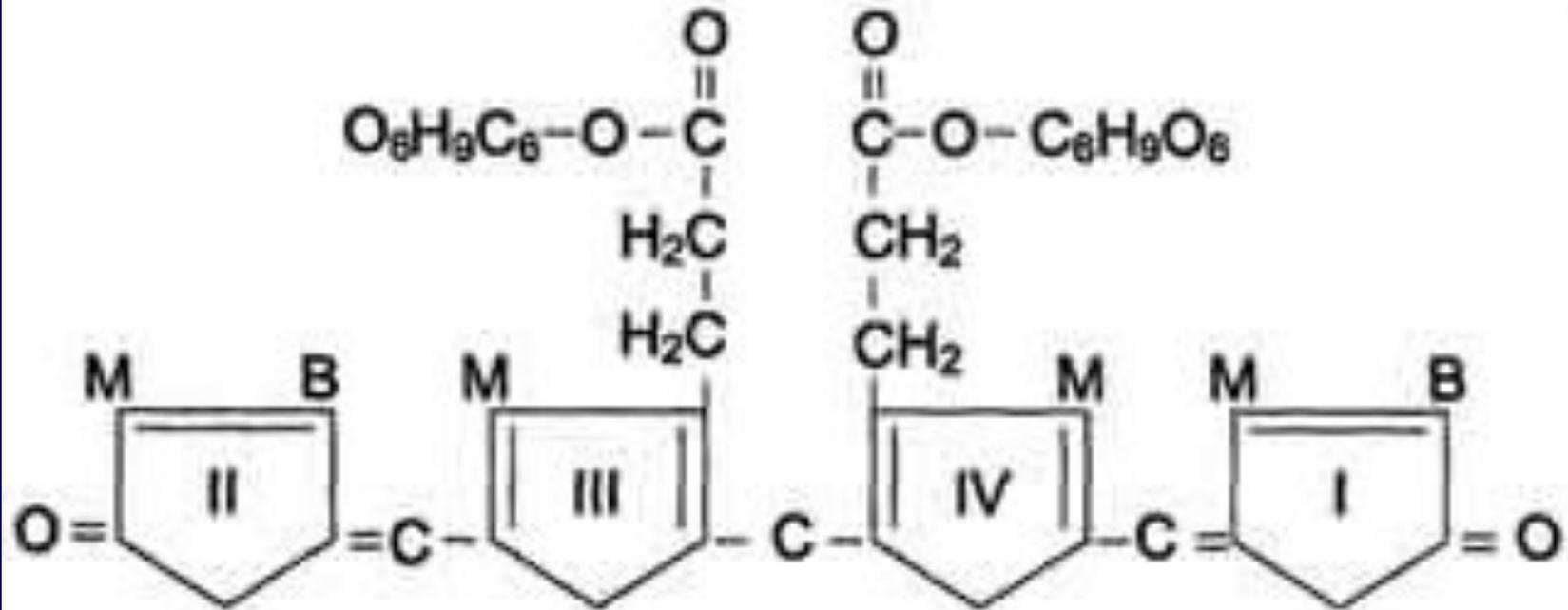
l'apoferritine



la ferritine

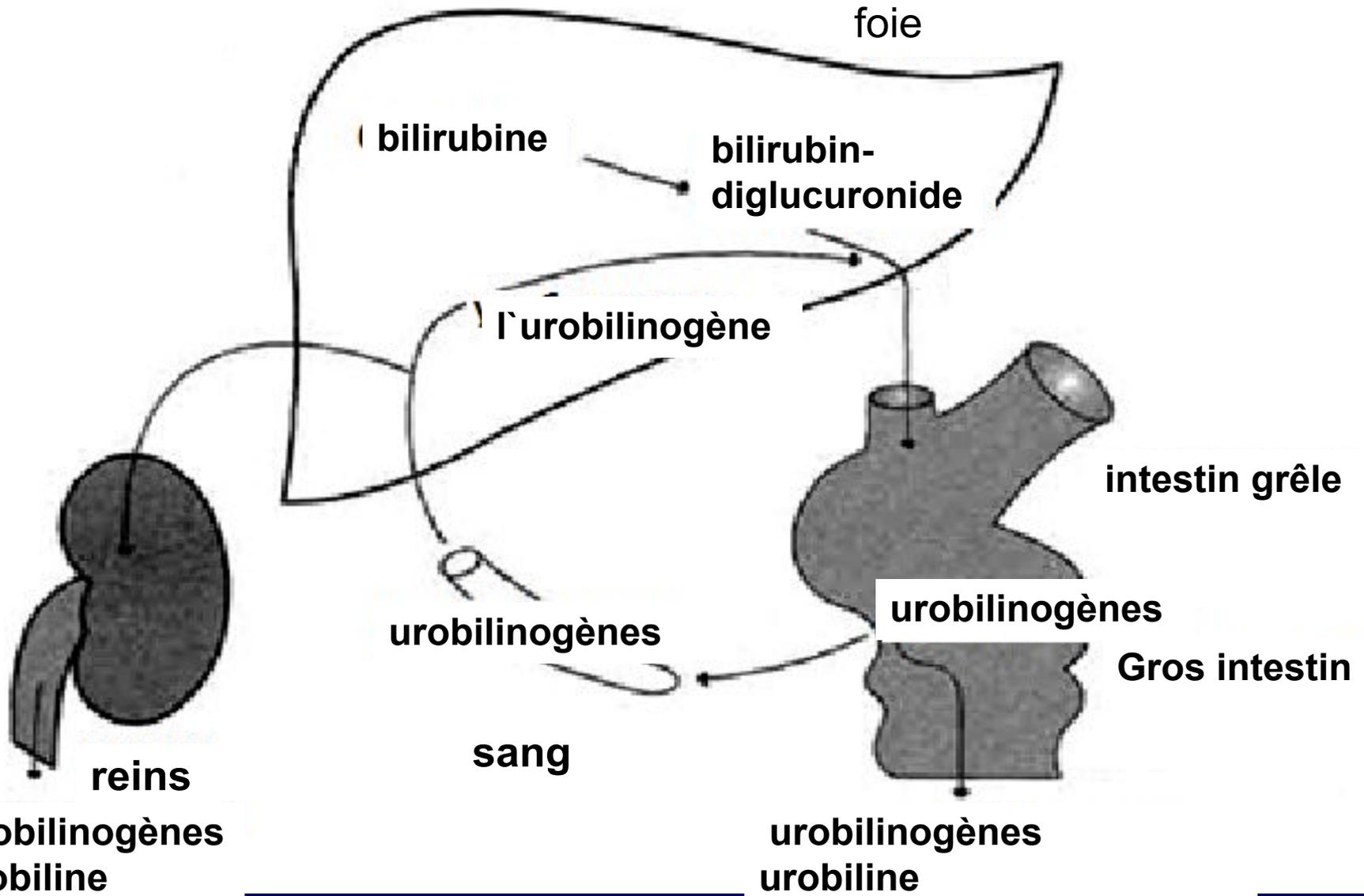
La décomposition de l'hémoglobine



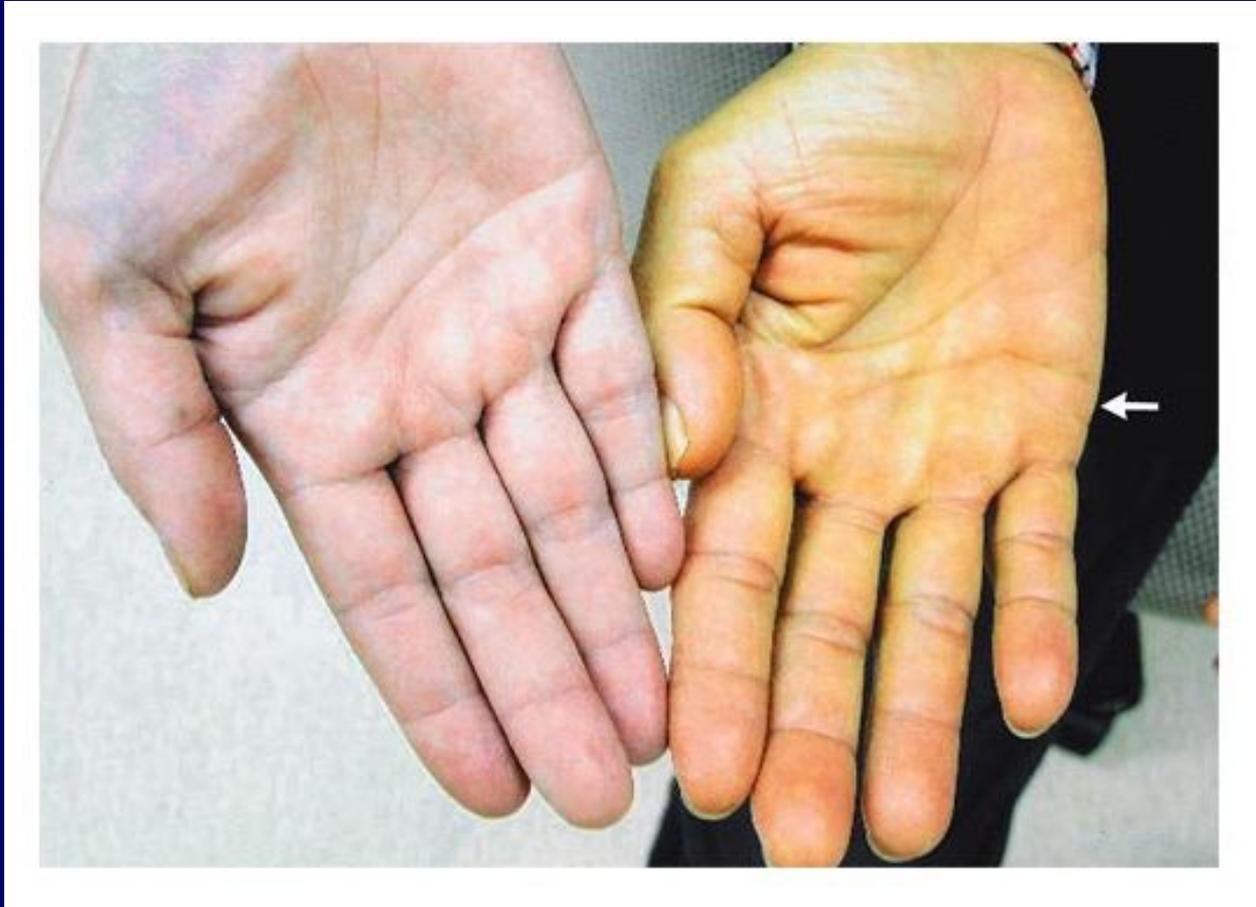


**la bilirubine conjuguée
(droite)**

Le catabolisme de la bilirubine



Types des ictères (jaunisses)



DIAGNOSTIC DIFFÉRENTIEL

Type de l'ictère	sang			urine		fèces
	bilirubine			Bilirubine conjuguée	urobilinogène	stercobilinogène
	générale	Non-conjugée	conjuguée			
L'ictère hémolytique	↑	↑	N ou ↑	0	+	↑
L'ictère hépatocellulaire	↑	N ou ↑	↑	↑	0	0
L'ictère cholestatique (hépatocanaliculaire;)	↑	↑	↑	↑	+	↓

N – norme;



– élévation;



– abaissement;

0 – non déterminé;

+ – déterminé

La synthèse de l'hémoglobine

Synthèse de l'hémoglobine

succinyl-CoA + glycine



acide 5-aminolévulinique



porphobilinogène



hème

+

globine



hémoglobine