

welcome back to electrochemistry class

so let us look back what we have studied

so far in this few lectures so if we look at the topics that are expected to be covered in this series of lectures are like conductance in electrolytic solution

that we have taken up in details and we have also tried to explain how this conductance of solution will vary due to several factors like concentration may be like dilution and maybe temperature in this regard we talked about also specific conductance and specific and molar conductivity then we also

discussed this variation of conductivity with concentration and important important thing

that important concept we have learnt over here here is ohm's law of independent migration

of ions that at infinite really dilute situation all the ions are free to move and there is

probably i mean most likely there is no inter ionic attraction that is internal attraction

is a minimize and ions can move ions can freely move and therefore therefore you know molar

conductivity of an electrolyte that is basically molar conductivity some of the molar

conductivities of the constituent ions then we have try to learn

electrolysis electrolysis means it is it is electro means while you apply some

electricity from outside through a pair of electrodes then your material is lysed that

is broken down into pieces like water if you electrolyze it is broken down into pieces like

hydrogen and oxygen that in that case you need to supply electricity from outside means you need

to supply energy

so that

so that the reaction i mean this breaking reaction lysis reaction

can take place and also we talked about this laws of electrolysis that is

paradise law of electrolysis just we gave the elementary idea then we also talked about

this dry cells like dry cell then also we talked about this electrolytic cells then galvanic

cells then lead acid cell that is lead acid battery which is used in like in car

etc also we have talked about electromotive force of a cell and

electromotive force is

nothing but the reversible cell potential when you draw you know zero current from the from

the cell then the cell reaction net cell reaction ΔG becomes i mean the cell reaction or

the electrode reaction become reversible that is perfect reversibility condition is

maintained and therefore thermodynamics of reversibility reversible

thermodynamic

principle can be simply applied to that and we also talked about standard electrode

potential that standard electrode potential is the potential when in activity of the of the electroactive substance is an unity or maybe it is an it is an i mean one molar concentration or may be unit an concentration then you also used nernst equation although derivation was not done over here just the statement of nernst equation that connects the cell potential connects the cell potential with the reaction quotient of the reaction that is involved in the in the cell and also we have given we have discussed a number of applications an in connection with this electro chemical i mean this emf measurement like ph of the solution how ph can be measured using emf measurement or maybe how can we follow the precipitation i mean this redox reaction that we also have discussed over here and also how to find out this solubility product of a sparingly soluble salt and that we discussed in in the light of this emf measurement now how emf can be measured it is not with the help of a voltmeter but it is it is a basically a potentiometric measurement where you draw a zero current so it is called the poggendorfs an compensation method so poggendorfs compensation method has been applied to apply to determine the emf of the cell various types of electrodes i mean half cells were also considered and we also attempted construction of this of cells ok construction of cells based on the requirement based on the net reaction we have given one or two example of that then we we an found out the relation between the gibbs energy change and the emf of of the cell ok these are the things that we have covered so far now an some more things are yet to be covered over here an first of all an one is fuel cell another is corrosion these are the two important aspects and and maybe we will just talk a little about redox reaction because this electrochemistry basically deals with this it is nothing but an the redox reaction in i mean it is an electrode process ok it is it is the redox reaction so therefore therefore what what is happening that when you when you construct a cell when you construct an electro chemical cell then there occurs an there occur some chemical reaction and at the electrode at the electrode what is happening that in one electrode there will be you know oxidation and in the other electrode there will be reduction now i already have discussed this point that when you dip one metal when you dip one metal in its electrolyte solution then either the

metal has the tendency to accept electrons from here from this and becoming it will

become more negatively charged it will acquire more negative potential over over here and this

one becomes positive or in other words this metal will lose electron and then it will

get dissolved over here ok in this way this acquires a negative potential with respect to the

solution and the reverse may also take place that that you have the electrode which is dipped into

the electrolyte solution and what will happen that that the the ions over here ions over here

will accept electron and electron from here and get reduced and it will it will be deposited

on on to this metal

so in that case in in that case it will become your positive richer so

when you connect these two then it will form it will form an

electrochemical cell only thing

is that while constructing this electrochemical cell you have have to keep this in mind keep

this in mind that your net cell potential net cell potential will be will be greater than

0 if it is greater than 0 means the cell reaction as represented for this e cell will be spontaneous

that is ΔG will be negative that is reaction will be will be spontaneous in the direction shown

with respect to this e cell greater than zero

so redox reaction means basically in in in in in

case of redox reaction what we do that we pick up a specific redox reaction and then what happens

that we we dip one electrode in that redox couple

so that this redox reaction is taking place by

means of the fact that this electron exchange will occur through this electrode and as a result

of which one electrode will acquire some positive charge and the other electrode will acquire

some negative charge and when these two are connected that means the potential relative

potential of these two negative potential of these two are same

so therefore with

respect to this one it is negatively negative it will acquire some negative potential

and this will acquire a positive potential therefore from external source if you connect with

with with wire then current will flow from this direction from here to here and electrons will

flow like this

so therefore this redox reactions are are very important

so if there is no redox

reaction then you know if it is say for example is say a precipitation

reaction say for example

$AgNO_3 + Cl^-$ that gets you $AgCl + Nitrate^-$

so directly you cannot

you know directly you cannot since it is not a redox reaction you cannot ah

you know form a cell like this but what you have to do you have to have have a an indirect way of measuring the parameters for this reaction that you will construct a redox process such that the net reaction will be like this so therefore redox reactions are of immense importance as far as this you know electrochemistry study of electrochemistry is concerned next is another important thing that we should keep in our mind that while while we were discussing this electrolysis then what happens that that you have two electrodes and you apply some ah potential difference between these two electrodes so it is grossly stated that that negative ions will be attracted by the positive electrode and negative ah negative ions will be attracted by the positive electrodes and positive ions will be attracted by the negative electrodes so that generally happens when these ions are this ions are are in close proximity to the electrodes so that it can it can you know it can fill the potential potential fill a potential gradient but if if this one is placed at a very long distance then practically this ion has the option to move in any direction in any direction means it can move in this direction or that direction okay so but those ions those negative ions which are which are close to this electrode will be attracted and ah and if the you know potential is such that that this ah you know electron transfer is favorable then this this ion will be discharged i mean this ion will be will ah i mean we will will you know lose one electron over here and then it will it will be it will be discharged in the same way this will happen for the plus so therefore therefore discharge is taking place or the redox process is taking place very close to the electrode but here it has got the provision to move randomly but statistically what happens that if these these ions are you know converted to their corresponding discharged counterpart then on an average the the concentration concentration of the negative ions will be reduced over here the system so system will face if you know a concentration gradient there a concentration gradient will be produced so in order to you know equilibrate this gradient i mean in order to minimize this gradient again you know ah negative ions will be will be coming ah you know in the vicinity of this so this way things will happen and net effect is that as if the negative ions are attracted by the positive electrode from

from any position

so it is it it is not as simple as that

so so only there when when when these

ions face the potential difference or a potential of this electrode only that happens when

this is coming at some appreciably close ah separation because

so that it can fill the field

otherwise at a larger separation you may not be able to or the iron may not be able to feel

the field

so these are the few things that that that you should remember next we

will let us ah let us turn our attention to ah to those two topics one is this ah

fuel cell and the other one is ah corrosion fuel cell now we have we have learnt ah cells

cells means the the device that can supply electricity like the battery or the lead

accumulator or lead acid cell that is pbpo2 this this thing now what happens that in case of ah in case of normal cell

like say for example lake lances cleanses ah this dry cell we have discussed

this point that this dry

cell is good until until all the reactants are are exhausted or you cannot keep this dry cell

for very long time this is because of the fact that this this is this will discharge there will

be a self discharge that is there is there is internal resistance

so electricity will will

you know ah we will will you know flow across the electrodes against that internal distance and

it will automatically get discharged if you keep the cell for long time and that is also going

to happen in case of this lead acid accumulator that that they are also there they are also

internal you know discharges is possible

so therefore

so these electrodes i mean this this

ah cells are good until the the the you know ah chemical substances ah are you know available over

there i mean those chemical substance which are involved in the chemical reaction to produce the

the the electrical energy

so therefore

so supply of electrical energy from the chemical reactant

ah basically from the from the chemical reactant is stored within this cell and then then then

what is what is happening the reactants are reactants are consumed and one situation will happen that

all the reactants will be consumed

so what will happen then if all

the reactants are consumed then nothing is remaining

so the cell reaction is

not going to is not going to proceed any further process in the further means no further cell

reaction will take place

so cell will stop you know functioning
so cell will become dead
so when reactants are consumed cell will become dead and then what do you do we
throw the cell
cell away if we go to the market we buy new sets of um cells and then plug in
those cells in in the
appropriate device
so therefore there is no way there is no way to reuse the cell or at least
the outside i mean the cover of the cell ok what happens in case of your lead
accumulator you can
recharge it and then you know you can use it again
so a number of cycles can be used
for recharging again and again but here it is not possible but suppose if the
situation is such that
there is a mechanism by which you can take out the used chemicals and then you
can feed the cell
with new set of chemicals unreacted chemicals then what is going to happen
then you can expect
that cell to gain you know its power again that means cell will start to work
again okay so
that's why as if you are i mean this thing means as if you are filling the
cell like you fill
your motorcycle or you fuel your car you go to the gas station and you know
pay money and then you
fill it you fill in fuel it means either you put petrol or diesel in the in
the fuel chamber or
fuel tank
so so that means when once this fuel in the fuel chamber is exhausted you put
new
new fuel in and then the system will continue to work again
so that is why that
so if there is a
mechanism by which you can you know refill it you can take out
the the bad ones and
you can take the new ones
so ah
so therefore ah so that means you are filling the cell and therefore
you can drive this
so you can fill the cell fill the cell and ultimately drive the cell
fill the cell or drive the cell
so this principle i mean this idea was first
demonstrated by group in eighteen thirty nine first this idea was implemented
it was demonstrated by group in 18 39 see see the idea was so
old that in that time ah people could think about that whether we can
fuel the cell ok
so it was i mean at that time it was known that water i mean as a result
of electrolysis of water water is decomposed electrolyzed to form H_2 and O_2
okay
so what group
tried was to was to recombine two you know ah these two i i mean this ah these
two recombinants
recombine water and i am sorry i mean hydrogen and oxygen to form water okay
so ah so
basically these two gases are allowed to recombine in a specific fashion

so basically

it will be the reverse of electrolysis

so reverse of ok reverse of electrolysis ok

so um so

recombine H_2 plus O_2 to produce water and this will cause a potential difference ok against two electrodes dipped over there

so a potential difference difference will be there across two electrodes present ok

so what is happening over there again you need to consider anode process and anodic and cathodic process

so anode process anode process anode process is H_2 gas

that gets you $2H$ plus plus twice electron and corresponding potential is

0 volt because H_2 to H plus remember this standard hydrogen electrode

case that where you know electrode potential at all temperature is assumed to be zero

so

so so that is the concept and cathode cathode process cathode process is half O_2 gas plus twice H plus plus twice

electron that gets you water and here E° is equal

to plus one point two three two three volt plus one you remember if you

think about the reverse reaction then it is minus one point two three volt which we already

have used many times

so what is the net reaction net reaction is H_2 gas plus half

O_2 gas that gets you H_2O liquid where E° is one point two three volt ok

so this is the this is

the basic idea then in 1959 first working hydrogen oxygen based first working hydrogen oxygen based fuel cell was invented ah invented by Francis T. Bacon ok now alkaline electrolyte nowadays

alkaline electrolyte is used alkaline is used in modern cells now what is the reaction anode reaction anode reaction is H_2 gas plus

$2H$ minus gets you $2H$ plus $2O$ plus twice electron again E°

is equal to zero volt then cathode reaction half O_2 gas plus two water plus twice electron that gets you

to which minus E° is equal to plus one point two three volt and

net is net is the same reaction net net reaction is H_2 gas plus half O_2 gas that gets water with E° equal to 1.

23 volt

so what is the what is the you know

pictorial representation of this

so pictorial representation will be like this it is like this that in one side you put oxygen in the other side you put

hydrogen H_2 to fuel ok you have porous electrode

so that this hydrogen can diffuse in and also oxygen can diffuse and

this is a porous electrode

so porous electrode and this one is anode this is cathode

so minus this is plus if you put this

against some external load then like then electrons will flow this way

current will flow this way the other way and the this anode reaction will be $2H_2$

that gets to four H plus plus four electron cathode reaction will be

four H plus plus O_2 plus four electron that gets through water that means it

is it is alkaline

so therefore any which and each plus will be moving
this direction

so this H_2 will will diffuse in and then it will be converted to
 H^+ plus and then it will move from this direction to this direction

so here you in input of

air that means O_2 and here excess of air excess of air and unused O_2 is coming
out so

this is a porous cathode this is porous anode and this porous cathode porous
anode

and the net net process is this ok

so only problem is that that

already i have discussed at one point that this oxygen consumption this
one this oxygen consumption process it's a it's a slow kinetically
slow process

so kinetically kinetically slow

so that you know puts a

problem for efficient problem against its efficient functioning okay

so therefore this

porous cathode if we replace this porous cathode with with some expensive you
know platinum cathode

then it has been found ah that such problems are mostly you know eradicated

so only problem is that

platinum is a costly metal

so it will increase the pricing of this device pricing of
this the cell

so therefore these are the this this is one of the one of the important
drawbacks for this this particular fuel cell

so what is happening that you are

feeling you are failing and this is the this is the ah this is the you know you
know excess of air or oxygen this is taken

out and also the reaction product is water so you see that as i when i started
discussion with

this then i was telling one thing that if you can remove the reaction product
and if you then you know feed the cell with new set of chemicals

have the same chemicals but new batch of chemicals then it will be filled i

mean this you are you are

feeling it ok you are feeling it and then after this reaction is taking place
again what will

happen you take out water

so this way you keep on doing

so you are failing you are gaining energy

and then you are taking out the reaction product this way this will continue

so therefore only

thing is that only problem is with platinum but you see you know this oxygen
it is readily

available from here hydrogen fuel you can you can get from electrolysis of you
know

water acidic acidic water you can you can get

so theoretically theoretically its

potential differences theoretically theoretically its potential difference is
about

one point two three volt at two ninety eight kelvin but in it has been found
because

of ah several other ah problems drawbacks ah and that also depends on the pressure of this air air then hydrogen and then nature of the electrode and and so if these are considered then then it has been found i mean if you consider all this um if those are the those create trouble then it will be you will be means actually what you get is that open circuit circuit the voltage is 1 volt around 1 volt it is not more than 1 volt and if you put load with load this reduces to about 0.5 to 0.8 volt ok so this this you know um this is called you know a fuel cell so so you are foiling it and you are getting energy and it is it is continuing means you are continuously fueling and you are getting the energy so that way it is it is it is happening ok so that completes the basic you know discussion with respect to the fuel cell then we will move on to move on to another another important issue that that is called corrosion corrosion that is corrosion means because of corrosion you know suppose you have a shining you know iron uh you know material iron iron pot in your hand and say you are not using that for long time and you are keeping in keeping that nice container iron container outside is shining ok you are keeping it here here means in open ear that may be may be which the problem is more during you know any time so you will be finding that after some days after some days this this shining color shining shining of shining nature of this material this pot is gone and some spots come brown spots come which are called the rust ok so therefore the therefore the you know deterioration of this material is there and this happens particularly if the place is you know is having damp or during you know rainy time but during winter time the situation is little better that the probability or the possibility that the materials will lose its last year as a result of rusting that is that is reduced or if you keep in you know airtight con container or maybe if you keep in a container material in a container just removing the ex i mean removing this moisture by keeping some hygroscopic material inside that you will take take you know moisture like calcium oxide or like that so so then then this probability or this chance of this corrosion will be reduced so basically it is the chemical reaction that is taking place at the smooth surface of ah various metals ok

so so technically corrosion means spontaneously the metals will return to their to their ore condition poor condition means as if you

are returning back your metal to i mean to their ore condition that means to their compound state ok

so electrochemical ah corrosion electrochemical corrosion means this ah corrosion as a result of chemical reaction at the surface and some small cells are formed and the net free energy for the cell reaction is such that the process is spontaneous and ultimately the end product is that you say that the surface is corroded

so therefore electrochemical corrosion corrosion is is is very important

so basically $M \rightarrow M^{n+} + n e^{-}$ so

this is the process

so and this is facilitated in presence of a suitable electron acceptor that means this electron is free if electron is accepted by by something then the process will

be facilitated that metal will form M^{n+} and therefore smooth surface of the metal will get corrosion

so in presence of this is facilitated by facilitated by the presence of presence of suitable electron acceptor ok and this is also known in corrosion language

that it is these are called depolarizer depolarizers sometime what happens that a

thin film of water or moisture a thin film of moisture moisture that is muscle which

is in the form of adsorption absorbed moisture can also be can be very dangerous in respect of corrosion that it

will promote corrosion of the promote corrosion of the metal surface

so so basically the corrosion system

the system where corrosion is taking place that may be regarded as that may also

be called as it may be

so corrosion system corrosion system or the system where corrosion takes place can be regarded as regarded as a short circuited circuited electrochemical cell cell in which in which the anodic process anodic reaction could be we say for example metal to

say metal two plus equals plus two is electron the celebrated example may be iron because the corrosion problem of corrosion is mostly with iron because we complain mostly

about corrosion i mean i mean the effect of corrosion or i don't like it i mean that people

does not like it that ok it is corroded

so it looks bad

so mostly ah that is involved with

that is involved with this iron

so iron two plus equated plus twice electron and the cathodic processes may be and the cathodic process

so this is an this

is an ah anodic process that is that is at as if it is an anode reaction

so anodic process and cathode corresponding cathode reactions could be $H^{+} + e^{-} \rightarrow H_2$ plus electron that is the that is an acceptor that gets to half H_2 gas

so this is an acceptor

so therefore in presence of acid that electron is accepted that that means the electron the electron which is which was liberated from the metal while it produces this m plus or m two plus that will be that will be accepted by h plus to produce water ah sorry to produce hydrogen or maybe m two plus plus twice electron then it gets you m solid where m is the metal ok

so corrosion is a is a is a two step process as i was discussing that one one part is

a cathodic part another part is a is an anodic part

so cathodic cut means you know metal

will lose the electron and someone will be there to someone means some other agent will

be there to accept the electrons

so that the drive of the driving force of the process will be in in forward direction

so corrosion is a it is a two step it is having two steps having two steps say for example if

we talk about this iron corrosion then first one is that in presence of you know

moisture a film of moisture what happens that that the surface is coated with moisture means the surface metal surface is having adsorbed moisture on it

so therefore first step is iron that makes iron two plus iron ferrous ion plus twice electron

so it it it dissolves okay and the the metal becomes this metal becomes excess negative charge because of this one it is having excess of negative charge suppose if

there is a mechanism that by which you pump this electron out then the process will be more more

favorable but if there is a mechanism by which you put more electrons from outside or or if you make

a situation that that that the environment is such that this particular system is reluctant to

you know reluctant to remove this electron from here that means once the electron is you

know accumulated over there it is very difficult to you know get rid of that then the process

will not be very favorable

so so iron will try to iron two plus or iron will try to remain is in its elemental ah state ok

so so what happens second step is

so correction corrosion

will will continue smoothly as the depolarizer or or the electron acceptor is removing removing that that this electron okay say for example acid as i mentioned acid

so two

h plus plus twice electron gets you h two or or like this is a b means like say for

example if you have a more noble metal more noble metal

so metal ion

so what will happen Cu^{2+} then plus two is electron that gets you Cu okay or maybe even even if there is oxygen situation is like if there is oxygen and this oxygen is available in air so if there is oxygen what is may be a problem O_2 so O_2 can again create an additional travel like this plus four water that produces four electrons plus four H^+ minus ok and this four which minus will combine with i mean this H^+ minus will combine with iron two to produce this hydrous ferrous oxide this will produce hydrospheres oxide which is called the rust so so it will look bad i mean the the surface will look bad so then then how to get rid of this how to get rid of this means so these are the possible situations that you have like problem is that that you have a copper sulphate solution that you know some somehow copper sulfate solution has spilled on the iron surface and little bit of moisture is there then what is going to happen thing is that that the tendency of copper and because this is in in the lower side i mean it is easy to reduce than this one so this one will be difficult to reduce so so the thing is that this reaction will proceed at the same time this reaction will also process so metal to metal ion and here metal into metal this coupled process will take place because the net thermodynamics of this process is very favorable so that's why more if more noble metals are present then it is a trouble ah more noble means more noble than iron or the corrosion of the concerned metal ok and also suppose if you if there is a spillage of acid then it will you know induce you know more corrosion suppose you have a little corrosion if by some means some acid has spilled onto the onto the little corroded surface then the corrosion will be more efficient and then corrosion will spread all over the surface of this iron so this nice ah you know shiny surface will be will be you know damaged so therefore corrosion is a real trouble ah to us and at the same time suppose you have you can you can paint the ah the i mean the shining iron surface with the help of a of a specific pain the point is that that so it is this paint is on the surface surface of the of the this metal so if your paint is not that good then what will happen when water will fall on the on the surface of the paint it will

penetrate and go inside and
will stay in the in between this iron surface and this this paint coating
so that means it is
it is practically a thin coating of moisture water and if that moisture stays
for long time what will
happen
so this surface i mean this metal surface on which this nice coating of this
paint is
there
so in between that there will be corrosion
so when corrosion is there this hydrosphere
oxide will come will will will prod will be produced over there and this
hydrosphere
is oxide is basically having more volume ok
so it will it will have bigger
larger volume
so it will just bulge out of ah out of the surface
so therefore
therefore it will it will look that that some as if some blister looking thing
on the surface
of i mean across the across the paint ok
so therefore if you give a little pressure then
it will break and then paint will also go and ultimately the good look of this
painted stuff is
lost
so therefore it is very important that if you use very efficient i mean
material ah very
efficient in the sense that if you use a water repellent material like teflon
or or
similar material then the possibility that this water moisture will go inside
and reach
the metal surface that will be reduced i am not telling that it will be it
will be removed
but it will be reduced ok
so that is why that is why you know you know covering the metal
surface with appropriate thing is very important ok or maybe in some cases oil
if you have oil you
know oil coating that is also good but the point is if oil is having some
moisture then its a real
difficulty ok
so so um sometime if you cover the surface with ah thin oxide then then that
will
inhibit the anodic dissolution process ok so ah anodic dissolution process
means $m \rightarrow m^n + n e^-$
this anodic dissolution processes is inhibited if you use this oxide film or
oxide paint and
if metal is ah now the point is if the metal is this metal surface say say on
which this
corrosion is taking place if this is that one is little biased with negative
potential
or if it is having excess of negative charge then the tendency that an
additional metal atom
that it will it will lose an electron and give it give it to this metal
surface and it will dissolve

this tendency will be reduced because already the metal is having excess of negative charge
so so
dissolution um will be you know will be difficult in this way dissolution will be difficult
so so
if you have if you have a coating onto this metal surface the metal i mean the coated metal i mean
the metal which with which you will you will coat the iron surface say for example if it is more
ah reactive than iron like say for example if you if you have a coating of zinc zinc metal on iron
then zinc will tend to get i mean zinc itself will tend to get corroded because with with moisture
i mean it will just become zinc two plus plus twice electron and this twice electron means this
twice electron will stay on to twice electron will stay onto the metals ah metal surface metal
means originally the iron surface
so then then what will happen that iron to iron two this process will be inhibited because this is already having excess of ah excess of negative charge so
therefore ah um
so the i mean basically the metal sheet metal sheet i mean irons say for example
iron sheet will have will have negative charge on it because of this reaction because it has
dissolved
so it ha it has left two electrons onto the metal surface
so therefore therefore look i
mean corrosion which which may be a local process that local process will be will be sluggish or
in some cases it may be reduced to a to a greater extent
so that is why you know coating of like
zinc coating on this iron iron sheet has been has been found in in many many cases ok
so thats
why
so these are the ways by which you can i mean you can you can reduce the extent of corrosion
but corrosion is a is is really a problem because i mean in atmosphere you have oxygen in
atmosphere you have moisture ok moisture the amount of moisture may be less or may be more
but it is there
so if you keep on exposing your metal surface this metal if the it is a reactive
metal metal surface ah with respect to this ah moisture then what will happen that a thin film
of air will be there and that will create trouble
so its a real real ah difficulty ah you know in ah
day to day life
so therefore like
so things will be corroded and it will it will guess get rust

and if there is rust then then the longevity as far as the longevity of the substance like ah you know mater you know ah you know material ah like like ah say for example ah so you know like if it is car or motorcycle or maybe may be iron maid substance things so all will be will be eventually damaged if it is exposed to moisture or suppose if it is kept in open ah sky and during you know rainy time it will get you know water and then you will be finding that after rainy season is over it it gets a thin coating of this brownish ah rust on it and if you keep keep it for longer time then it will continue to increase so that is why you need to scrap it and then probably with appropriate oxide coating or may be appropriate metal coating like drink ah can prevent its further damage and may be life of this this thing will be will be you know increased so so ah while some summing up ah what we have studied in this ah particular ah piece of ah lecture that we have tried to sum up the some of most of the things that we already have studied over here during the earlier lectures now in this piece of lecture we have you know talked about this ah this corrosion and also we have talked about fuel cell which is a very important concept and so so this fuel cell we we talked i mean the basic of the basics of that ah has been talked about and also ah the corrosion that is that is a and it is a problem as far as electrochemistry is concerned problem in the sense that materials are getting damaged due to this electrochemical you know processes on the surface so so how this can be eradicated whether it is fully eradicatable or not so ah that is a question but at least we can we can try to minimize it so this much for today so in in the next lecture probably that is the that is the final lecture in this electro chemistry session we will take up some numerical problems along with some questions we will discuss some questions possible and possible answers ah to make we make you acquainted with the possible you know questions maybe maybe may come in your mind ok so and also possible the miracle problems that you may attempt with attempt with you know attempt while discussing or while while while reading ah some of the books that is available in market

ok

so so

so that will be taken up in the in the next next lecture possibly that is the final

lecture

so till then have good time thank you you

Prutor@IITK