

Production of bio oil and bio char through thepyrolysisroute

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ABSTRACT:

Energy frameworks are perceived as a promising answer for the energy lack and ecological contamination emergencies. To fulfilltheexpandingneedforenergy, differentpotential frameworkshave been researched forthecreationofalternateenergy sourcesareinvestigatedrigorously. The current study explains the pyrolysis process for the generation of bio oil and bio charfrom the pyrolysis process for the pyrolysis pyrolysi pyrolysis pyrolysis pyrolysis pyrolysisocess. The study investigates the effect of the pyrolysis process parameters on the bio oil yield and the bio char yield. Observationsmade by the study reveal that these process parameters significantly affect the bio char yield as well as the bio oil yields. Bio oilyield increased with the rise in the heating time and heating temperature. The reason may the higher temperature or heating ratesuits the chemical reactions which remove off the volatiles preset in the material. Thus the gaseous increases component which

oncoolingbecomeliquidproduct. The biocharyield is found to reduce with the rise in either of the heating time or heating temperature in the current study.

Keywords:Energystorage,Pyrolysis,Thermaltreatment, Correspondingauthor:AvitaAgarwal

I. INTRODUCTION

Orgyisbeingrequestedanddevouredaroundtheworld.Meanwhile,thebroaddepletionofpetroleumderivativeshasmade mankindface genuine energy emergency and ecological disturbances in ongoing many years. Specifically, the radical emanations of ozoneharming substance and other burning poisons have caused non-unimportant outcomes, for example, a dangerous atmosphericdeviation and air contamination, undermining the future improvement of human culture. As per the Worldwide Energy Specialistmeasurements, the all out CO2 emanations from fuel ignition expanded to 32.3 billion metric tons in 2013, a number expected toincrementasthenon-OECD(AssociationforMonetaryCo-activityandAdvancement)non-

industrial nations are expected to burn-

throughmore energy in the forth coming many years. In the present circumstance, investigating substitute energy assets an dpractical energy frameworks has turned into an exceptionally pressing mission confronting all nations as of now [1].

In light of these expressed difficulties, numerous endeavors have been dedicated to new energy framework innovative work, and different elective advancements have been proposed and researched. Hydrogen energy frameworks began to draw in perceptible consideration during the energy emergency of the 1970s and have gone through colossal improvement from that point forwardHydrogen is viewed as an extremely encouraging energy transporter on account of its high level inborn properties, like its highenergy thickness of 14,300 J/(kgK) [2], feasibility, harmless to the ecosystem burning items, and broad long haul assets Despitethefactthathydrogenisthe

mostbountifulcomponentontheplanet, it for themost partexists inwater and hydrocarbons, where it is bound to oxygen and carbon and henceforth not normally prepared to utilize subsequently; various cycles have been created and used to deliver unadulterated hydrogen from different assets. Among these accessible hydrogen creation measures, the steamtransforming response (SRR) of petroleum gas followed by the water gas shift response (WGSR) is perceived as the most prudent cycle and is answerable for more than 90% of the complete hydrogen creation as of now . Meanwhile, other promising advances, including water parting and biomass preparing, are quickly creating a result of their capability to change over spotless and sustainable power assets to hydrogen. The utilization of biomass preparing for hydrogen creation is direly required, not just onaccount of the enormous stockpile of sustainable assets yet additionally as a result of the capacity to save the climate by usingunloaded mining and agrarian squanders. To accomplish an effective clean hydrogen energy society, delivering hydrogen from acarbon obliged technology, like biomass, is just the initial segment of the entire hydrogen energy framework. The division anddecontamination, the capacity and conveyance and the proficient usage of the hydrogen gas delivered are additionally vital stagesin understanding the advantages of utilizing the hydrogen energy produced from manageable assets [3]. Among these means, the division and decontamination establish a basic interaction in the hydrogen energy framework from both the specialized and monetarypoints of view. Presently, different gas detachment advancements are promptly accessible for isolating hydrogen from the blended gases delivered, and film division technology enjoys shown unmistakable benefits of low energy utilization, harmless to the ecosystem properties and the alluring capability filling in asamultifunctional layer reactor [4].

Thinking about hydrogen's extraordinary potential as a perfect energy transporter, the hydrogen energy framework was proposed and perceived as a long-lasting answer for the oil based fuel consumption and ecological emergency. In a hydrogen energyframework, the energies delivered from various essential energy sources (sunlight based, wind, biomass, and so on) are at firstmoved to a hydrogen transporter through an assortment of responses (gasification, water parting and improving, and so on) Theenergy transporter hydrogen is then shipped, put away lastly used by the terminal clients, as displayed in. In this framework, hydrogen fills in as the mediator to move energy between the energy asset and the energy use clients Right now, the hydrogenenergy situation is n't just standing out according to the examination viewpoint, yet additionally a few stater unadditional stater and the state of thministrationshavemade enormous speculations to fabricate the foundation for understanding a hydrogen energydriven society. One astoundingoccasion is that the public authority of Iceland has defined the objective of hydrogen-economy country theextended of2030 turning into а total in time byutilizingtheirgeothermalenergythroughhydrogenenergyframeworks[5].

Furthermore, the advancement of hydrogen-

fueledvehiclesisempoweredbynumerousareas,likeJapanandChina.Notwithstanding, there are four significant specialized provokes that should be looked before a proficient hydrogen energyframeworkcanbeacknowledged,includinghydrogencreationanddecontaminationthroughacarbon-

compelled measure, hydrogen conveyance and dissemination through wide-running and very much oversaw foundation, hydrogen stockpiling in solid and safeplants and hydrogen use through a productive interaction.

Gasification is a two-venture measure in which a strong fuel (biomass or coal) is thermochemically changed over to a low-ormedium-energy-content gas. Flammable gas contains 35 MJ/m³. Air-blown biomass gasification brings about roughly 5 MJ/m³;oxygen-blown in 15 MJ/m³. In the principal response, pyrolysis, the separated and unstable parts of the fuel is disintegrated attemperatures as low as 600 °C. Remembered for the unpredictable fumes are hydrocarbon gases, hydrogen, carbon monoxide,carbon dioxide, tar, and water fume. Since biomass powers will in general have more unstable parts (70-86 % on a dry premise)than coal (30%) 6,7], pyrolysis assumes a bigger part in biomass gasification than in coal gasification. Gas stage warm breaking ofthevolatileshappens,lesseningthedegreesoftar.Burn(fixedcarbon)anddebrisarethepyrolysisresultsthatarenotdisin tegrated.In the subsequent advance, the burn is gasified through responses with oxygen, steam, and hydrogen. A portion of the unburnedscorch mightbe combusted to deliverthehotnessrequiredfortheendothermic pyrolysisresponses.

Gasification combined with water-gas shift is the most broadly polished cycle course for biomass to hydrogen. Warm, steam andfractional oxidation gasification innovations are a work in progress all throughout the planet. Feed stocks incorporate both devotedharvestsandruralandbackwoodsitembuildupsofhardwood, softwoodand herbaceousspecies.

Warm gasification is basically high seriousness pyrolysis in spite of the fact that steam is by and large present. An illustration of this is the Sylvagas (BCL/FERCO) low-pressure, in a round about way warmed circling liquid bed[8]. $Biomass+Energy \rightarrow CO + H_2 + CH_4 + \cdots$

Other relevant gasifier types are bubbling fluid beds being tested by Enerkem, and the highpressure high-temperature slurry-fedentrainedflowTexacogasifier.

All of these gasifier examples will need to include significant gas conditioning, including the removal of tars and inorganic iimpurities and the conversion of COtoH2by the water-gasshift reaction

 $C0{+}H20{\rightarrow}C02{+}H2$

Significant attention has been given to the conversion of wet feedstocks by high-pressure aqueous systems. This includes the supercritical-gasification-in-water approach by Antal and coworkers as well as the supercritical partial oxidation approach by GeneralAtomics.

Pyrolysistohydrogenandcarbonisbeinginvestigatedasapracticaltechnologyforcarbonsequestrationalbeitmostworki sappliedto flammable gas pyrolysis. Biomass or biomassderived intermediates could be handled along these lines. Organic transformationthrough anaerobic assimilation is presently being rehearsed by laborers all throughout the planet bringing about methane that canbehandledtohydrogenbyregularsteamimprovingcycles[9].

1.2StorableIntermediates

Pyrolysis creates a fluid item called bio-oil, which is the premise of a few cycles for the advancement of fills, synthetic substances and materials. The response isendothermic:

$Biomass+Energy \rightarrow Bio-oil+Char+Gas$

The oil can be framed in 66 wt.% yields. Chornet and collaborators have fostered the idea of utilizing the

lingering parts of the oilforhydrogenafterco-itemshavebeendelivered.SynergiststeamimprovingofBio-oilat750-850°Coveranickel-basedimpetusisatwo-venturemeasure that incorporates he shiftresponse:

 $Bio-oil+H_2O\rightarrow CO+H_2$

 $CO+H_2O\rightarrow CO_2+H_2$

The overallstoichiometry gives a maximum yield of 1 7.2 gH/1 00 gbio-oil(11 .2% based on wood). $CH_{1.9}O_{0.7}+26H_2O \rightarrow CO_2+2.21H_2$

Local organizations of pyrolysis plants could be established to give oil to a focal steam improving office. The interaction is viable with other natural was test reams, for example, aquous-steam fractionation measures utilized for ethanol creation and trapoil.

Methanolandethanolcanlikewisebedeliveredfrom biomassbyanassortmentofinnovationsandutilizedforonboardtransforming for transportation. Methane from anaerobic assimilation could be changed alongside gaseous petrol. Methane

couldbepyrolyzedtohydrogenandcarbonifmarketsforcarbondarkwereaccessible.Frameworksinvestigationhassho wnthatbiomassgasification I shift change is financially troublesome contrasted with gaseous petrol steam improving aside from

exceptionallyminimalexpensebiomassandexpected natural motivations. The pyrolysis with significant coitem approachy ields hydrogen in the values cope of \$6-\$8/GJ, which is promising for close to term applications 10].

II. EXPERIMENTAL WORK

The experiments we reconducted to develop the energy rich products from the wastematerials and convert these into electrodes and check their performance.

Biomass derived char was synthesized using the pyrolysis of the biomass waste. The selected biomass was fed to a muffle

furnaceandthenthechamberwasfilledwithnitrogengas. Thenitrogengasflowratewascontrolledandfromoutsideanditw asdeterminedwith the help of a nitrogen flow meter. Once the whole reactor chamber was filled with the nitrogen gas, the heater was switchedON, and the heater was programmed to reach a t a definite temperature at a fixed heating rate. Once everything was set, biomassstarted to heat. The heating of the biomass was carried out at various temperature, and time combinations. The Table 1 shows thevariouscombinationsoftimeandtemperaturewhich were used to synthesize the char.

RunNo.	Timeofheating(min)	Temperature(⁰ C)
1	10	650
2	20	650
3	30	650
4	40	650
5	50	650
6	60	650
7	10	700
8	20	700
9	30	700
10	40	700
11	50	700
12	60	700
13	10	750
14	20	750
15	30	750
16	40	750
17	50	750
18	60	750
19	10	800
20	20	800
21	30	800
22	40	800
23	50	800
24	60	800
25	10	850
26	20	850

27	30	850
28	40	850
29	50	850
30	60	850
31	10	900
32	20	900
33	30	900
34	40	900
35	50	900
36	60	900

III. RESULTS AND DISCUSSIONS

Effectoftemperatureonpyrolysisproductyield

It is well known fact that the pyrolysis products yield depends upon the pyrolysis parameters. The experiments performed veryclearly confirmed this. It is seen clearly that with the rise in the temperature, the product yield is significantly. changing The yield of each product depends up on the temperature. As the temperature increases, the bio oil yield starts increasing and the contract of the temperature increases and the contract of temperature increases and temperaharyield goes down. The reason for this behavior may be understood from the fact that as the temperature increases up, the reaction within the materials increases and the thermal cracking of various types of bonds present in the biomass starts. The thermal cracking of these bonds least othe breaking of bonds and thus the volatile material p[resent in the biomass starts to convert themselves and the start start starts the start start start start start starts and the start starinto the gas eous form. These gasses liberates from the materials and these starts to get away from the materials. The gas the material starts the starts to get a start of the starts thcomesoutandis collected into the condenser. Clearly as the temperature will increase more and more volatiles will be released from within thebiomassmaterial andthusthebiooilyieldwillbeenhanced.

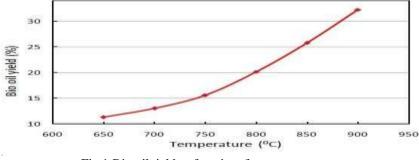
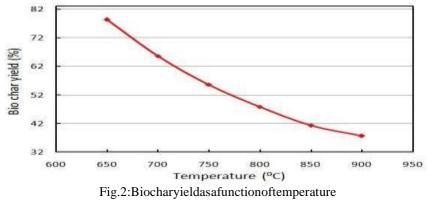


Fig.1:Bio oilyieldasafunctionoftemperature

However, when the bio oil yield will go up, clearly it will occur at the cost of the char yield. As more and more volatiles will comeout of the char clearly it will reduce the char yield. The similar observations can be made from the results obtained from the experiments. A graph plotted between bio oil yields vs. the temperature is shown in Fig. 1. It shows that the rise in the temperaturehas increased the bio oil yield significantly. The reason of such a behavior is already explained. Similarly, the char yield vs.temperature is shown in Fig. 2. The Fig. 2 shows the expected behavior, where it is seen that as the temperature rise, the char yieldgoesdown.Thereasonbeingthattheriseintemperaturemayleadtotheevaporationofvolatilesandthusthebiocharyi eldlowers.



${\it Effect of reaction time on pyrolysis product yield}$

The effect of the heating time on the bio oil yield was also investigated in the present study. The results obtained from

theexperiments are shown in the graphical form in Fig. 3. This Fig. 3 shows the variation of the bio oil yield as a function of the heating time. It is evident from the fig. 3 that the rise in the heating time enhances the bio oil yield. The enhanced bio oil yield with rise intemperature is seen because longer heating time provides more time for the chemical reaction to take place and as the results of the sechemical reactions, the volatile material present in the bio mass converts its elfint othe gas and slowly it comes out. The more

the heating time the more will be the reaction time which further will liberate more volatiles which eventually will produce moregasses. These gasses will finally be condensed toget the biooil. The study showed that the biooily ield increased from 3.12% to

27.23% on increasing the heating time from 10 minto 60 min. This is almost 89.5 times rise in the bio oily ield which is signific antly high. Thus it is seen that the heating time affects the bio oil production up to a large extent.

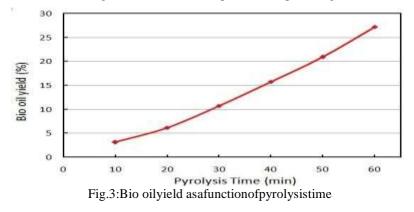
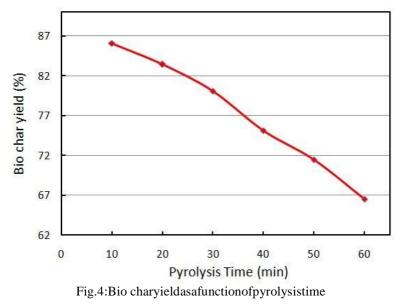


Fig.4isarepresentationofthebiocharyieldasafunctionoftheheatingtime.Thebiocharyieldinthepresentexperimentisse ento be lowered significantly with the rise in the heating time. It is observed that with the rise in the heating time from 10 min to 60min, the bio char yield is reduced from 86.62 % to 66.55 %. This is a reduction of almost 25 % in the bio char yield. This loweringin the bio char yield is seen as the more heating time provides sufficient time for the material under investigation to perform thechemical reactions. The resultant of these chemical reactions is the evaporation of the volatiles resent in the material. Thus theincrementinthe heatingtime reduces the biocharyield.



IV. CONCLUSIONS

Thestudydiscussed the impact of the pyrolysis process parameters on the bio oil yield and the bio charyield. Two different process parameters viz. heating temperature and heating time were chosen for this study. The study found that both of these parameters significantly affect the pyrolysis product yields. However, the effect of these process

parameters is different for the different pyrolysis products. Whereas, bio oil production enhances on increasing the heating time or the heating temperature, the hio charyield lowers with the rise in either of the pyrolysis process parameter. The increase in the bio oilyield may be attributed the pyrolysis process parameter of the pyrol pyrolothebetterchemicalreactionatlongerheatingtimeandheatingtemperatures.Improvedchemicalreactions mayleadtotheevaporationofthevolatile present in the material. These volatiles can further be condensed and the bio oil extracted. Thus can be higher temperature or heating time produces more gasses and more bio oil is extracted. On the other handwith the longer heating rate of the temperature of tempeorhigherheatingtemperatures, the charyield lowers as the volatileses capes away from the soild material reducing the bio charyield.

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