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A Comparative Study of Diffusion Coefficients from Convective and IR Drying of Woodchip

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Abstract - Convective and infared (IR) Halogen Drying processes are used in the woodchip biomass industry to test the moisture content of woodchip. Woodchip drying, though energy intensive, is necessary to increase the calorific content of woodchip, in turn increasing combustibility. The bulk process within the production of woodchip uses convective drying with agitation. Analysis of the diffusion in wood can be used to estimate the time to dry lumber to a specified moisture content value. Relationships between the effect of temperature and moisture content allow more accurate predictions and operational evaluations of driers.

The aim of this study was to investigate constant heat source convective and IR drying by comparing the drying curves when batch drying a sample of woodchip biomass whilst controlling the heat source temperature at 328K, 338K, 348K and 358K. This was achieved through comparison of preexponential diffusion coefficients and activation energy, determining the temperature dependency of these terms in convective and IR drying of wetted wood. Lower temperatures increased drying time for both convective and IR drying, with convective drying taking up to 5 times longer than IR. The preexponential diffusion coefficient and activation energy found for IR drying were $3.555 \times 10^{-4} \pm 0.824 \times 10^{-4}$ m².s-1 and $3.405 \times 10^4 \pm 0.065 \times 10^4$ J.mol⁻¹. The convective drying preexponential diffusion coefficient and activation energy *calculated* was $2.948 \times 10^{-7} \pm 0.376 \times 10^{-7}$ $1.619 \times 10^4 \pm 0.037 \times 10^4$ *J.mol*⁻¹ respectively.

Nomenclature

M	[kg]	Mass
X (DB)	[-]	Moisture Content- Dry Basis
W	$[g.s^{-1}M^{-1}]$	Rate of Drying
D	$[m^2.s^{-1}]$	Diffusion Coefficient
L	[m]	Thickness
C	[mol.m ³]	Concentration
J	$[kg.s^{-1}m^{-3}]$	Mass Flux

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t	[s]	Time
T	[°C]	Temperature
MR	[-]	Moisture ratio
D_{Eff}	$[m^2.s^{-1}]$	Effective Diffusion Coefficient
E_a	$[J.Mol^{-1}]$	Activation Energy
k	[W.m ⁻¹ K ⁻¹]	Conductive Heat Transfer
κ		Coefficient
A	$[m^2]$	Surface Area
R	$[J.K^{-1}.mol^{-1}]$	Ideal Gas Constant
q	$[W.m^{-2}]$	Heat Transfer
h	[W.m ⁻³ K ⁻¹]	Convective Heat Transfer
		Coefficient
σ	$[W/m^2K^{-4}]$	Boltzmann constant
ϵ	[-]	Surface emissivity
S	[-]	Saturation
φ	[-]	Porosity
K_{H}	[-]	Henrys Constant
Subscripts		•
e		Equilibrium/Final
0		Initial
i		Species i/Result i
D		Drying
S		Surface
∞		Ambient
p, va, L		Porous Media, Vapour, Liquid

Keywords: Drying, Convective, Infrared, Activation Energy, Diffusion.

1. Introduction

Woodchip production consists of processing wood through three main stages (i) sourcing timber, (ii) chipping of logs, and (iii) drying the chipped wood. With the calorific (MJ/kg) value of woodchip being linearly proportional to its moisture content [1], drying wet chip

is necessary to achieve greater combustion per weight of biomass. Biomass boilers are therefore rated to a defined moisture content of wood feed. Other benefits of drying include quality, characterisation of wood allowing comparison between various origins, reduced energy consumption for transport and storage, driven by a reduction in mass and fungal build up. On the other hand, the fuel sourced from woodchip is generally in higher demand over the winter period when the supply faces a higher moisture content due to both the colder and wetter weather. Therefore, in winter months drying is required to not only improve the fuels quality but also preserve and keep a consistent supply.

Both convective and infrared (IR) drying are used to find the moisture content of woodchip, though British Standards recommend convective drying [2]. Convective drying is in general slower than IR drying as energy, in the form of heat, is first transferred to the air and then the woodchip, whereas in IR, drying energy is transferred directly to the woodchip and water via radiative heat transfer. The rate of heat transfer by convection is commonly described using Fourier's law of conduction and Newton's Law of Cooling, comparatively IR heat transfer is described using the Stefan-Boltzmann's law of radiation (Equations 1, 2 and 3 respectfully)[3].

$$q = -hA(T_s - T_{\infty})$$

$$q = -k(T_s - T_{\infty})$$

$$q = -\epsilon \sigma (T_s^4 - T_\infty^4)$$

where g is heat transfer (W.m-2), h is the convective heat transfer coefficient (W.m-3K-1), A is the heat transfer area (m2), k is a conductive heat transfer coefficient (W.m⁻¹K⁻¹), ϵ is surface emissivity, σ is the Stefan-Boltzmann constant (W.m-2K-4), T_s is the solid surface temperature (K) and T_{∞} is the ambient temperature (K).

These show the impact of temperature on heat transfer. For IR drying the effect of temperature increase is amplified by the fourth power, compared to that of convective drying for which the equation of heat transfer contains no exponent. The driving force for heat transfer is the temperature gradient along the woodchip and between the atmosphere and the woodchip.

2. Mathematical Models

The falling rate period of drying shows the behaviour of moisture diffusing through material. After an initial evaporation of the surface moisture, the rate of mass loss decreases as moisture has to diffuse from the centre of the material to the surface of the solid, obeying Fick's law of mass transfer;

$$J_i = -D \frac{dC_i}{dx}$$

Where the mass flux J_i is defined by the diffusion coefficient, D, and the concentration gradient, $\frac{dC_i}{dx}$. Transforming the second Fick's law [4], by considering the rate of accumulation of a substrate, Ci. in this case water vapour in a control volume, the resulting relationship is;

$$\frac{dC_i}{dt} = \frac{d}{dx} \left(D \frac{dC_i}{dx} \right)$$
 5

Expressing the ratio of moisture, MR, within the woodchip at time t from the current M_i, initial M₀ and final masses M_e;

$$MR = \frac{M_i - M_e}{M_0 - M_e}$$

Using the assumptions of:

- Symmetric uniform distribution of moisture within the initial sample.
- Symmetric mass transfer with respect to the centre of the solid.
- Constant diffusion coefficient.
- Negligible shrinkage.
- Instantaneous evaporation at the surface (i.e. the concentration on the face from which the diffusing substance emerges is maintained at effectively zero).
- Steady state conditions for the finite interval of time is defined after IR drying at 105°C [5].

The drying rate, W_D, is defined as the amount of moisture removed from the dried material per unit time and unit surface area [6]; $W_D = \frac{-M_e dX}{Adt}$

$$W_{D} = \frac{-M_{e}dX}{Adt}$$

Where X is the moisture content on a dry basis, Me is the equilibrium dry mass, A is area and t is time in seconds. For a constant surface area, A, and mass of dried solid, Me, the rate, WD, can therefore be determined through differentiation of the moisture ratio over time. $W_D A = (M_0 - M_e) \left(\frac{dMR}{dt}\right) = \frac{dM_t}{dt}$

$$W_{D}A = (M_{0} - M_{e}) \left(\frac{dMR}{dt}\right) = \frac{dM_{t}}{dt}$$

To define the moisture ratio diffusion coefficients and activation energy can be calculated from singular pieces of chip of known thickness L.

Using the following model by Crank [7];

$$MR = \frac{M_t - M_e}{M_0 - M_e} = \frac{8}{\pi^2} \sum_{n=0}^{\infty} \frac{1}{(2n+1)^2} e^{-(2n+1)^2 \left(\frac{\pi^2 D_{eff}}{L^2}\right)t} 9$$

Using time, t (s), the effective diffusion coefficient, D_{eff} (m.s⁻¹) and thickness, L(m).

For long drying times;

$$MR = \frac{8}{\pi^2} e^{-\left(\frac{\pi^2 D_{eff}}{L^2}\right)t}$$
 10

Temperature Dependence

The impact of temperature on effective diffusivity can be estimated using the Arrhenius profile of effective diffusivity, Equation 11.

$$D_{\alpha\beta\beta} = D_{\alpha} e^{\left[-\frac{E_a}{RT}\right]}$$

 $D_{eff} = D_0 \, e^{\left[-\frac{E_a}{RT}\right]} \qquad \qquad 11$ Within these equations the pre-exponential diffusion coefficient, D_0 (m².s⁻¹) and activation energy, $E_a(J.mol^{-1})$ are calculated constants, while temperature, T(K), and the ideal gas constant, R(8.314J.K-1.mol-1) are predefined and time is a dependent measured variable.

3. Materials and Method

The woodchip was supplied by Bowland Bioenergy Ltd. This wood had been externally stored as logs and chipped, with no prior drying other than natural weathering. A single chip of softwood was used which was then soaked in deionised water for 12 hours resulting in a wetted chip weighing ≈3g prior to the drying procedure.

3. 1. Drying Procedure

Convective Drying

Adjacent to a set of scales, with an accuracy of ± 0.0005 g, a variable temperature heat gun was mounted (shown in Figure 1) which passes air at 1.6m/s over a sample tray. The weight on this tray was recorded every 5 seconds, with the heat gun causing an initial constant fluctuation in the weight and variations around this caused by changes in the air flow. The temperature of the air was measured 4cm from the outlet of the heat gun and 5cm from the centre of the 10cm ø sample tray. This temperature stayed at the set temperature ±5°C and has an on/off controller set to turn off +6°C from the set point which turns the heat source off until the temperature drops below this value. The ambient temperature of the room was 22°C at the start of every experiment and the equipment was uncovered, increasing this ambient temperature close to the sample over time.

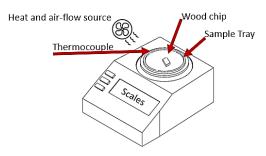


Figure 1. Experimental Setup - Convective.

IR Drying

The infrared (IR) halogen drier contains scales with a weight accuracy of ± 0.0005 g, and was programmed to stop when a change of 0.001g per 99 seconds was reached and the total weight recorded every 5 seconds. The moisture balance was setup to have a target temperature of 145% of the set temperature for the first instance and when the set temperature is reached the temperature is maintained. This setup resulted in some overshoots in temperature control by a few degrees in the first minute and then remained at the set drying temperature ± 1 °C, with tighter temperature control than the convective drying. The ambient temperature was found to have no discernible effect on the results of IR drying due to the enclosed chamber of the dryer. The halogen heater had a heat duty of 400W. For Both Experiments

A single woodchip weighing ~3g was used for investigating both convective and IR drying. This dimensions woodchip has known 6.13×29.87×25.61mm³. The drying temperatures used were 328K, 338K, 348K and 358K.

In both cases mass data were logged using a bidirectional RS-232 cable connected to a laptop with appropriate data logging software. The mass over time, M_t, was recorded until the mass over time levelled out for 30minutes or until the IR drier automatically stopped and after this original session stopped saved to a file. The first mass recorded in this session is the starting mass M_0 .

To obtain the equilibrium value, Me, the IR drying apparatus was used with the temperature set to 105°C and weight recorded every 5 seconds, and the balance was again programmed to stop when a change of 0.001g per 99 seconds was reached. The final mass value from this session was M_e relating to the previous saved session.

Once Me was obtained the woodchip was then soaked again in deionised water, until the mass reached $M_0\pm0.5g$ after draining, which took up to 24 hours. The drying procedure for the next set temperature was then repeated to obtain M_t and M_e data for each experimental setup.

4. Results Analysis

A fit to Equation 15 to experimental results for the moisture ratio has been evaluated using MATLAB and Non-linear Least Squares regression analysis. To identify the goodness of the fit commonly used statistical criteria were used: these include the sum of the square estimate errors (SSE), Equation 12, residual squared (R2), Equation 13, and root mean squared error (RMSE), Equation 14.

$$\begin{aligned} \text{SSE} &= \sum_{i=1}^{n} \! \left(\text{MR}_{i} - \widehat{\text{MR}}_{1} \right)^{2} & 12 \\ \text{R}^{2} &= 1 - \frac{\sum_{i=1}^{n} \! \left(\text{MR}_{i} - \widehat{\text{MR}}_{i} \right)^{2}}{\sum_{i=1}^{n} \! \left(\text{MR}_{i} - \widehat{\text{MR}}_{i} \right)^{2}} & 13 \\ \text{RMSE} &= \left[0.5 \sum_{i=1}^{n} \! \left(\text{MR}_{i} - \widehat{\text{MR}}_{i} \right)^{2} \right]^{0.5} & 14 \end{aligned}$$

$$\sum_{i=1}^{n} (MR_{i} - MR_{i})^{2}$$

$$DMSE = \left[0.5 \sum_{i=1}^{n} (MR_{i} - MR_{i})^{2} \right]^{0.5}$$
14

$$RMSE = \left[0.5 \sum_{i=1}^{n} \left(MR_i - \widehat{MR}_i\right)^2\right]^{0.5}$$
 14

These results show the goodness of fit to the models within Table 1. The closer to 1 R² is and the smaller the SSE and RMSE values are, the better the goodness of fit [8].

5. Results And Discussion

The measured mass vs time data for IR and convective drying shows that convective drying takes longer than IR at the same temperature shown in Figure 2 and Figure 3. The drying time was up to 5 times longer for Convective drying than IR drying. IR drying also resulted in a lower final mass.

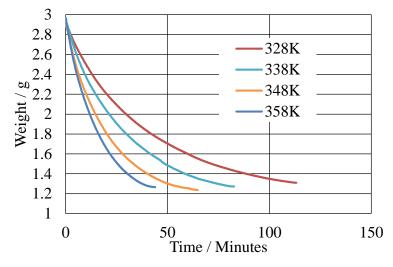


Figure 2. Plot of Experimental IR Drying Curves.

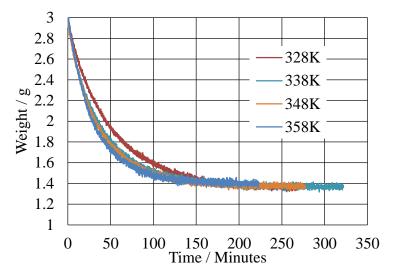
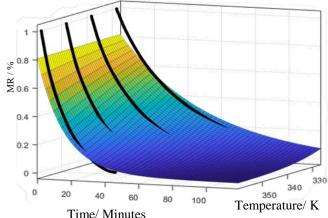


Figure 3. Plot of Experimental Convective Drying Curves.

The thickness of the woodchip, L in Equation 10, was measured as 0.00613 m. Equations 10 and 11 can therefore be combined making Equation 15, which can be fitted to the data using MATLAB Curve Fitting Tool.

$$MR = \frac{8}{\pi^2} e^{-\left(\frac{\pi^2 D_0 e^{\left[-\frac{E_a}{8.314T}\right]}}{0.00613^2}\right)} t$$
 15

It is noticeable that rate, or steepness of the curves in Figure 2 and Figure 3 increases with temperature. Therefore the effective diffusivity term will also increase with temperature, as suggested by the Arrhenius equation (Equation 11). From Equation 15 a lower activation energy and pre-exponential diffusion coefficient decrease the drying time, the wider distribution of the IR drying curves in Figure comparison to Figure suggests a greater activation energy, and the faster drying time suggests a larger preexponential diffusion coefficient.



Time/ Minutes
Figure 4. IR Moisture Ratio Results and Equation 15 Fit.

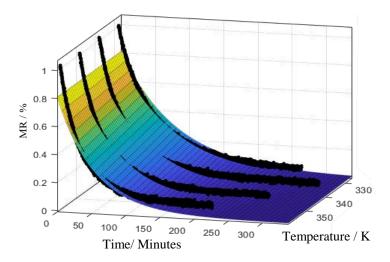


Figure 5. Convective Moisture Ratio Results and Equation 15 Fit.

Table 1. Pre-Exponential Diffusion Coefficient and Activation Energy Results (with 95% confidence bounds).

	$D_0 / m^2.s^{-1}$	E_a / J.mol ⁻¹
IR	3.555×10^{-4}	3.405×10^4
	$\pm 0.824 \times 10^{-4}$	$\pm 0.065 \times 10^4$
Convective	2.948×10^{-7}	1.619×10^4
	$\pm 0.376 \times 10^{-7}$	$\pm 0.037 \times 10^4$

The activation energy and pre-exponential diffusion coefficient for IR drying, from Table 1, are greater than the convective results. The order of magnitude of pre-exponential diffusion coefficient was much greater for IR drying, whereas for the activation energy the orders of magnitude are the same for both IR and convective drying. Previous experiments have found that for Casuarina Equisetifolia wood the preexponential diffusion coefficient, D_0 , and activation energy, E_a , were determined as 2.15×10^{-2} m² .s⁻¹and 4.527×10^4 J.mol⁻¹ respectively [9]. The IR drying results obtained in this analysis are of the same order of magnitude and a similar value to those reported in [8]. The most commonly reported data relating to Equation 15 are for foodstuffs, the general range reported for the pre-exponential diffusion coefficients of foodstuffs are generally within 10^{-11} – 10^{-6} m² s⁻¹ [10] which the convective pre-exponential diffusion coefficient for convective drying is within. For food materials, activation energy is 1.27 to 11×10^4 J.mol⁻¹[11] which both results are within. The diffusivity value found for IR drying of woodchip is outside of this range, but between this range for fruit and the results for drying of Casuarina Equisetifolia wood.

Something to note from Equation 10 is that, though this model is widely used [12,13,14], the equation itself will never be an exact fit to moisture ratio (MR) based on the definition of MR. As MR is the percentage moisture in the sample based on the initial moisture content. From time zero the value of MR is 1, however, Equation 10 will always be $\frac{8}{\pi^2}$ when t is zero and after time zero MR will be below this value, shown by the offset of the curve fits in Figure 5 and Figure 4. This is further supported by the evaluation of goodness of fit shown in Table 2 with a large SSE value and an R² value below 0.95, below the R² values found from previous fits of woodchip drying to other experimental models[15,16] with multiple terms. The model used was formed from the first term of Equation 9, a different method to fit this equation would be to fit the experimental data to multiple terms of the Fourier series equations [17].

Table 2. Statistical Analysis of Goodness of Fit to

$8\pi 2e - \pi 2D0e - Ea8.314T0.006132t$			15.	
	SSE	\mathbb{R}^2	RMSE	
IR	29.48	0.9474	0.04747	
Convective	16.97	0.9724	0.03602	

Equation 15 is exponential approaching zero, as the equilibrium mass after secondary heating (used as the dry mass) was less than that at the end of the measured mass over time period, shown in Table 3, the final plotted results for MR are offset from zero. This difference in mass from the plotted period and IR drying at 105°C is due to the activity of water; at lower temperatures and moisture content water activity is lower[18], reducing the total moisture loss over the drying period.

Table 3. Experimental Initial Mass, Final Mass During Experiments, and Equilibrium Mass.

Temperatur	IR		Convective			
e / K	M_0	M_{end}	M_{e}	M_0	M_{end}	M_{e}
328	2.926	1.309	1.230	2.973	1.363	1.246
338	2.960	1.272	1.226	2.909	1.360	1.251
348	2.930	1.237	1.224	2.935	1.365	1.254
358	2.968	1.266	1.225	2.984	1.375	1.255

Table 3 shows the starting masses for all experiments were within a range of 0.1g, for IR drying the weights at the end of the temperature-controlled experiment varied by 0.5g, for convective the lowest measured weight was taken which differs by 0.015g. The

equilibrium weights for all experiments were within 0.03g of each other showing a good repeatability.

6. Conclusions

This study showed the drying behaviour of a single woodchip, exhibiting falling rate characteristics and suggesting that the drying process was diffusively dominant. The temperature dependency of the drying process in the model used was determined through the Arrhenius equation.

Key Findings

- Convective drying takes longer than IR drying.
- The pre-exponential diffusion coefficient and activation energy found for IR drying using Equation 10 proposed by Crank was found to be $3.555 \times 10^{-4} \pm 0.824 \times 10^{-4}$ m².s⁻¹ and $3.405 \times 10^{4} \pm 0.065 \times 10^{4}$ J.mol⁻¹, for convective drying $2.948 \times 10^{-7} \pm 0.376 \times 10^{-7}$ m².s⁻¹ and $1.619 \times 10^{4} \pm 0.037 \times 10^{4}$ J.mol⁻¹ respectively.
- Crank's equation could not effectively represent MR data due to the equation for MR using the initial mass as an 100% reference mass.
- IR drying resulted in a higher mass loss over the drying period than convective drying.

Highlights and Limitations

In the convective drying experimental test fluctuations in weight from airflow patterns made it difficult to produce live weight measurements of samples as noise within results through this method increased the measurement error. The ambient air temperature increased over time with convective drying, making it difficult to alter the sample temperature rather than the heat source temperature, the heat source temperature did not equate to the temperature of the air that reached the sample and so temperature control in IR drying was more effective due to the contained space.

7. Recommendations

The Crank model used contains a lumped diffusion term governing moisture transport, however, water moves through woodchip via diffusive moisture transport and capillary moisture transport. Thus, to properly account for this physical phenomena, separate terms for diffusive and capillary transport of liquid water and water vapour could be used to further describe the movement of moisture. For example the diffusion through a porous media $D_{\rm p}^{\rm s}$ can be split into a liquid and

vapour term, as shown in Equation 16 proposed by Millington and Quirk in 1961[19].

$$D_{p}^{*} = \phi^{\frac{4}{3}} \left(S_{va}^{\frac{10}{3}} D_{va} + S_{L}^{\frac{10}{3}} D_{L} / K_{H} \right)$$
 16

Where D_{va} is vapour diffusion, D_L is liquid diffusion, S_L and S_{va} are vapour and liquid saturation, ϕ is the porosity and K_H is henrys constant.

A different method to fit data to the Crank model could be to use multiple terms of the Fourier series expansion of Equation 9[20] (note the Crank model used was formed from only the first term of Equation 9).

The moisture content also depends on several other factors for instance, it can be affected by airflow, temperature, pressure, relative humidity, and structure on diffusivity. The full effects of drying type and conditions on moisture loss over time can be found using computational and experimental investigation. This analyses could consider effects such as; responses to high temperature drying, low temperature drying, convective drying, IR drying, microwave drying or freeze drying using different models and situations.

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