Laboratory Gasification Memo
Partial Pressure of Steam and Residence Time Experiments

Summary

An experimental campaign was completed to explore the suspected dependence of carbon conversion on the partial pressure of steam as well as the residence time. Partial pressure of CO_2 was held constant at 7 psi, and total pressure was set to 50 psig. It was found that the partial pressure of steam had an effect on both carbon yield and carbon release at 1350 °C and 1450 °C but did not have a statistically significant effect on methane or tar yields at either temperature. The maximum residence time had an effect on carbon conversions only at 1450 °C, and shorter residence times lead to higher conversions in those cases. Possible explanations and recommendations for further analysis and experimentation are given in this memo.
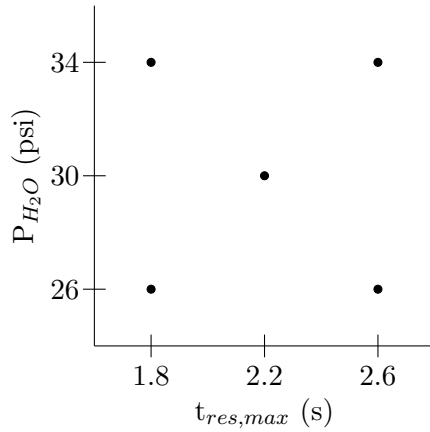
Figure 1: Two factorial experimental matrix used to vary maximum residence time and partial pressure of water.

Experimental Methods

Because of the difficulty in manually designing an experimental matrix in which the $\Delta H_{max}/A$ is explicitly controlled as a factor, a large number of potential inlet conditions were simulated using Sundrop Fuel's analysis software. Argon flow rate was set to 2 SLPM, and total pressure was 50 psig. The biomass mass flow rate was randomly chosen using an even distribution between 2 and 4 lbs/hr. Steam flow rate and temperatures were chosen in the same manner between 12 and 24 mL/min and 300 and 500 °C, respectively. A makeup flow rate of nitrogen was set to be between 0 and 20 SLPM. The total flow rate of entrainment gas was set such that there were

6 SLPM for every lb/hr of biomass flow.

Once the total inlet molar flow rate was known for the potential run, the portion of the entrainment gas which was CO_2 was set using Equation 1 such that the partial pressure of CO_2 was 7 psi. The remainder of the entrainment gas flow rate was made to be nitrogen.

$$\dot{n}_{CO_2} = \dot{n}_{tot} \frac{P_{CO_2}}{P_{tot}} \qquad (1)$$

Temperatures for the outer reactor wall were set to be 1350 °C and 1450 °C for two separate experimental matrices. The inner diameter of the tube was 1.5". The maximum residence time, or space time, was calculated assuming inlet conditions using Equation 2. Once the experiment had been completed, the minimum residence time was calculated using outlet molar flow rates and assuming the product gases immediately reached the outer wall temperature of middle of the SiC tube. Equation 3 calculates the minimum residence time.

$$t_{max} = \frac{VP}{\sum_{i \neq biomass} \dot{n}_i RT_{mix}} \qquad (2)$$

$$t_{min} = \frac{VP}{(\sum_{i \neq H_2O} \dot{n}_i + \dot{n}_{H_2O,0})RT_{wall}} \qquad (3)$$

Before the experiment took place, the heat duty ($\Delta H_{max}/A$) was calculated with Equation 4 assuming complete conversion would take place and the products reached the reactor outer wall

temperature. lPotential experiments were filtered such that the $\Delta H_{max}/A$ was 60 $\pm 2\%$ kW m^{-2}. Based on previous observations, any variations in experiments caused by changes in heat duty within this range would not be detected with the current setup. More runs were filtered out to ensure that the adiabatic mixing temperature of the reactants would not get near the condensation temperature of steam at 50 psig (about 148 °C). Finally, points were chosen to match the desired targets for space time and total inlet steam partial pressure shown in Figure 1. All experimental set points can be found in Appendix 9.

$$\Delta H_{max} = \sum_{products} n_i \left[\Delta H_i^\circ + \int_{T^\circ}^{T_{out}} C_{p,i} dT \right] \quad (4)$$

$$- \sum_{reactants} n_i \left[\Delta H_i^\circ + \int_{T^\circ}^{T_{in,i}} C_{p,i} dT \right]$$

Two measures of carbon conversion are discussed in this memo. The first is the fraction of carbon in the biomass which is converted to either CO or CO_2, as these are the two species which are the precursor to synthetic liquid products in the planned commercial process. This measure is referred to as carbon yield, although it has been referred to in the past as good conversion, and is given in Equation 5.

$$Y_{CO+CO_2} = \frac{\dot{n}_{CO,out} + \dot{n}_{CO_2,out} - \dot{n}_{CO_2,in}}{\dot{n}_{C_{biomass},in}} \quad (5)$$

The second measure is carbon release, which has been referred to in the past as total conversion. This was calculated using Equation 6 and is a representation of the fraction of carbon in the biomass which is converted to any gaseous species detected by the mass spectrometer.

$$X_C = \frac{\dot{n}_{C_{gas},out} - \dot{n}_{CO_2,in}}{\dot{n}_{C_{biomass},in}} \quad (6)$$

Tar loading is a measure of the mass of tars detected by the mass spectrometer (C_6H_6, C_7H_8, and $C_{10}H_8$) in a standard volume of product gas. This value was calculated using Equation 7.

Table 1: ANOVA results on effects of designed experimental campaign for carbon yield.

Effect	Prob $<$F	
	1350 °C	1450 °C
$t_{res,max}$	0.1974	0.0057
P_{H_2O}	0.0039	0.0019
$t_{res,max} \times P_{H_2O}$	0.0812	0.7289

$$C_{tar} = \frac{\dot{m}_{C_6H_6} + \dot{m}_{C_7H_8} + \dot{m}_{C_{10}H_8}}{\dot{V}(\frac{P}{P_{std}})(\frac{T_{std}}{T})} \quad (7)$$

Finally, the last measure discussed in this memo is methane yield. It is a representation of the fraction of carbon in the biomass which is converted to methane, and it was calculated using Equation 8. A table defining all variables used in this memo can be found in Appendix G.

$$Y_{CH_4} = \frac{\dot{n}_{CH_4}}{\dot{n}_{C_{biomass},in}} \quad (8)$$

Results and Discussion

Carbon Yield

ANOVA results for carbon yield, given in Tables 1 and 2, showed that there was an effect of partial pressure of steam on conversion of biomass carbon to CO and CO_2 at both 1350 and 1450 °C. There were effects of both the minimum and maximum residence times on the carbon yield at 1450 °C, but neither measure had an effect at 1350 °C. At both temperatures, a shorter residence time led to higher carbon yields, as shown in the plots in Appendix A. At 1350 °C, there seemed to be an interaction effect between the minimum residence time and the partial pressure of steam on the carbon yield.

It's interesting to note that the results for both residence times mirrored what was seen in a previous experimental campaign where maximum residence time and $\Delta H_{max}/A$ were explicitly controlled as factors. The results sparked interest previously, because general kinetic insight would lead one to predict that longer space times should lead to better conversions, not vice versa. The

Table 2: ANOVA results on effects of minimum residence time and partial pressure of water on carbon yield.

Effect	Prob <F	
	1350 °C	1450 °C
$t_{res,min}$	0.0694	0.0001
P_{H_2O}	0.0060	<0.0001
$t_{res,min} \times P_{H_2O}$	0.0400	0.2459

fact that the same results were observed in a different experimental matrix hints at the possibility that there was another factor effecting the carbon yields that was correlated with the residence times.

One possible factor that would be very tightly correlated with the residence time is the gas velocity in the reactor. Simplifying the flow pattern in the reactor and assuming plug flow, the minimum and maximum velocities are functions of the reactor length together with the maximum and minimum residence times, respectively. These relationships are shown in Equations 9 and 10. The minimum velocity would be closest to what is seen at the entrance of the reactor, and the maximum velocity would be nearest to what may be occurring at the outlet of the reactor.

$$u_{min} = \frac{l}{t_{res,max}} \qquad (9)$$

$$u_{max} = \frac{l}{t_{res,min}} \qquad (10)$$

If the velocity of the gas was affecting the conversion of carbon in the biomass to CO and CO_2, it could have been due to heat transfer effects; higher gas velocities would lead to higher convective heat transfer coefficients. The effects may be apparent at 1450 °C and not 1350 °C, because the driving force for convective heat transfer is higher when the ΔT between the reactants and reactor wall are higher. These results may be further evidence that the overall heat transfer rate between the tube wall and the reactants is a more important driving force in carbon conversion than the residence time. This conclusion could be further supported through estimating

Table 3: ANOVA results on effects of designed experimental campaign for carbon release.

Effect	Prob <F	
	1350 °C	1450 °C
$t_{res,max}$	0.0804	0.0039
P_{H_2O}	0.0256	0.0051
$t_{res,max} \times P_{H_2O}$	0.8947	0.7544

Table 4: ANOVA results on effects of minimum residence time and partial pressure of water on carbon release.

Effect	Prob <F	
	1350 °C	1450 °C
$t_{res,min}$	0.0363	<0.0001
P_{H_2O}	0.0388	0.0002
$t_{res,min} \times P_{H_2O}$	0.4340	0.1932

radiative and convective heat transfer coefficients for completed gasification experiments with computer models and seeing the effect the coefficients have on carbon yields.

Carbon Release

ANOVA results for the effects of residence times and partial pressure of steam on total carbon release are given in Tables 3 and 4. Similar to carbon yield, carbon release had a dependence on partial pressure of water at both 1350 and 1450 °C. Both maximum and minimum residence time affected the carbon yield at 1450 °C. However, unlike carbon yield, carbon release showed a statistically significant dependence on the minimum residence time at 1350 °C that was not present for the maximum residence time. Plots in Appendix B show results for the experiments.

This apparent dependence of the total carbon release on the minimum residence time, but not maximum residence time, could be a result of the fact that the minimum residence time is calculated using conditions at the exit of the reactor. Higher conversions would mean more gas was released from the solids, leading to higher volumetric flow rates and higher velocities at the

Table 5: ANOVA results on effects of designed experimental campaign for tar loading.

Effect	Prob <F	
	1350 °C	1450 °C
$t_{res,max}$	0.8412	0.4901
P_{H_2O}	0.4180	0.5533
$t_{res,max} \times P_{H_2O}$	0.2223	0.4228

Table 6: ANOVA results on effects of minimum residence time and partial pressure of water on tar loading.

Effect	Prob <F	
	1350 °C	1450 °C
$t_{res,min}$	0.7755	0.3675
P_{H_2O}	0.4392	0.4980
$t_{res,min} \times P_{H_2O}$	0.3548	0.4106

Table 7: ANOVA results on effects of designed experimental campaign for methane yield.

Effect	Prob <F	
	1350 °C	1450 °C
$t_{res,max}$	0.0573	0.7161
P_{H_2O}	0.0678	0.3773
$t_{res,max} \times P_{H_2O}$	0.1209	0.5752

exit of the reactor. Although, if there was an actual physical effect, it could give interesting insight into the importance of different heat transfer modes in different parts of the reactor.

It is believed, and has been supported with computer models, that there is a zone at the entrance of the reactor which resembles a continuous stirred tank reactor. This would mean this zone has very turbulent gas recirculation patterns and higher convective heat transfer coefficients. The minimum residence time is probably more closely correlated to the average gas velocity throughout the rest of the reactor, because the bulk of the gas release that would increase the velocity likely takes place at low temperatures near the entrance of the reactor. This could contribute to the face that the minimum residence time was found to be a more statistically significant factor in carbon conversion. Also, the fact that the minimum residence time was much more statistically significant at 1450 °C could be due to the fact that higher conversions at this temperature led to much lower amounts of solids at the bottom of the reactor and increased the importance of convective heat transfer to the gas from the tube wall in this area.

Tar Loading

ANOVA results, shown in Tables 5 and 6, did not show any effects from minimum residence time, maximum residence time, or the partial pressure of steam on the production of tars. Results are plotted in Appendix C.

The results differed slightly with what was seen in a previously completed experimental matrix changing $\Delta H_{max}/A$ and maximum space time. These experiments showed an effect of residence time on the tar loading at 1350 °C such that longer space times led to lower tar levels.

Methane Yield

Similarly to tar loading, methane yield did not show any dependencies on the factors tested in these experiments. Results from ANOVA are in Tables 7 and 8, and they are plotted in Appendix D. Again, these results differed from the previously completed experimental matrix where residence time had an effect on methane production at both temperatures, and longer residence times led to less methane. However, the partial pressure of water was nearly statistically significant at 1350 °C, and it's possible that a wider range steam pressures could reveal effects on the production of methane.

Conclusion

The experimental campaign showed the effects of residence time and partial pressure of steam on the gasification of biomass. Partial pressure of steam had an effect on both carbon yield and carbon release at 1350 and 1450 °C. Maximum residence time only had an effect on carbon yield

Table 8: ANOVA results on effects of minimum residence time and partial pressure of water on methane yield.

Effect	Prob $<$F	
	1350 °C	1450 °C
$t_{res,min}$	0.0482	0.6862
P_{H_2O}	0.1054	0.3490
$t_{res,min} \times P_{H_2O}$	0.3255	0.7152

and carbon release at 1450 °C. However, minimum residence time had an effect on the total carbon release at both temperatures. Neither residence time or partial pressure of steam had an effect on methane or tar yields in this set of experiments.

Further analysis and experiments can shed some light on the physical effects behind the results. Better understanding what the convective heat transfer in different areas within the reactor could shed some light on why it appears there are decreasing conversions with increasing residence times.
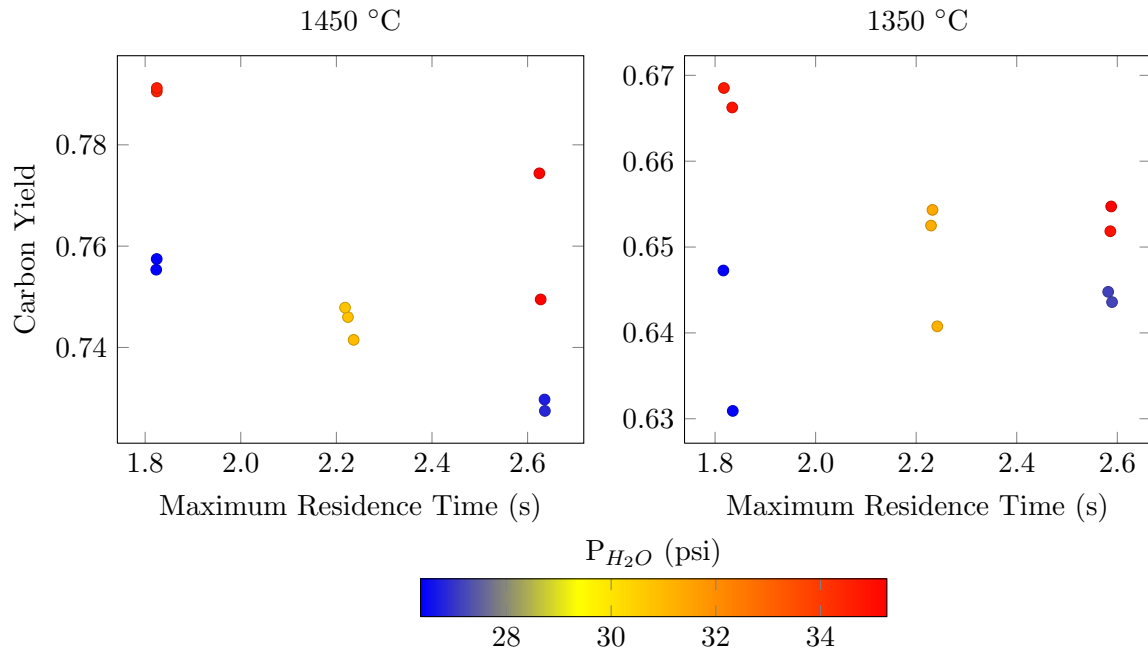
A   Carbon Yield Plots

Figure 2: Carbon yield plotted against maximum residence time and colored by the partial pressure of water. At 1450 °C, there was a dependence on both maximum residence time and partial pressure of steam. However, at 1350 °C, there was only a dependence on partial pressure of steam. These observations are backed up by ANOVA results.
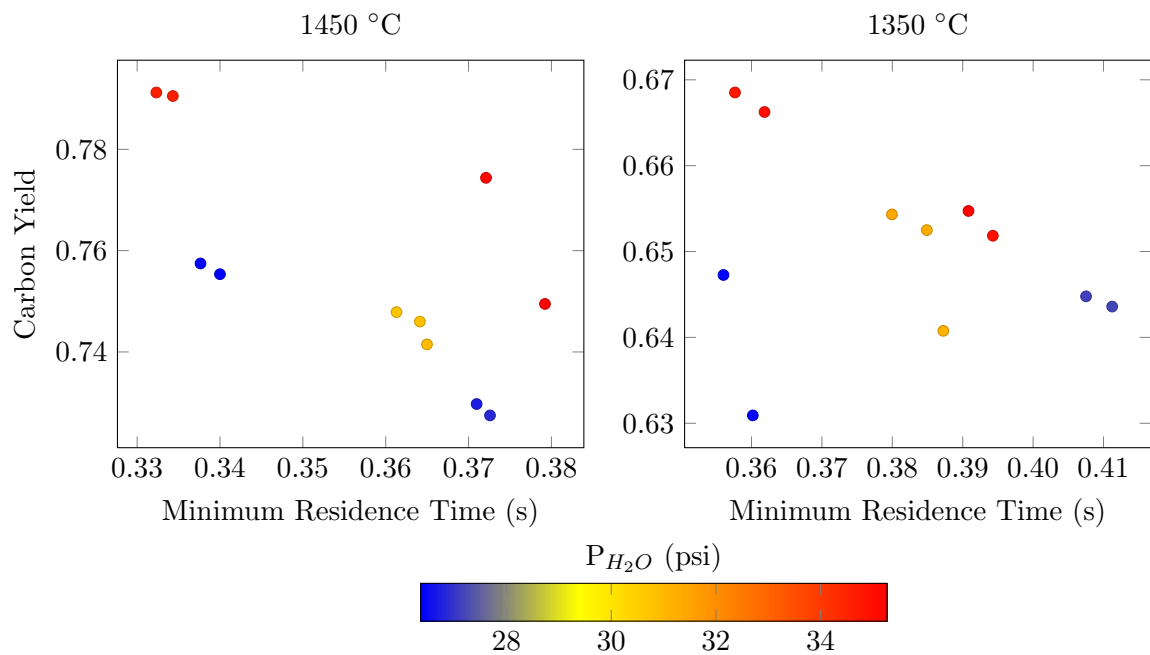

Figure 3: Carbon yield plotted against minimum residence time and colored by partial pressure of steam. Similar results were observed when looking at minimum residence time rather than maximum residence time. ANOVA results showed an interaction effect between the minimum residence time and partial pressure of steam at 1350 °C.
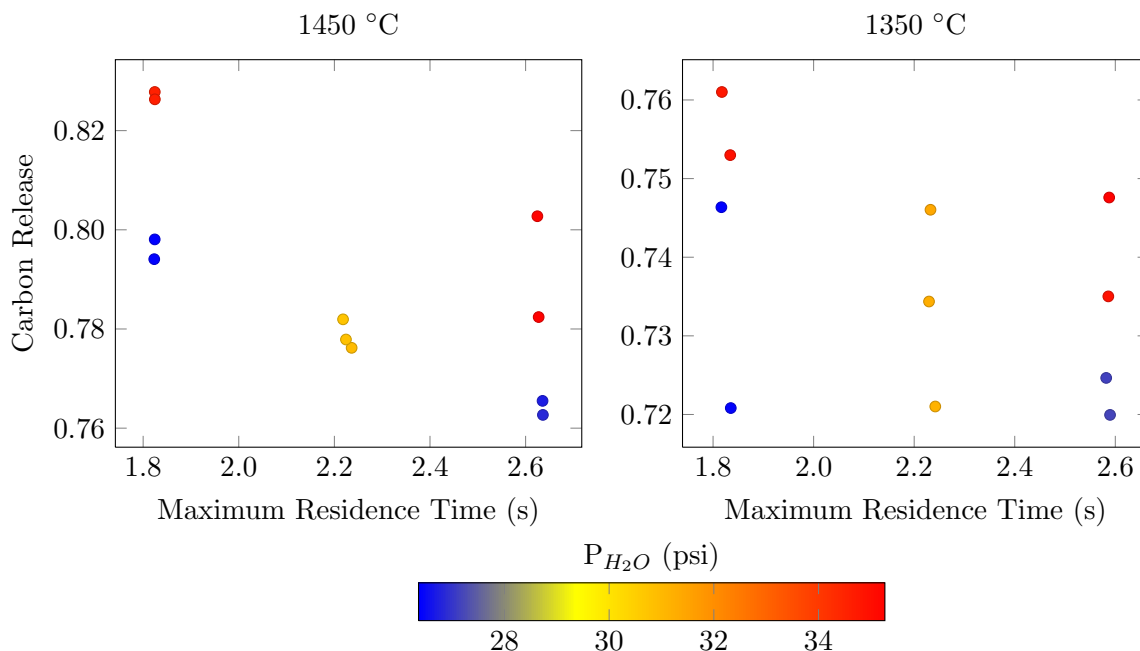
B Carbon Release Plots

Figure 4: Total carbon release from biomass plotted against maximum residence time, colored by the partial pressure of water. Similar to carbon yield, carbon release showed dependence on both partial pressure of steam and maximum residence time only at 1450 °C. ANOVA results showed a dependence on only partial pressure of steam at 1350 °C.
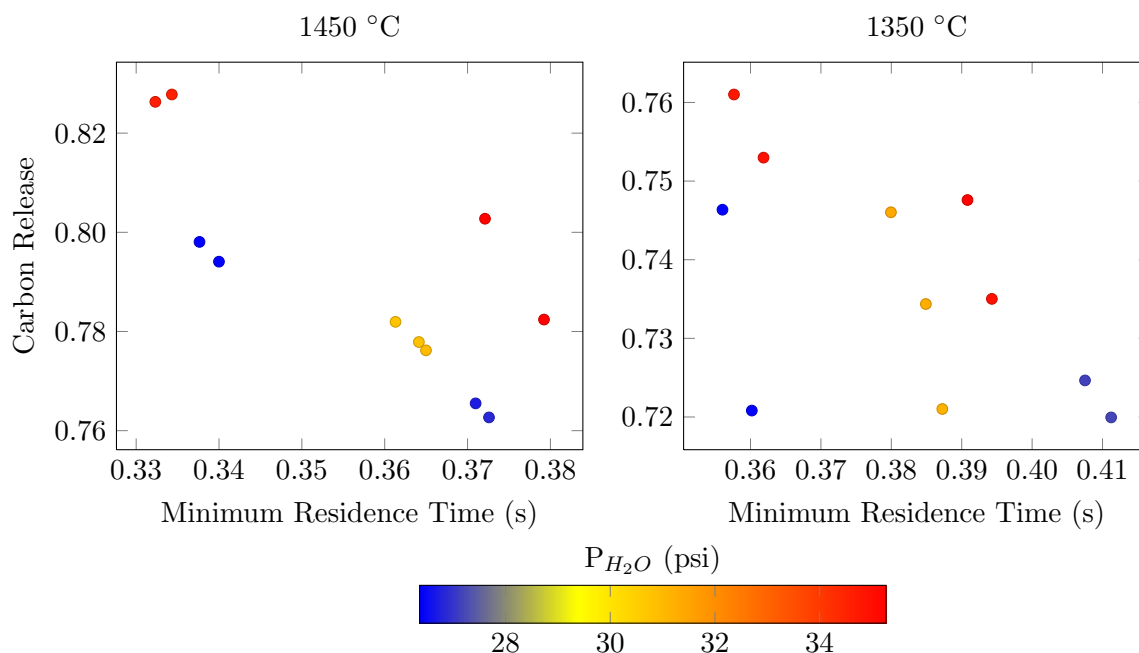

Figure 5: Carbon release vs. minimum residence time and colored according to partial pressure of steam. Unlike the maximum residence time, minimum residence time was found to be an important factor in carbon release at both 1350 °C and 1450 °C according to ANOVA. Partial pressure of steam also showed an effect on carbon release at both temperatures.
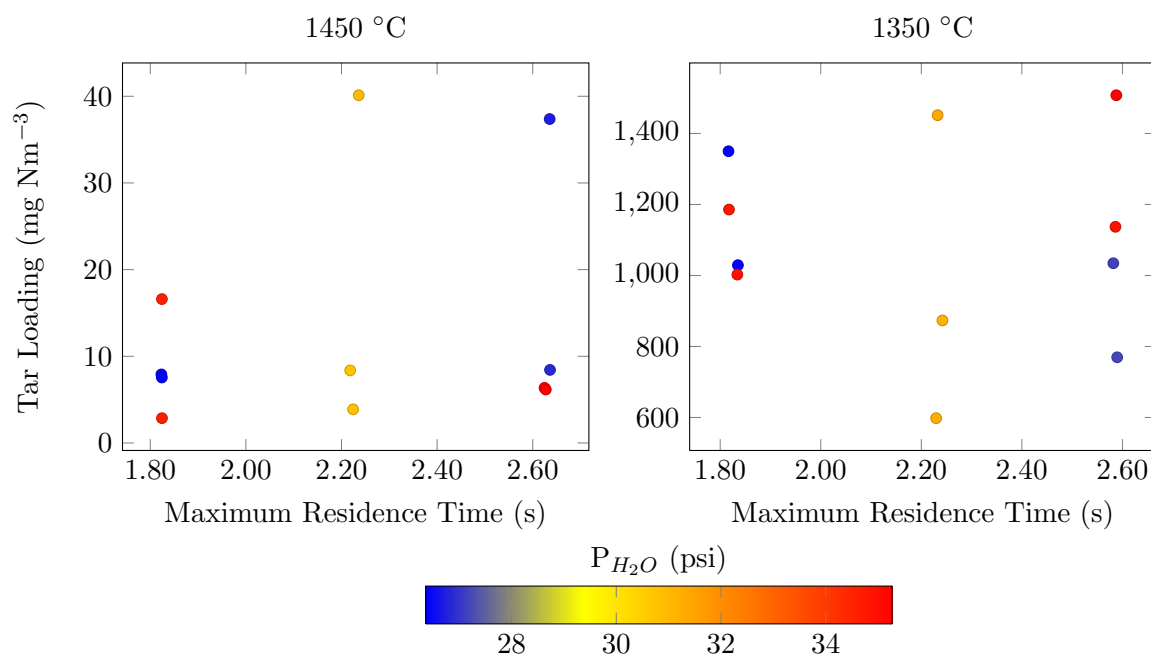
C Tar Loading Plots

Figure 6: Tar loading vs. maximum residence time, colored by the partial pressure of steam. ANOVA showed that neither partial pressure of steam or maximum residence time had an effect on the tar loading at either 1350 °C or 1450 °C.
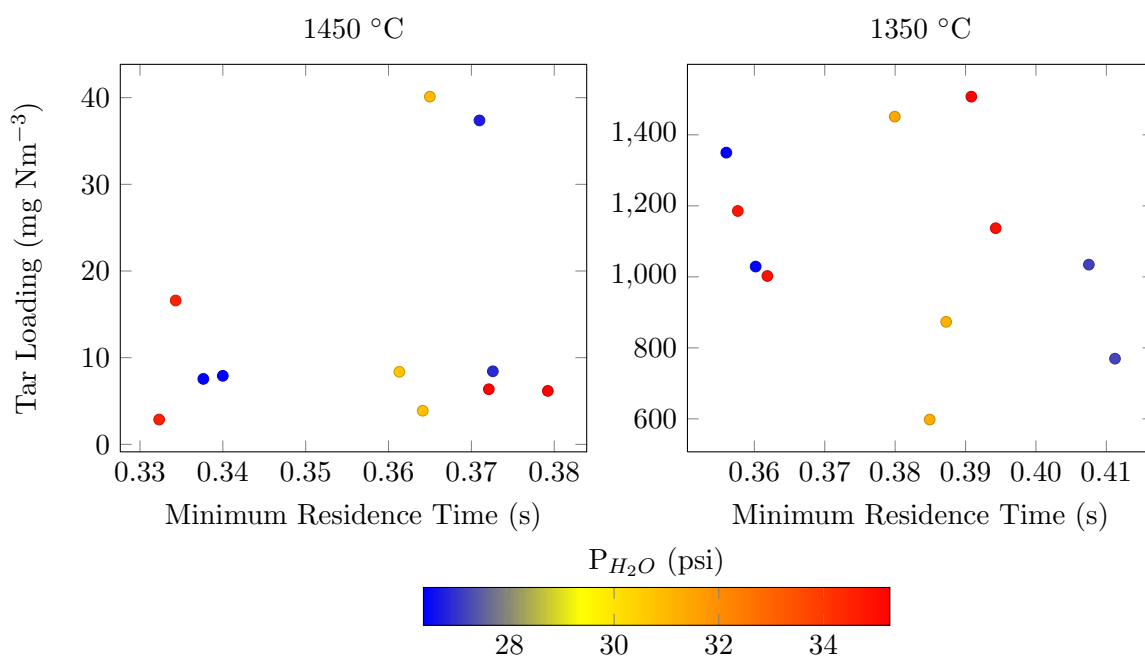

Figure 7: Tar loading plotted against minimum residence time and colored against partial pressure of steam. Again, there were no statistically significant effects found from ANOVA results at either temperature.

D   Methane Yield Plots

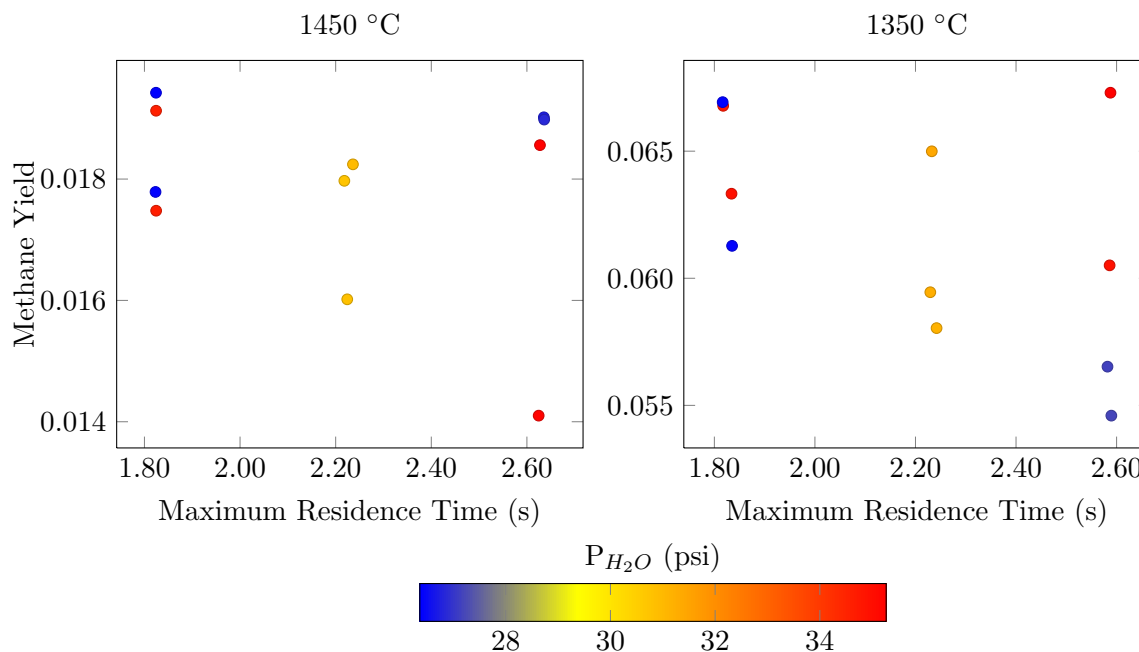Figure 8: Methane yield vs. maximum residence time, colored by partial pressure of water. ANOVA results showed no statistically significant dependence of methane yeild on either maximum residence time or partial pressure of steam.
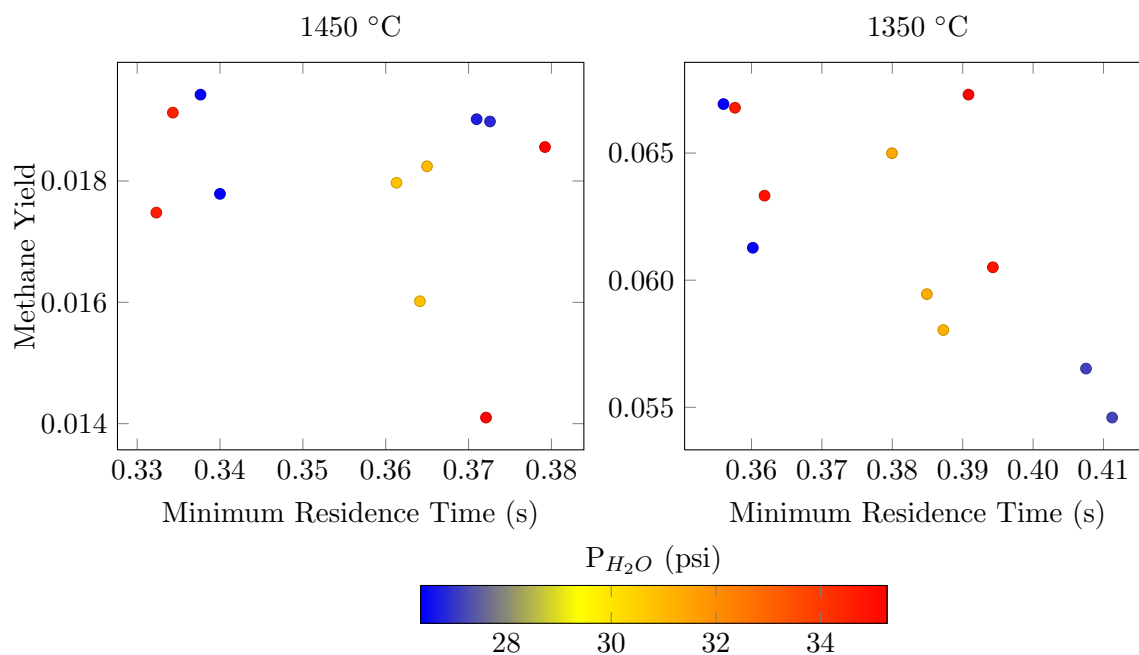

Figure 9: Methane yield plotted against minimum residence time and colored according to partial pressure of steam. The only significant effect found through ANOVA was minimum residence time at 1350 °C.

E Experimental Setpoints

Table 9: Setpoints for partial pressure of steam vs. gas velocity experiments. Pressure is 50 psig, and argon flow is 2 SLPM for all runs. Partial pressure of CO_2 is 7 psi, and $\Delta H_{Max}/A$ is .

Target P_{H_2O} (psi)	Target $t_{res,min}$ (s)	Temp. (°C)	Biomass (lb/hr)	Ent. N_2 (SLPM)	Ent. CO_2 (SLPM)	Makeup N_2 (SLPM)	Steam (g/min)	Steam (°C)
26	2.6	1450	3.2	14.4	4.7	5.1	12.9	433
26	1.8	1450	2.4	8.1	6.5	19.7	17.9	483
30	2.2	1450	3.0	12.9	5.0	5.0	16.0	486
34	2.6	1450	3.0	13.4	4.7	0	17.0	354
34	1.8	1450	2.5	8.4	6.3	10.9	22.9	411
26	2.6	1350	3.0	11.8	6.0	6.7	21.5	463
26	1.8	1350	2.7	9.6	6.7	18.9	18.5	449
30	2.2	1350	3.1	13.4	5.4	6.0	17.4	404
34	2.6	1350	3.2	14.1	5.0	1.2	18.0	306
34	1.8	1350	3.4	16.1	4.6	2.6	12.5	490

F Experimental Results

Table 10: Selected results from the experimental campaign.

Run ID	Temp. (°C)	Max Res Time (s)	Min Res Time (s)	P_{H_2O} (psi)	$\Delta H_{max}/A$ (kW m^{-2})	Carbon Yield	Carbon Release	Tar Loading (mg Nm^{-3})	CH_4 Yield
561	1450	2.24	0.365	30.9	58.5	0.741	0.776	40.1	0.0182
562	1450	1.82	0.334	34.5	58.4	0.791	0.828	16.6	0.0191
563	1450	2.64	0.371	26.7	60.1	0.730	0.766	37.4	0.0190
564	1450	1.82	0.340	26.4	57.4	0.755	0.794	7.91	0.0178
565	1450	2.62	0.372	35.2	57.8	0.774	0.803	6.36	0.0141
566	1450	2.64	0.373	26.8	59.6	0.727	0.763	8.43	0.0190
567	1450	2.22	0.364	30.8	58.3	0.746	0.778	3.88	0.0160
568	1450	1.82	0.332	34.6	58.4	0.791	0.826	2.86	0.0175
569	1450	2.63	0.379	35.3	58.0	0.749	0.782	6.17	0.0186
570	1450	1.82	0.338	26.4	57.8	0.757	0.798	7.55	0.0194
571	1450	2.22	0.361	30.7	58.6	0.748	0.782	8.37	0.0180
572	1350	2.23	0.385	31.1	58.7	0.652	0.734	598	0.0595
573	1350	1.82	0.358	34.4	59.4	0.669	0.761	1190	0.0668
574	1350	2.59	0.411	27.4	58.5	0.644	0.720	769	0.0546
575	1350	2.59	0.391	34.8	59.3	0.655	0.748	1510	0.0673
576	1350	1.82	0.356	26.7	57.9	0.647	0.746	1350	0.0669
577	1350	2.23	0.380	31.3	59.3	0.654	0.746	1450	0.0650
578	1350	1.84	0.360	26.7	58.6	0.631	0.721	1030	0.0613
579	1350	2.59	0.394	34.5	59.0	0.652	0.735	1140	0.0605
580	1350	1.83	0.362	34.5	59.2	0.666	0.753	1000	0.0633
581	1350	2.58	0.408	27.4	58.8	0.645	0.725	1030	0.0565
582	1350	2.24	0.387	31.0	58.8	0.641	0.721	873	0.0580

G Variable Legend

Table 11

Variable	Definition
A	Surface area of the inside surface of the reactor tube
C_{tar}	Tar loading in the product gas in mg Nm^{-3}
ΔH_{max}	Maximum enthalpy change of the reactants assuming complete conversion
$\dot{m}_{i,out}$	Mass flow rate of species i flowing out of the reactor
$\dot{n}_{C_{biomass},in}$	Molar flow rate of carbon in the biomass flowing into the reactor
$\dot{n}_{C_{gas},out}$	Molar flow rate of carbon in all gaseous species flowing out of the reactor
$\dot{n}_{i,in}$	Molar flow rate of gaseous species i into the reactor
$\dot{n}_{i,out}$	Molar flow rate of gaseous species i out of the reactor
P	Pressure
Pi	Partial pressure of species i
P_{std}	Standard pressure
T	Temperature
T_{mix}	Adiabatic mixing temperature of all reactants, excluding biomass, at the reactor inlet
T_{std}	Standard temperature
$t_{res,max}$	Maximum residence time, or space time
$t_{res,min}$	Minimum possible residence time
$\dot{V}$	Volumetric flow rate
X_C	Carbon release from carbon in the biomass
Y_{CH_4}	Methane yield from carbon in the biomass
Y_{CO+CO_2}	Carbon yield to CO and CO_2 from carbon in the biomass