

Welcome back students.

We will continue with this unit.

Module #2 standard free energy change.

An equilibrium constant coupled reactions

and additive nature of standard free

energy change the outline for this

lecture is standard free energy change

and equilibrium constant and the additive

nature of standard free energy change.

The learning outcomes are you will

be able to describe the relation

between standard free energy change

and equilibrium constant and you will

gain knowledge about the additive

nature of the standard free energy in

the previous video we have already

learned the terms gives the energy.

Another state functions.

Now using these terms

will find new relations.

The equilibrium constant.

An equilibrium state concentration of reactants and products, rate of their properties and reverse reactions are equal and no further net change occurs in the system.

So if we take the reaction a moles of A + B moles of E giving see moles of C + D moles of F.

The equilibrium constant for this reaction can be given as the molar concentration of C raised to the number of moles of C.

Into the molar concentration of the reactants raised to the number of moles of the reactants.

Upon the molar concentration of A raised to the number of moles of A and the molar concentration of E raised to the number of moles of E.

So this is the equilibrium constant.

Now the next term standard free energy change a reacting system continuously

changes to attain equilibrium.

The magnitude of the driving force

which moves the reaction towards

equilibrium can be expressed as

free energy change of the reaction,

that is ΔG .

So under standard conditions.

Say 25 degrees Celsius temperature when

reactants and products are initially

present at one molar concentration

at one atmospheric pressure,

the force driving this system

towards equilibrium is defined as

standard free energy change which

is denoted as ΔG° .

Each chemical reaction has a

characteristic standard free energy

change which may be positive,

negative,

or zero depending on the quality

and constant of the reaction.

So ΔG° indicates in which direction and how far by given reaction must go to reach equilibrium under standard conditions, this G° is a constant.

It has a characteristic, unchanging value for a given reaction, so we know that ΔG° and K will have got up relation and they are independent.

Now if we consider a reaction similar to the first one.

Very touchy of this reaction depends on the concentration of the reactants and the products temperature and pressure, so the relation can be given as ΔG .

G is equal to ΔG , not plus RT .

Concentration of series to see molar concentration of the race to the number of moles of the upon

molar concentration of a raised.

A number of moles of a an molar

concentration of B says to the number

of moles of B wherein ΔG° is

the standard free energy change.

R is the gas constant is the

absolute temperature,

so this is 1 equation we have got.

Now we have to find out the relation

between ΔG° and K equilibrium.

So since we're tackling

with biochemical reactions,

we keep a standard pH of seven.

An estimate as standard transformed constants

and returned with a prime cheap prime

and K equilibrium prime to distinguish

them from the untransformed constants

used by chemists and physicists.

OK, so we use this prime

for biochemical reactions.

The standard free energy change

can be calculated from the

equilibrium constant $K_{\text{equilibrium}}$.

So ΔG is equal to

$\Delta G^{\circ} + RT \ln$ after the

concentrations of the products divided

by concentrations of the reactants.

So since reaction is at equilibrium,

ΔG is equal to 0.

So since ΔG is equal to 0,

the equation will turn into ΔG° ,

ΔG° is equal to

$-RT \ln K_{\text{equilibrium}}$,

since $K_{\text{equilibrium}}$ is equal to

the ratio of the products to the

Reactance there is a G° ,

ΔG° is the standard transform

free energy change and we can.

Convert $\ln$ into.

Log by using two point minus 2.303 RT .

Log of $K_{\text{equilibrium}}$.

So now using this relation.

We can predict the reaction.

If the equilibrium is more than 1D G,

not will be negative and the

reaction will proceed forward.

If the equilibrium is equal to 1,

delta G not will be 0 and the

reaction will be at equilibrium.

If he equilibrium is less than one,

then delta G knot will be positive and

the reaction will proceed backwards.

Now will take an example to understand this.

Calculate the standard free energy change

of the reaction catalyzed by the enzyme.

For school, gluco mutates,

so we know this reaction.

Glucose 1/4 spade converts to glucose

6 phosphate using this enzyme.

So given that the reaction began

with \$20 million of glucose,

one phosphate and no glucose 6 phosphate,

and the final equilibrium mixture at

25 degrees Celsius and PH-7 contains

one millimolar glucose one phosphate.

In 19 millimolar glucose 6 phosphate.

So we have to find her.

Standard free energy change of this

reaction and the second question is,

does the reaction in the direction of

glucose 6 phosphate formation proceed

with the loss of or gain of the energy?

So now we solve this.

First we have to calculate

the equilibrium constant,

which we can get by the concentration

of the product upon the concentration

of the reactant.

Since we know the final concentration

was 19 millimolar,

we can divide it by one millimolar

and we will get the answer 19.

And now you calculate the standard

free energy change that is delta

and she not prime,

which is equal to minus ΔG°

enough K equilibrium.

We can put in the values and the answer

will get is minus 7.3 kilo joules per mole.

OK,

now we have the next question whether

there will be gain or loss of energy.

So since the standard free

energy change is negative,

the conversion of glucose one

phosphate to glucose 6 phosphate

proceeds with loss of the energy and

therefore it's an exergonic reaction.

Now we'll go to coupling of

reactions in biochemical reactions.

Coupling plays a very important role.

What are catabolic and anabolic

reactions in the previous video?

I had already told you.

What is it?

The exergonic reactions are termed
catabolism which means the breakdown
or oxidation of fuel molecules
and anabolism is the synthetic
reactions that build up substances.

The combined catabolic anabolic
processes constitute metabolic.

Now standard free energy
changes are additive.

If we have a sequential reaction
wherein A is giving B & B is converting
to see then the ΔG values
values are added so in the reaction
A converting to B we have ΔG_1
 ΔG_1 and we have we
converting to see we got ΔG_2 not.

So there that the total will be equal
to ΔG_1 not plus ΔG_2 total.

So I turn more dynamically,
if an unfavorable reaction can be
driven in the forward direction by

coupling it with an exergonic reaction.

How we will see in the next slide?

So if we have the synthesis of

glucose 6 phosphate glucose plus

inorganic phosphate gives us

glucose 6 phosphate plus water.

We know this reaction.

And for this,

ΔG is 13.8 kJ.

And this will not proceed

spontaneously in this direction,

because it requires energy to be absorbed.

ATP plus water will give you

ADP plus inorganic phosphate.

This is the second reaction wherein

ΔG is negative 30.5.

So these react.

This is an exergonic reaction.

These reactions share this

common intermediates,

inorganic phosphate and water,

and may be expressed as glucose

plus inorganic phosphate giving

glucose 6 phosphate and water.

And the second reaction,

ATP,

does water giving ADP plus

inorganic phosphate.

So summing the Delta G north

of both these reactions we get

minus 16.7 kilo joules per mole.

So the overall reaction becomes exergonic.

Energy stored in ATP is used to

drive the synthesis of glucose 6

phosphate even though its formation

from glucose and inorganic

phosphate is and the governing the

pathway of glucose 6 phosphate.

Formation from glucose by

phosphoryl transfer from ATP

is different from reactions.

Yeah, thank you.