

mailed  
4-4-97

# AIR CONTAMINANT EVALUATION TEST REPORT

## **COWLITZ STUD COMPANY** **Dry Kiln VOC Emission Factors**

February 25-March 13, 1997

Prepared for

Cowlitz Stud Company  
P.O. Box 219  
Randle, Washington 98377

By

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&  
David R. Rossman P.E.



RECEIVED  
AUG 11 1997  
SOUTHWEST AIR POLLUTION  
CONTROL AUTHORITY

## Introduction

Between February 25 and March 13, 1997, four samples of Cowlitz Stud Company lumber were dried in Horizon Engineering's laboratory dry kiln. Volatile organic compounds (VOCs) were continuously measured in the test kiln using the Dettinger Method. The laboratory test was done instead of a source test due to the expense and uncertainties involved in testing an actual dry kiln.

Charlie Allen, of Cowlitz Stud, and Heinz Dettinger, their consultant, arranged for the work. Horizon Engineering personnel David Broderick did the testing and Michael Wallace did the data processing. A copy of the test method has been included in the Appendix.

## Summary of Results

Table 1 summarizes the results of the testing. Figures 1 through 4 are plots of the calculated emission factors for the range of percentage H<sub>2</sub>O (wet basis) of the wood samples. It should be noted that the results are based on an actual board foot basis, not the nominal dimensions of each sample board.

Detailed results and sampling parameters are included in the Appendix.

Table 1  
Summary of Results

Results	Units				
Species		Pine	Pine	Doug Fir	Hemlock
Sample size	bd ft	8.4	8.4	7.8	8.0
Initial weight	lb	25.3	23.1	20.1	29.0
Weight loss	lb	7.5	5.3	5.5	12
Test time	hr	54	77	70	75
Avg VOC (dry)	ppmvC	181	145	86	42
Max VOC (wet)	ppmvC	361	317	362	66
Emission Factor					
@ 0% Moisture	lbC/Mbdft	0.78	0.86	0.51	0.25
@ 14% Moisture	lbC/Mbdft	0.27	0.19	0.23	0.10



# Cowlitz Stud Trial #1

## Pine

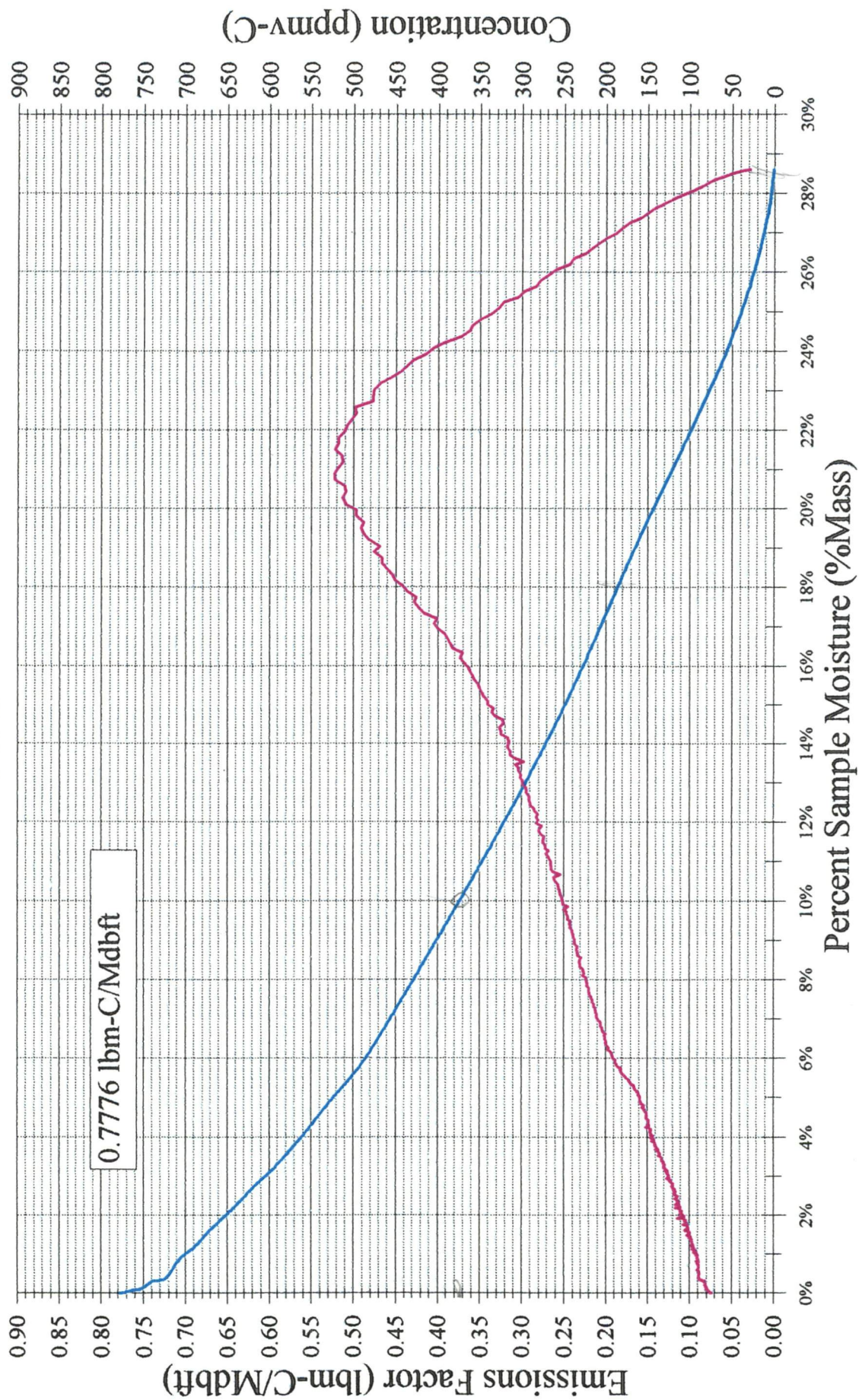
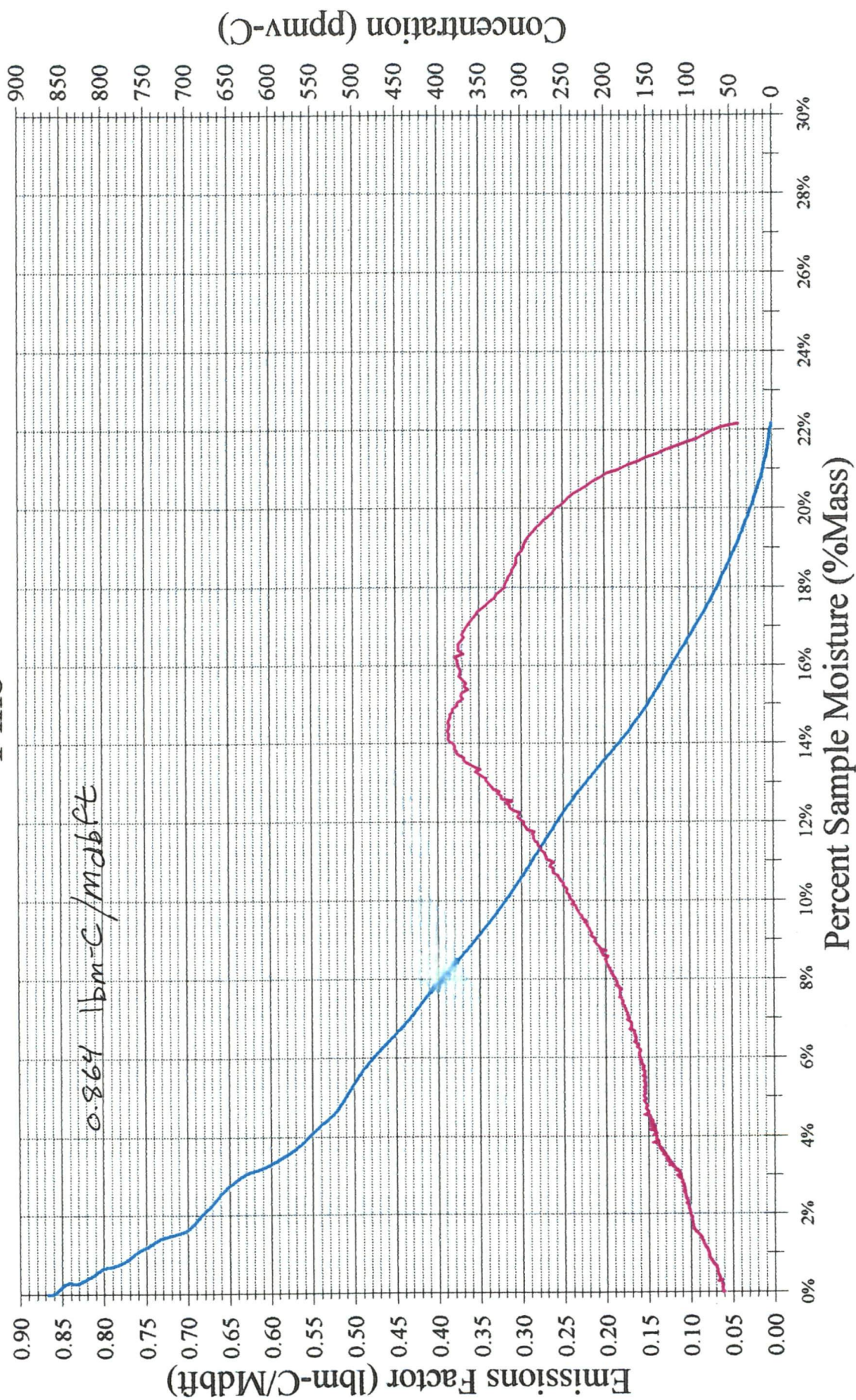


Figure 1



# Cowlitz Stud Trial #2 Pine



— Emission Factor — VOC concentration (ppmv-C Dry)

Figure 2



# Cowlitz Stud Trial #1 Douglas Fir

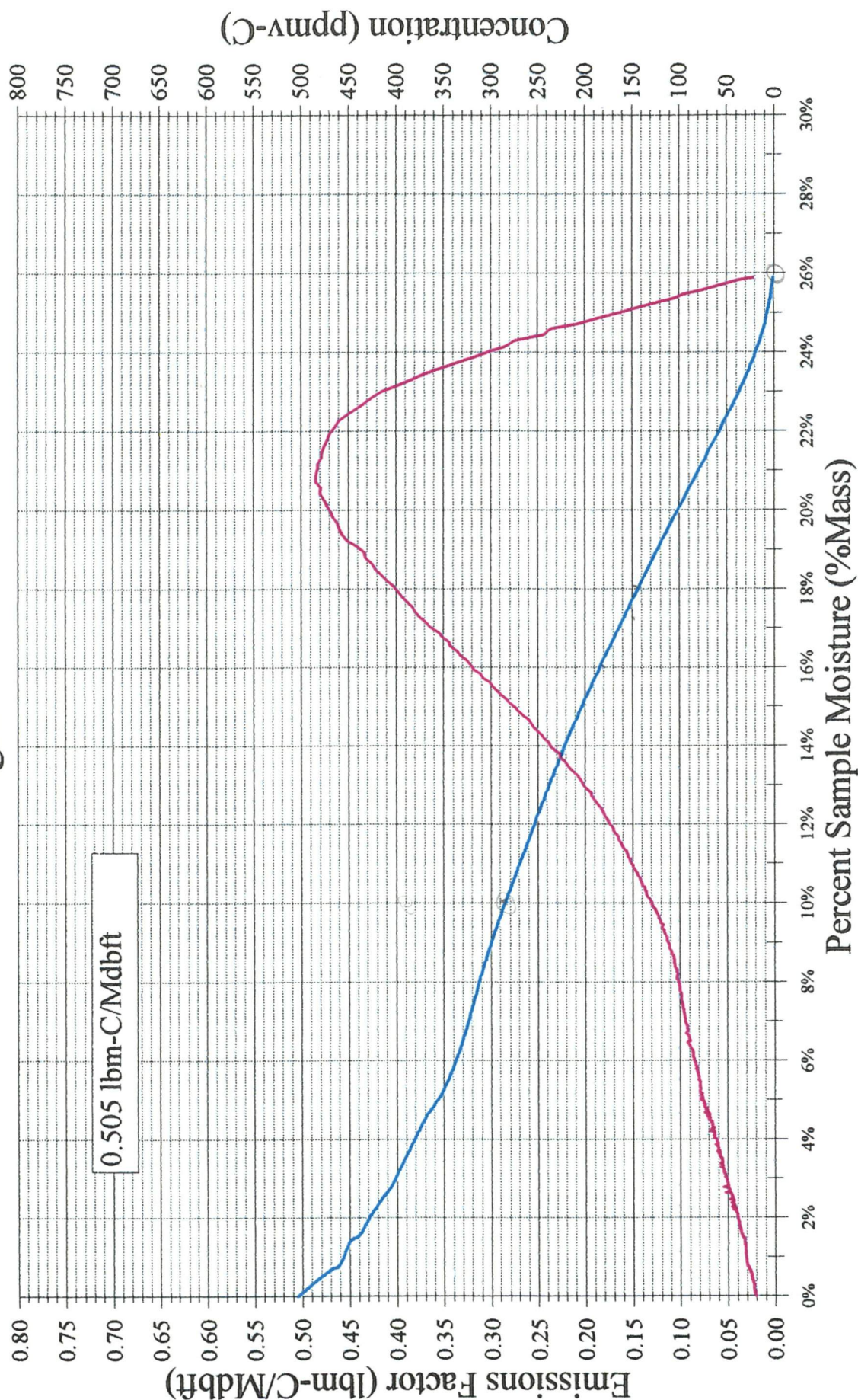


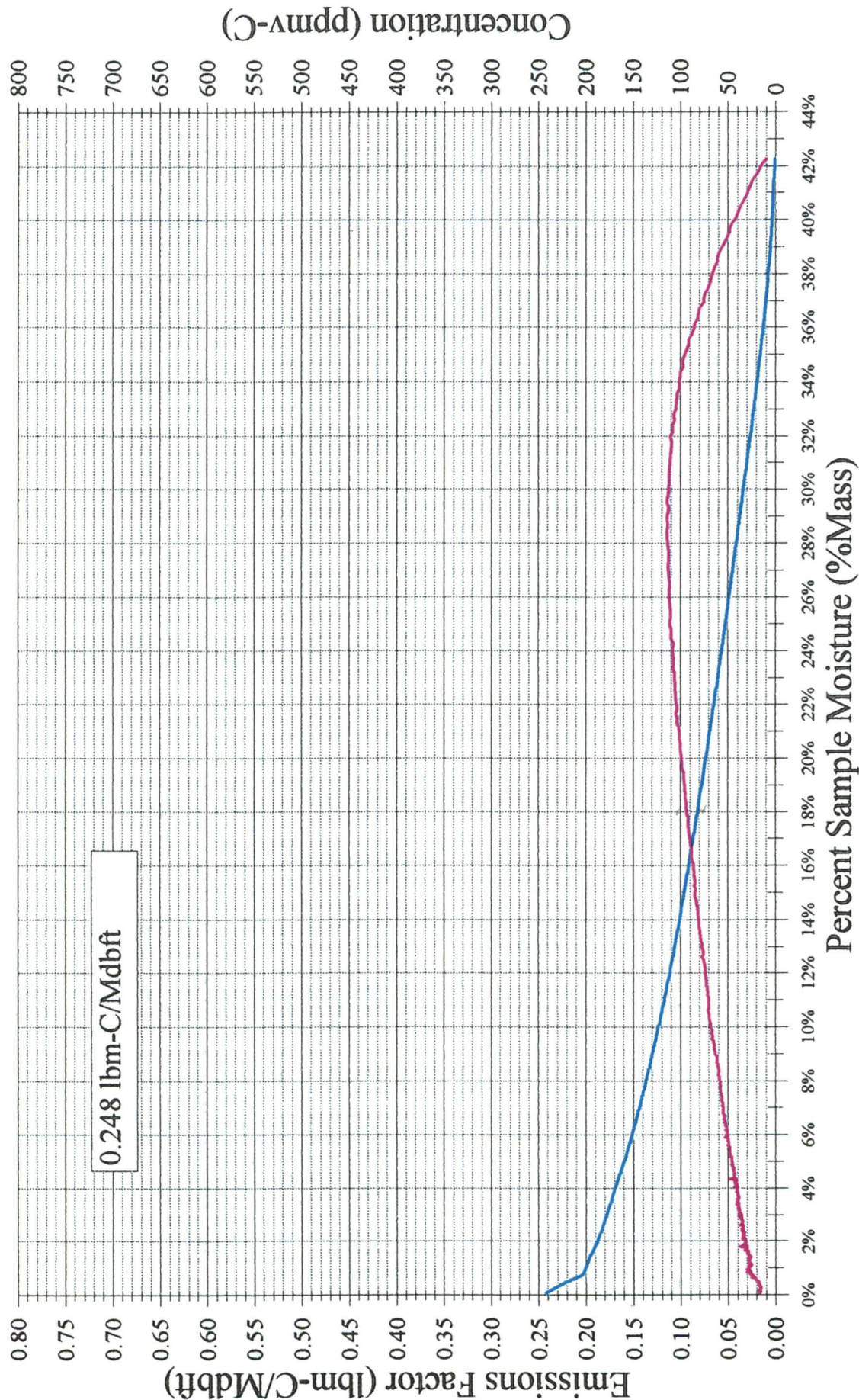
Figure 3

— Emission Factor — VOC concentration



# Cowlitz Stud Trial #1

## Hemlock



— Emission Factor — VOC concentration

Figure 4



### **Rationale for the Test Method**

Testing a dry kiln is difficult, costly, and there are many uncertainties when using the standard EPA Method 25A, because of the following conditions:

- a.) Lumber drying can take over 100 hours to process one load.
- b.) Most dry kilns have multiple vents and often have significant leakage around the loading doors.
- c.) The venting process is periodic. The vents open to release moisture and VOC's in an irregular pattern.

The multiple vent configuration of most dry kilns and the periodic venting makes it difficult to quantify the total exhaust rate. Leakage from doors and other gaps is difficult to measure and therefore will produce inaccurate results. In addition, tests would need to be repeated for every species of wood the plant dries.

### **Testing Method**

The Dettinger method, applied to the test kiln, employs EPA Method 25A in a controlled manner to measure VOC emissions. The method is assumed to be a worst case analysis, drying to the highest temperature for a normal drying cycle. The test chamber humidity is not controlled but inlet air humidity and volume rate is measured. Normal maximum temperature in a dry kiln is about 200°F but this varies for species and by kiln site. This method allows sample drying times of approximately 36 to 48 hours. Actual drying cycles may take between 36 to over 100 hours.

**General** The test kiln schematic is outlined in the test procedure. An industrial convection oven was used to dry the wood. A 0-100 lb load cell monitors the weight of the drying lumber.

A J.U.M. Engineering VE-7 total hydrocarbon analyzer with heated flame ionization detector and heated sample line was used to measure VOC concentrations at oven conditions. Data was recorded every five minutes by a Rustrak Ranger II data logging system. The accompanying software for the data logger was used to calculate averages for the gas concentrations. Graphic printouts of the data logger information are in the Appendix.

Sample for the analyzers was taken from a fixed sampling probe in the oven. Sample gas was routed through a heated sample line to the continuous analyzer.

**Calibrations** Zero, span, calibration error (linearity) and bias checks were made on the VOC monitor at the beginning and end of each test.

**Calculations** The results from the data logger are corrected for minor instrument drift according to the time when they were calibrated and when the test run was made. System calibration response (bias check) values are used as the basis for these corrections.

**Flow Rate** Air is supplied to the kiln at a constant rate and the total air flow for the entire drying cycle is calculated from dry gas meter readings, in cubic feet. The air in-flow, corrected to a dry standard volume (dscf), is the same as the out-flow dscf (the kiln is kept slightly pressurized). This in-flow rate was used in the pounds of VOC calculation.

**Moisture** The test kiln moisture was calculated from the weight loss of the sample plus the humidity (which is monitored) in the in-flow air. From the total water vapor volume and the total dry air volume, a percentage moisture (by volume) was calculated. Due to the nature of the scale used, the weight loss was not a perfectly smooth curve. The jumps in weight loss caused swings in the ongoing calculated internal moisture of the kiln. To remove these swings, Newton's best fit method was used to derive a smooth (conditioned) curve for the weight loss.

**Board Volume**      The sample boards, as received, were of uniform length. Each sample load was stacked tightly together and measured. The board foot amount of the sample was based on a board foot being 144 cubic inches of wood.

### **Discussion**

The final moisture content of the actual kiln dried lumber should be used to enter the plots of the results figures. Annual emissions of VOC (as carbon) can be calculated based on production of dried lumber.



## **APPENDIX**

### **Nomenclature**

Lab Data

VOC Concentration Plot

Temperature-Humidity Plot

Weight-Moisture Plot

### **Calibration Information**

Meter Box

Standard meter

Thermocouples

Drift Correction

### **Test Method**

Y of meter: 983

Pbar	30.0	300				
Date	2/28/97	3/3				

Bdft (note if dry or wet):

JUM # 2	actual	start bias	end bias
span C#4	952	951.2	949.0
mid C <sub>3</sub> H <sub>8</sub>	601.2	576.0	580.4
mid C <sub>4</sub> H <sub>4</sub>	90.5	91.4	92.0
zero N <sub>2</sub>	0.0	-0.4	-2.0
time & date		10:00am/2-28	15:45/3-3

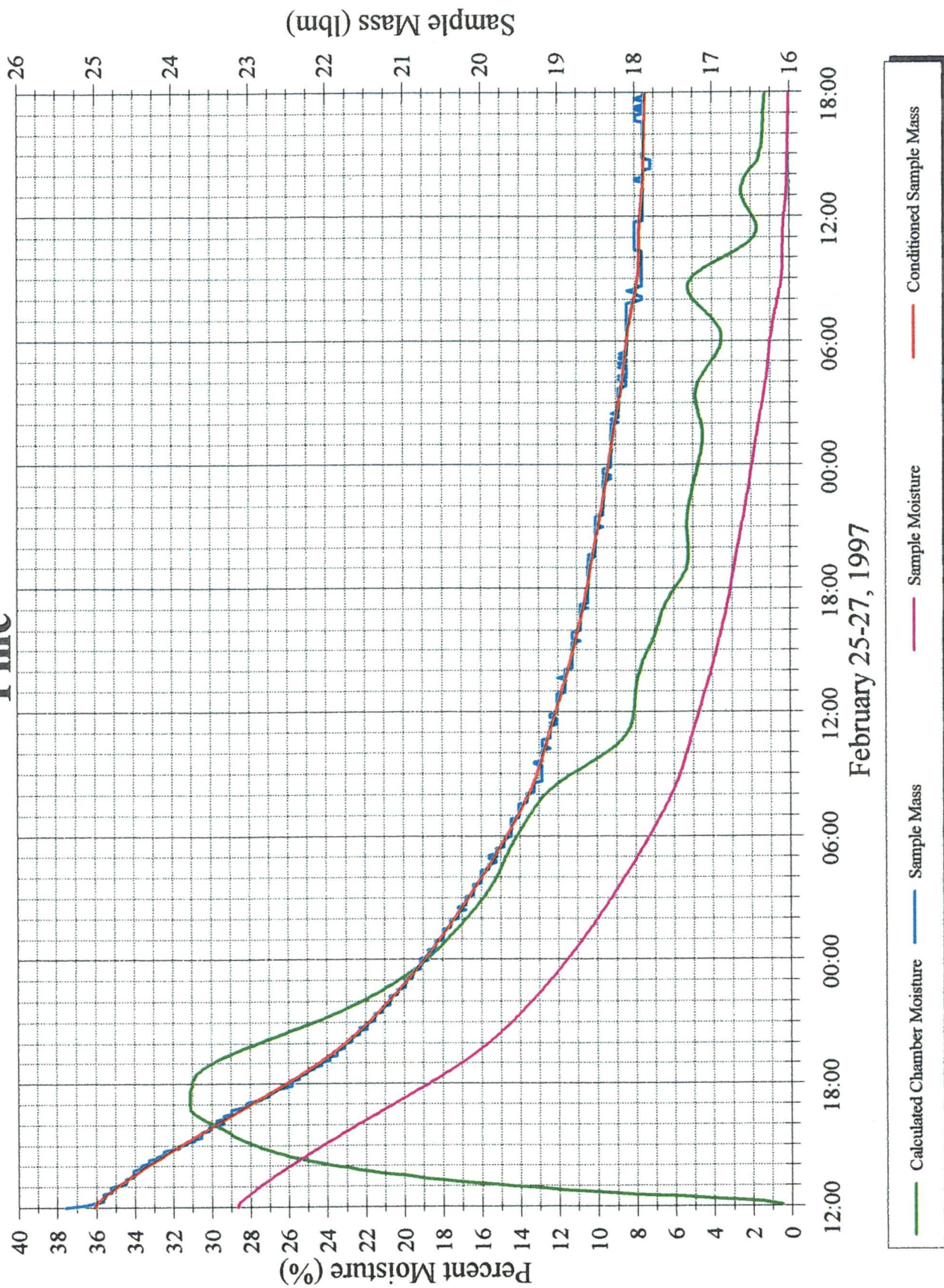
LOAD CELL	actual	start check	end check
high	49.7	51.0	51.0
zero	0.0	0.0	<del>0.0</del> -0.1
time & date		10:00am / 2-28	15:20 / 3-3

[illegible]

Roto Meter  
set at  
 $\approx 12 \text{ L/min}$

# Cowlitz Stud Trial #1

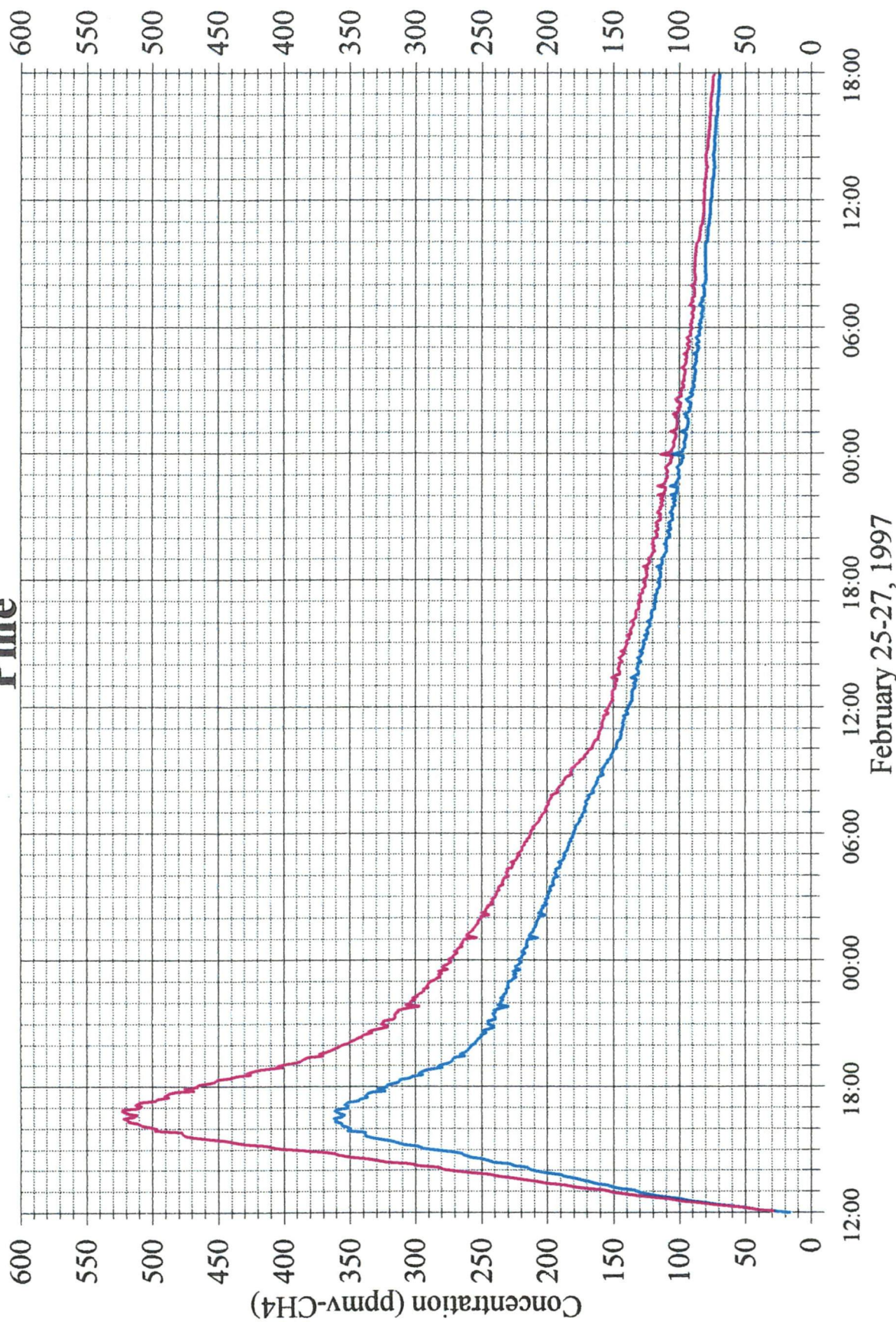
## Pine





# Cowlitz Stud Trial #1

## Pine

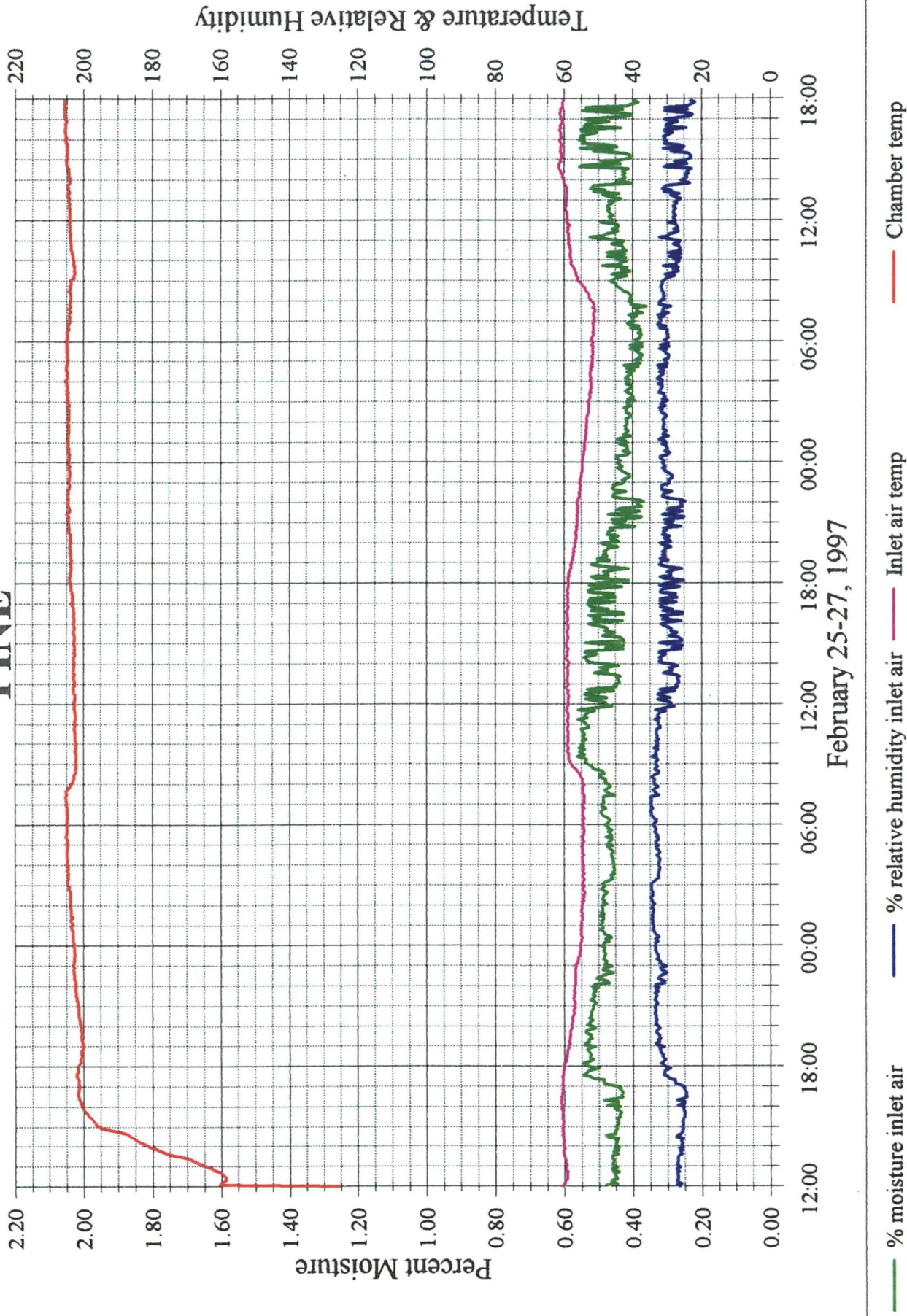


— Concentration Raw Data (Wet) — Corrected Concentration (Dry)



# Cowlitz Stud Trial #1

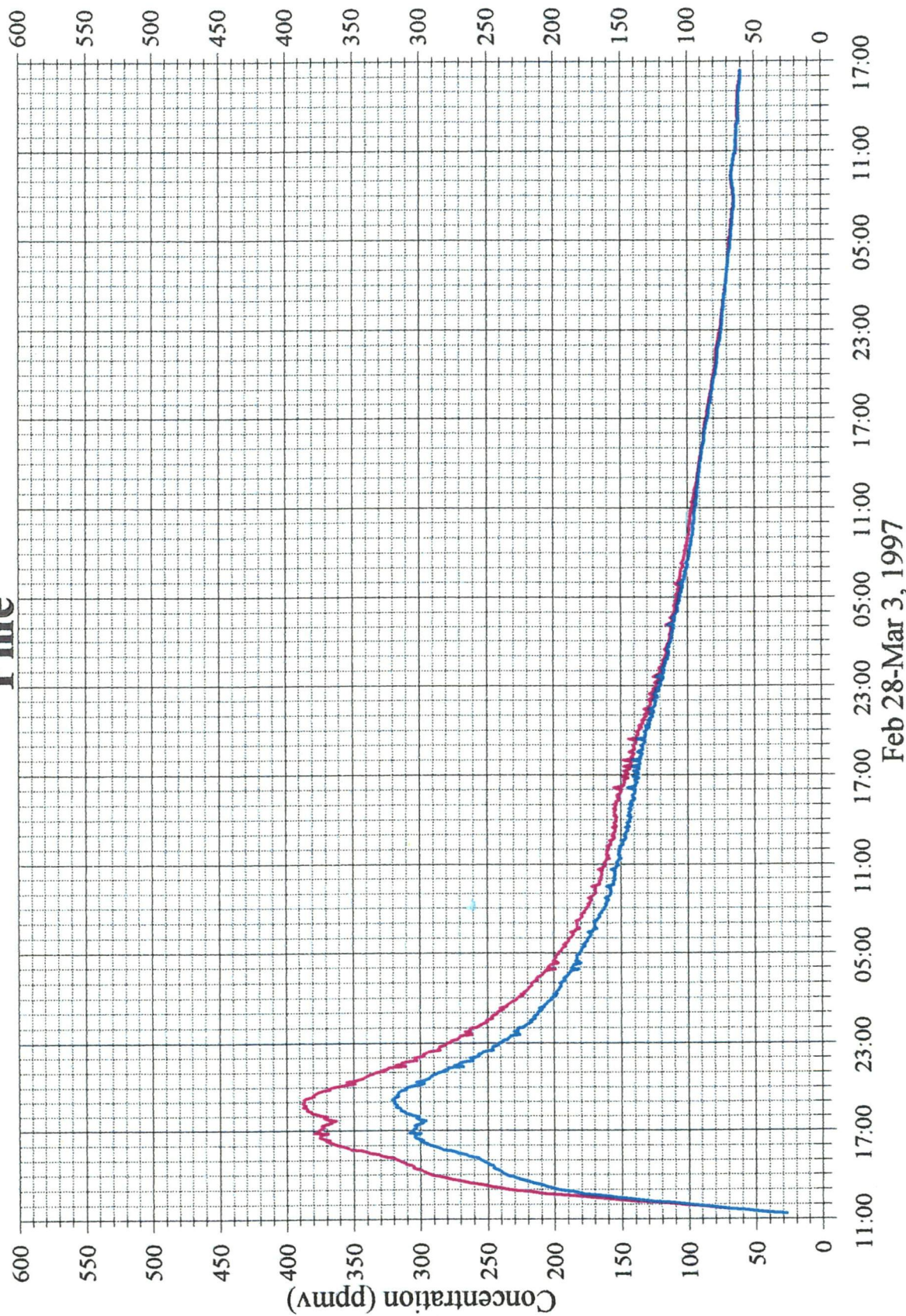
## PINE





# Cowlitz Stud Trial #2

## Pine

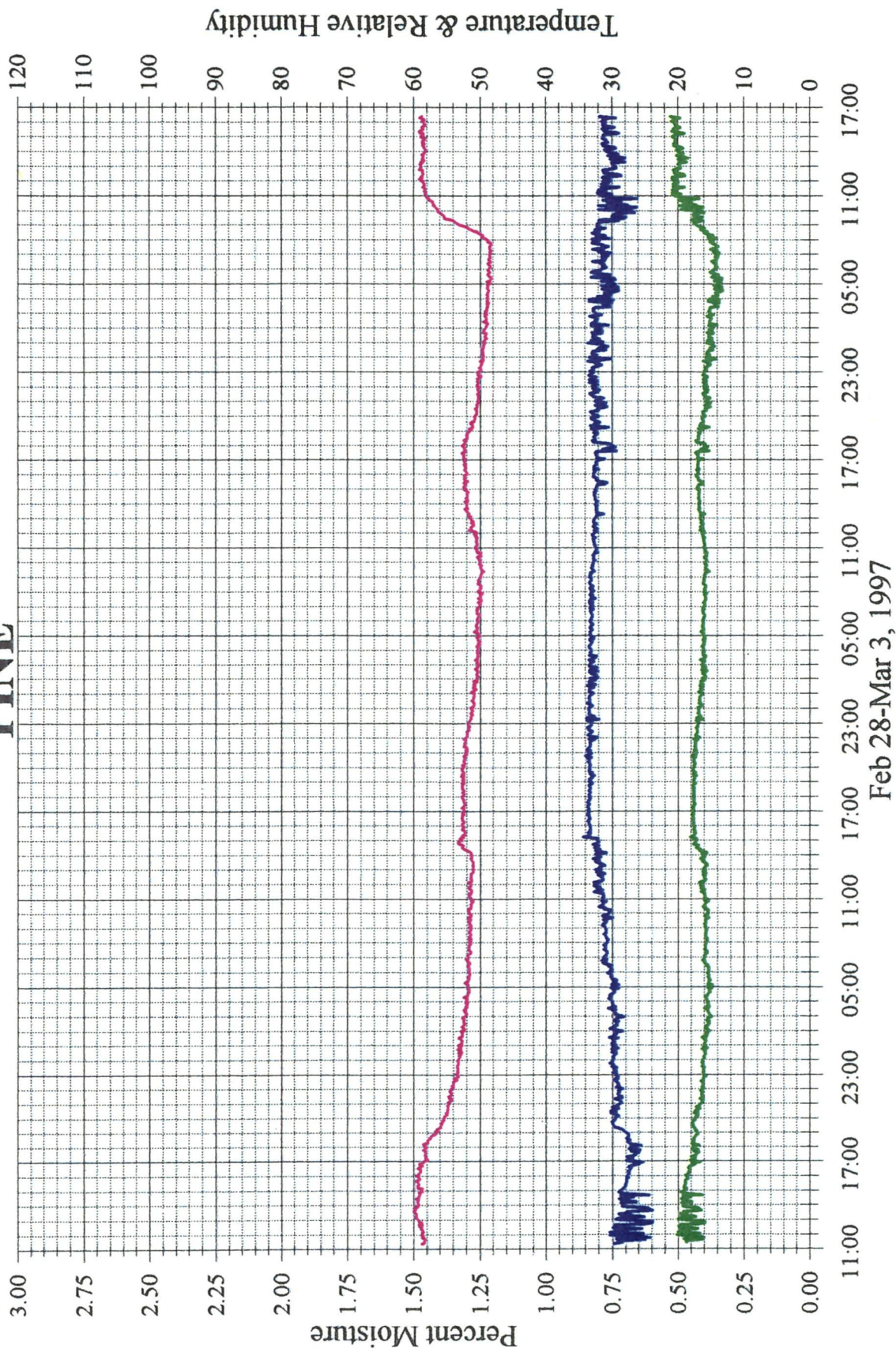


— Corrected Concentration (Dry) — Concentration Raw Data (Wet)



# Cowlitz Stud Trial #2

## PINE

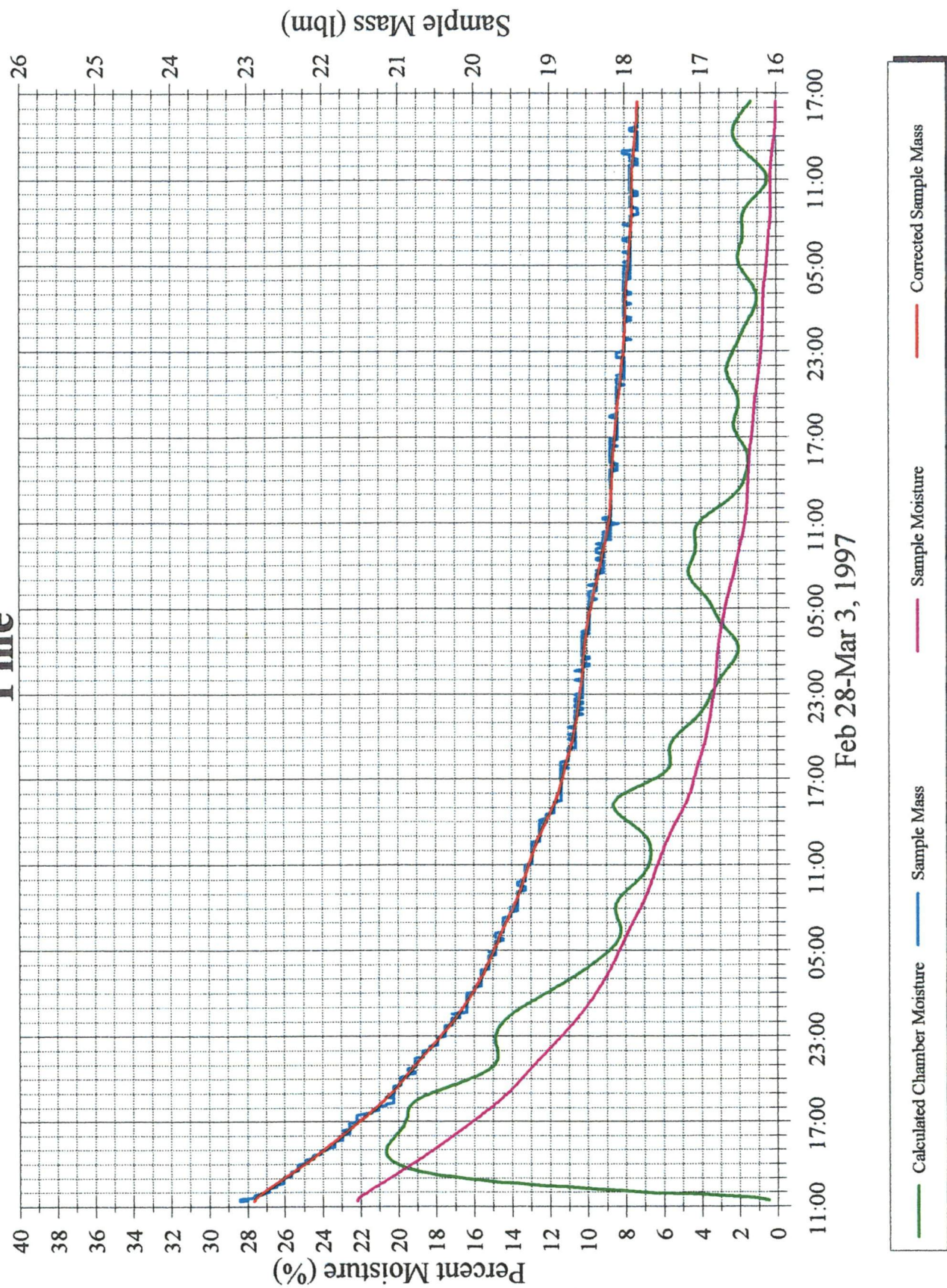


— % moisture inlet air — % relative humidity inlet air — Inlet air temp



# Cowlitz Stud Trial #2

## Pine



Client: Cowhite Studs  
 Species: D. Fur  
 Run: 3  
 Start Time: 16:00  
 Start Date: 3/7/97  
 Y of meter: .983

Pbar	30.1	30.1	30.1			
Date	3/7/97	3/9	3/10			

# of boards: 8

dim of boards:

dim of total load:  $6\frac{3}{8} \times 7\frac{3}{8} \times 2'$

Bdft (note if dry or wet):

JUM #	2	actual	start bias	end bias
span	CH <sub>4</sub>	952	953.2	947.6
mid	60.2	600.2	578	582.2
mid	CH <sub>4</sub>	90.5	92.5	91.1
zero	N <sub>2</sub>	0.0	0.0	-1.2
time & date			1552 3/7	1404 3/10

C<sub>3</sub>H<sub>8</sub>

LOAD CELL	actual	start check	end check
high	94.70	49.4	49.4
zero	0.0	-0.2	-0.1
time & date		15:45 3/7/97	1400 3/10

Meter Reading	Time	Date	Load Weight
299.525	16:10	3/7	20.2
309.195	16:25	3/7	19.7
261.500	13:49	3/9/97	14.79
352.100	1802	3/9	14.71
676.341	933	3/10	14.38
766.679	1359	3/10	14.46

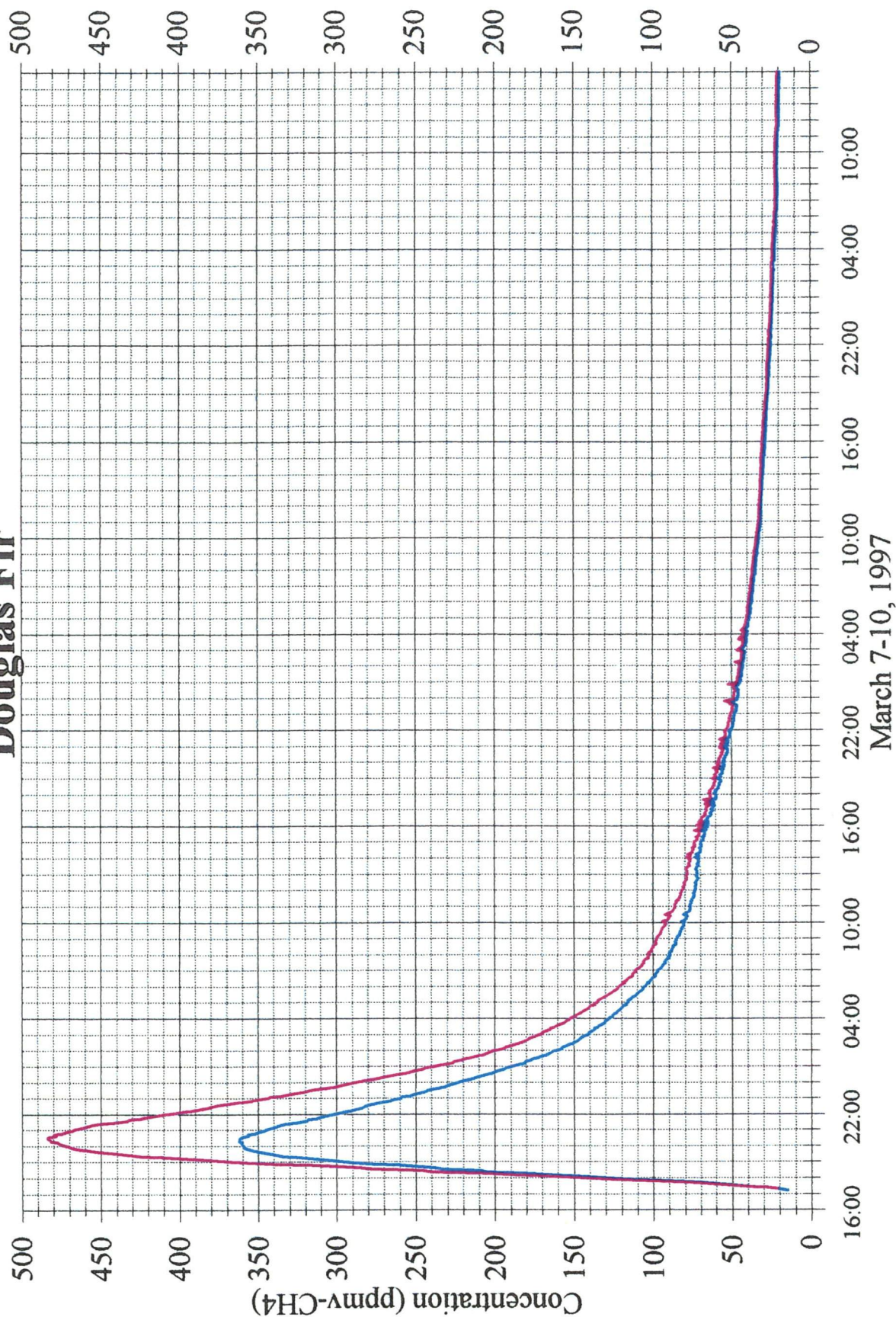
(17:10)  
(17:25)

14:49  
19:02  
10:38  
14:59



# Cowlitz Stud Trial #1

## Douglas Fir

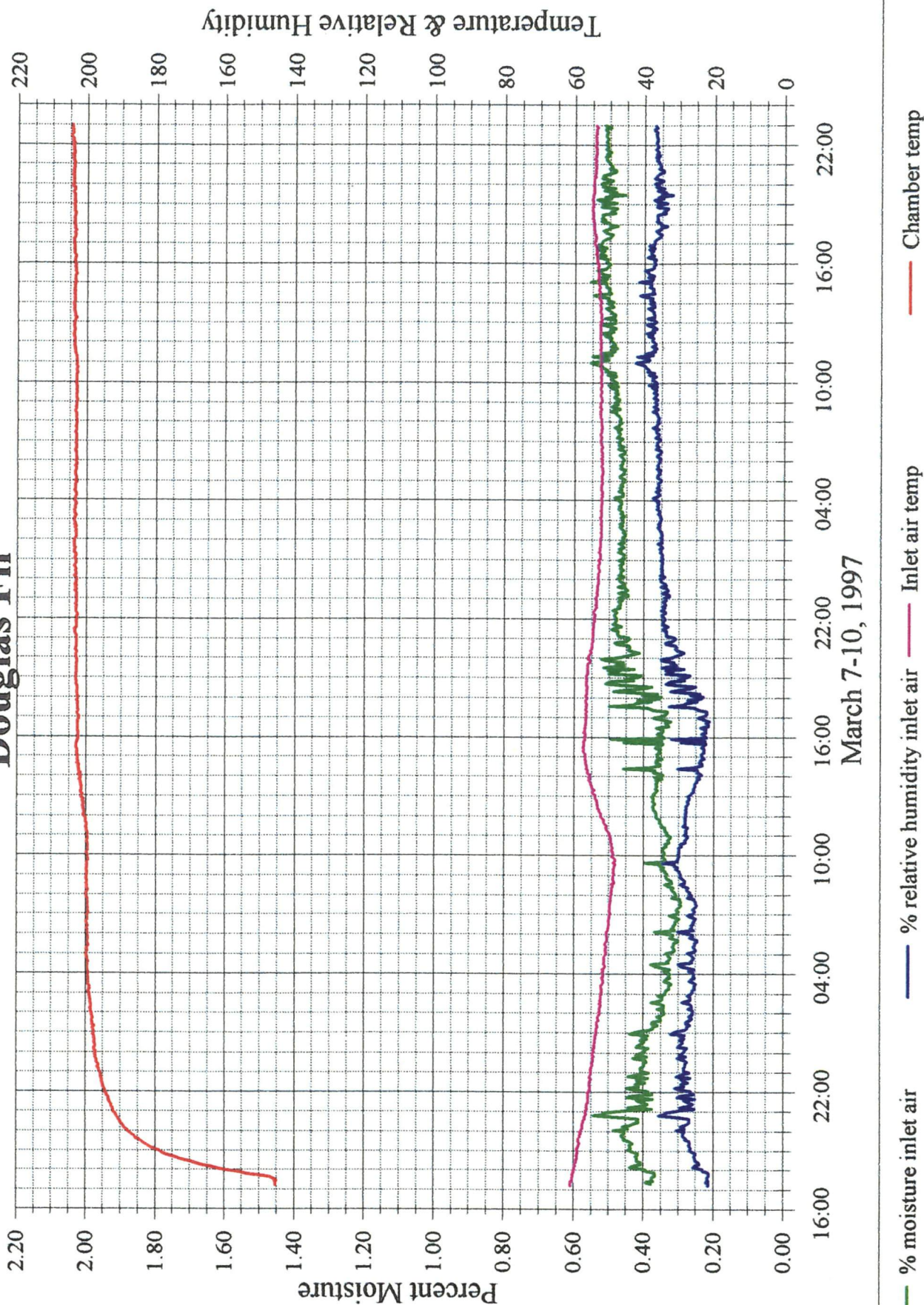


— Concentration Raw Data (Wet) — Corrected Concentration (Dry)



# Cowlitz Stud Trial #1

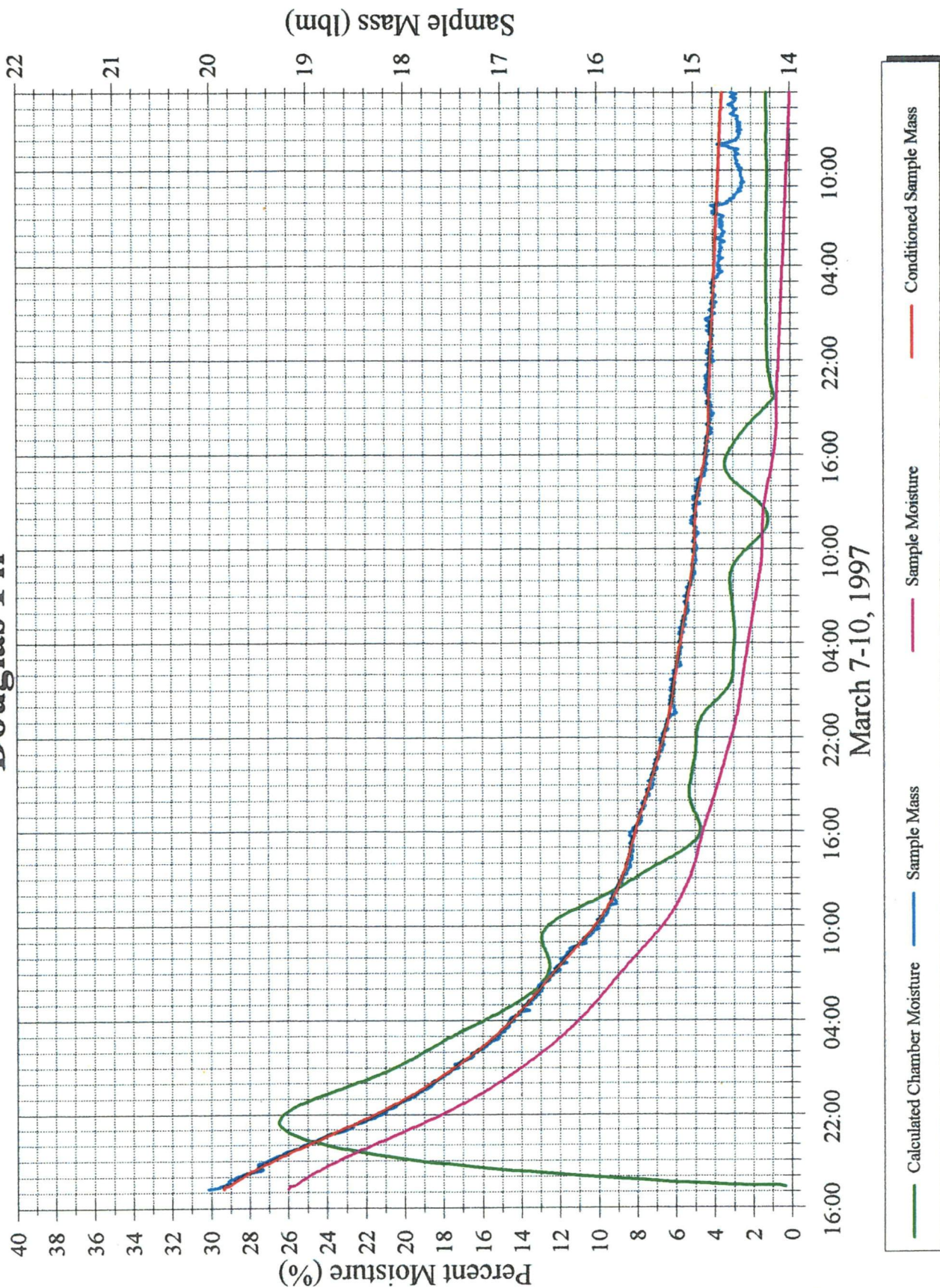
## Douglas Fir





# Cowlitz Stud Trial #1

## Douglas Fir





Client: Schwartz Studs  
 Species: Hemlock  
 Run: 4  
 Start Time: 14:25  
 Start Date: 3/10/97  
 Y of meter: 0.983

Pbar	30.1	30.0	30.00	30.1		
Date	3/10	3/11	3/12	3/13		

# of boards: 8  
 dim of boards:  
 dim of total load:  
 Bdft (note if dry or wet):

JUM #	actual	start bias	end bias
span CH <sub>4</sub>	952	954.9	952.0
mid C <sub>3</sub> H <sub>8</sub>	601.2	582.2	579.4
mid CH <sub>4</sub>	90.5	91.1	91.3
zero N <sub>2</sub>	0.0	-1.7	-2.5
time & date		3/10/97 14:15	3/13/97 17:30

LOAD CELL	actual	start check	end check
high	49.70	49.48	49.43
zero	0.0	-0.05	-0.15
time & date		14:17 3/10	17:25 3/13

Meter Reading	Time	Date	Load Weight
774.531	14:22	3/10	29.0
832.205	17:12	3/10	27.32
130.150	745	3/11	19.60
228.004	17:34	3/11	17.88
611.902	07:45	3/12	16.72
746.304	14:30	3/12	16.79
794.303	16:55	3/12	16.89
94.775	8:02	3/13	8.02 16.42
278.495	17:20	3/13	16.87

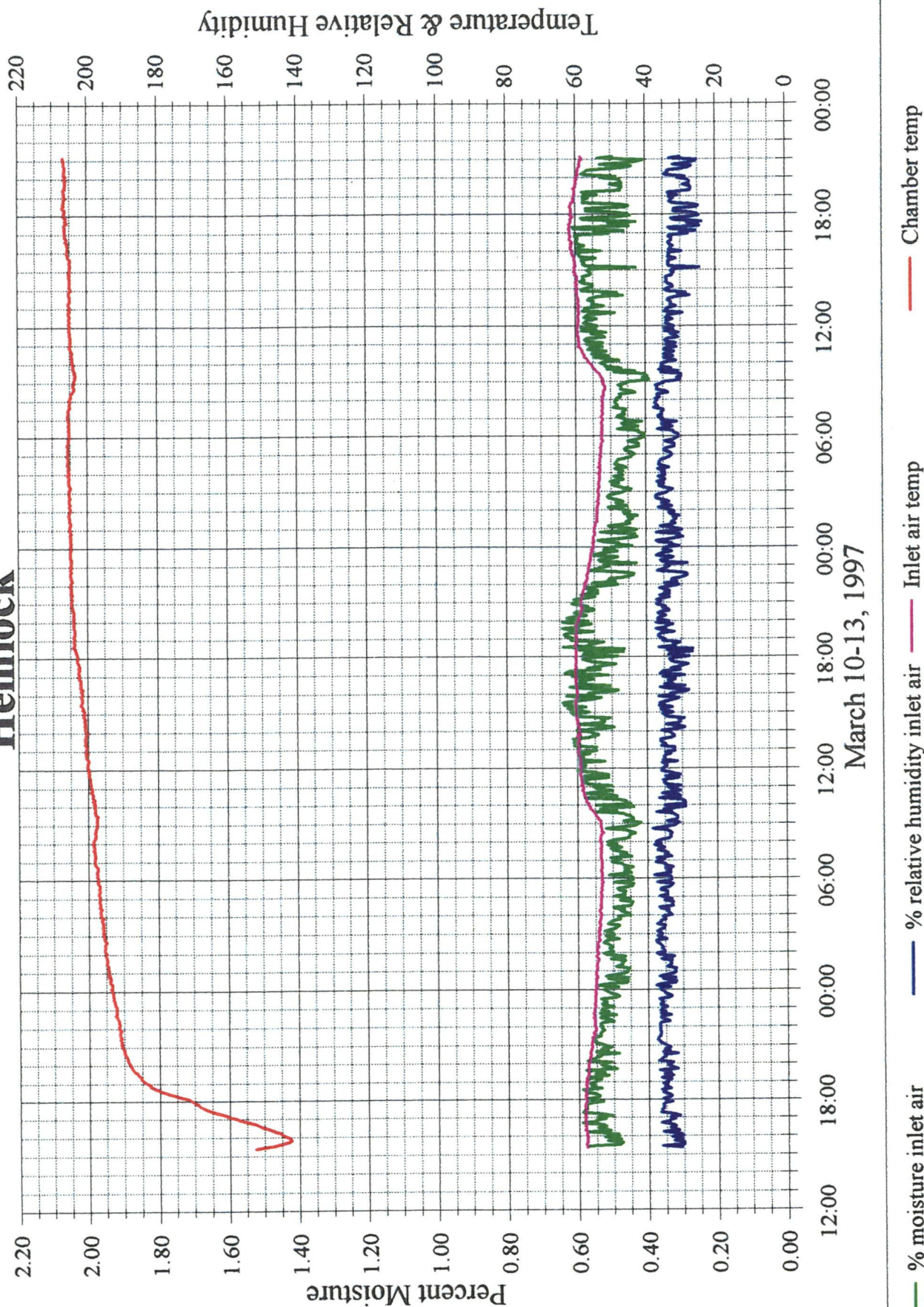






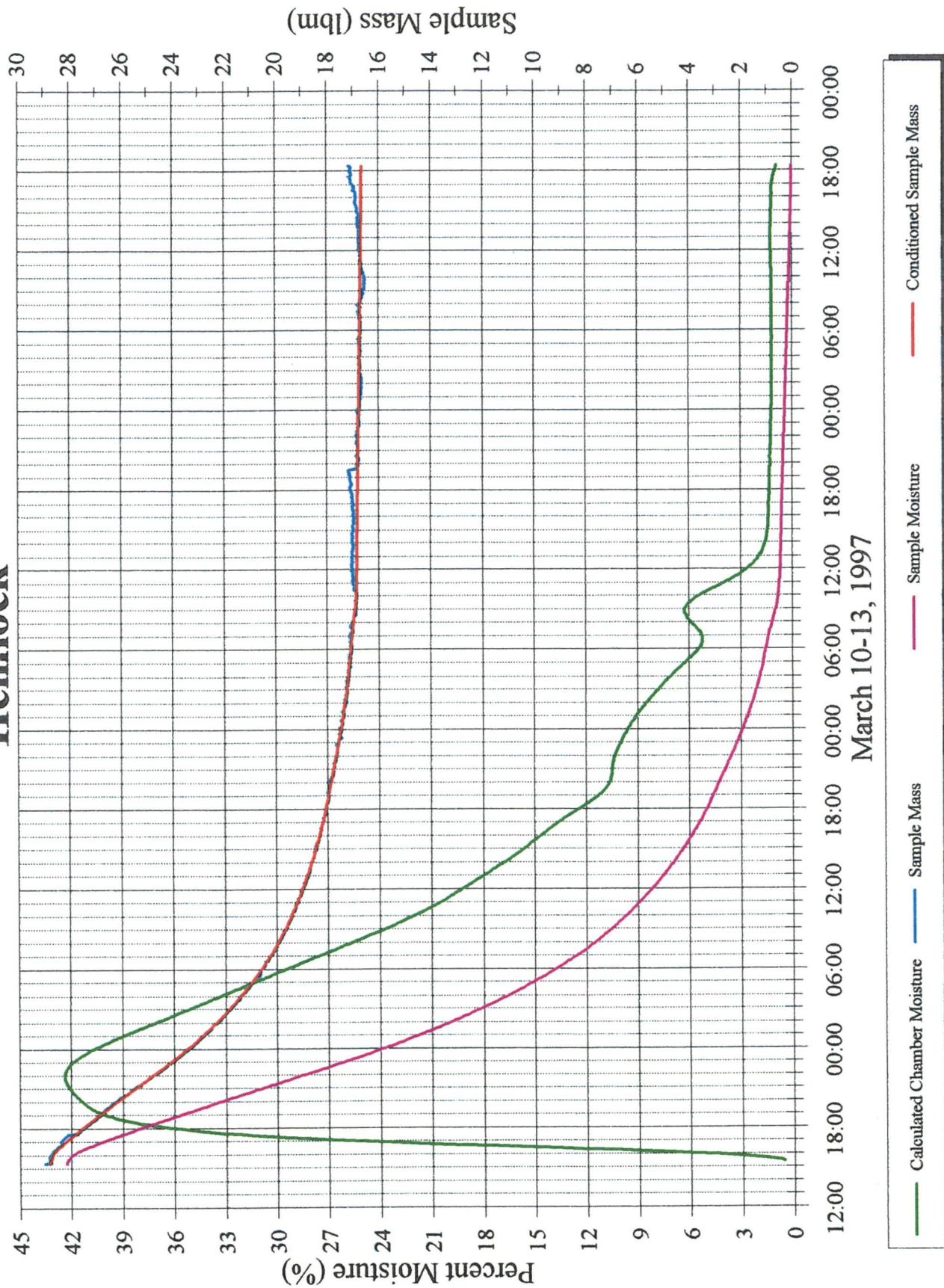
# Cowlitz Stud Trial #1

## Hemlock





# Cowlitz Stud Trial #1 Hemlock





# Standard Dry Gas Meter

Flow Rate CFH

Prof

21

O. 99.7

C. 99.8

36

O. 99.7

C. 99.6

41

O. 99.6

C. 99.6

60

O. 99.7

C. 99.5

75

O. 99.5

C. 99.6

Tested 12/21/95

By Greg Beck

## Free Standing Meter Calibrations

File

fs022796

Method

EPA #5.3.2 & 5.6

Location

Horizon Shop

Meter Box ID

FS-A 685998

Meter ID

None

Calibrated

jdf

Pb=

30.02 (in Hg)

Ta=

60 (oF)

Date

2/14/97

Old

9-23-96

New

02-14-97

Change (+/-)

Y=

0.98537

0.99578

1.06%

FS-A 685998	VAC	dH	Standard	Net	Field	Net	Standard	Meter	Field	Meter	To	Tm	Time		Allowable
FS-A 685998	(inH2O)	(inH2O)	Meter	(ft3)	Meter	(ft3)	Tw	Tw	Tdi	Tdo	(oR)	(oR)	t		Tolerance
FS-A 685998			(ft3)		(ft3)		(oF)	(oR)	(oF)	(oF)			(min)	Y	Y
FS-A 685998															0.020
Initial	N/A	N/A	0.0050	5.9950	892.6050	6.0170	61.0	521.2	61.2	61.2	521.4	521.4	23.133	0.99673	0.001
Final			6.0000		898.6220		61.4		61.6	61.6					
Initial	N/A	N/A	0.0000	6.0000	899.0040	6.0390	61.8	522.1	62.0	62.0	522.0	522.0	10.367	0.99335	0.002
Final			6.0000		905.0430		62.4		62.0	62.0					
Initial	N/A	N/A	0.0000	6.0000	905.0430	6.0130	62.6	522.8	62.2	62.2	522.5	522.5	11.433	0.99727	0.001
Final			6.0000		911.0560		63.0		62.8	62.8					
														0.99578	0.002

File	fs022796														
Method	EPA #5.3.2 & 5.6														
Location	Horizon Shop														
Meter Box ID	FS-D 2713329														
Meter ID	None														
Calibrated	jdf														
							Pb=	30.20 (in Hg)			Old	New	Change		
							Ta=	60 (oF)			09-23-96	02-14-97	(+/-)		
							Date	02/14/97			Y=	0.99517	0.98300	-1.22%	
FS-D 2713329	VAC.	dH	Standard	Net	Field	Net	Standard	Meter	Field	Meter	To	Tm	Time		Allowable
FS-D 2713329	(inH2O)	(inH2O)	Meter	(ft3)	Meter	(ft3)	Tw	Tw	Tdi	Tdo	(oR)	(oR)	t		Tolerance
FS-D 2713329			(ft3)		(ft3)		(oF)	(oR)	(oF)	(oF)			(min)	Y	Y
FS-D 2713329															0.020
Initial	N/A	N/A	0.0000	6.0250	6.6120	6.1730	63.8	524.1	62.6	62.6	522.2	522.2	12.033	0.97249	0.011
Final			6.0250		12.7850		64.4		61.8	61.8					
Initial	N/A	N/A	0.0250	5.9760	12.7850	5.9740	64.4	524.7	61.8	61.8	521.7	521.7	11.467	0.99462	0.012
Final			6.0010		18.7590		65.0		61.6	61.6					
Initial	N/A	N/A	0.0010	6.0000	18.5590	6.0710	65.0	525.3	61.8	61.8	521.9	521.9	13.917	0.98191	0.001
Final			6.0010		24.6300		65.6		62.0	62.0					
														0.98300	0.008

file

fs022796

Method

EPA #5.3.2 & 5.6

Location

Horizon Shop

Meter Box ID

FS-E 2713328

Pb=

30.20 (in Hg)

Meter ID

None

Ta=

60 (oF)

Calibrated

jdf

Date

02/14/97

Old

09-23-96

New

02-14-97

Change

(+/-)

Y=

1.00818

0.99878

-0.93%

FS-E 2713328	VAC	dH	Standard	Net	Field	Net	Standard	Meter	Field	Meter	To	Tm	Time		Allowable
FS-E 2713328	(inH2O)	(inH2O)	Meter	(ft3)	Meter	(ft3)	Tw	Tw	Tdi	Tdo	(oR)	(oR)	t	Y	Tolerance
FS-E 2713328			(ft3)		(ft3)		(oF)	(oR)	(oF)	(oF)			(min)		Y
FS-E 2713328															0.020
Initial	N/A	N/A	0.0010	5.9990	293.7780	5.9200	64.8	524.7	62.6	62.6	522.3	522.3	10.333	1.00871	0.010
Final			6.0000		299.6980		64.6		62.0	62.0					
Initial	N/A	N/A	0.0000	6.0250	299.6980	6.0620	65.2	525.3	62.2	62.2	522.1	522.1	12.167	0.98784	0.011
Final			6.0250		305.7600		65.4		62.0	62.0					
Initial	N/A	N/A	0.0250	5.9750	305.7600	5.9330	65.4	525.8	62.0	62.0	522.0	522.0	15.000	0.99980	0.001
Final			6.0000		311.6930		66.2		62.0	62.0					
														0.99878	0.007



Thermocouple Calibration

Date:	21-Aug-96	Deviation	@60 F	7.8	Pb=	29.95 in Hg	DRB				
Next Calibration:	17-Feb-97	Limit	@212 F	10.1	Ta=	80.0 oF	960820tc				
			@375 F	12.5							
	Probe/ID	Ambient			Boiling, Water			Boiling, Oil			Average
		Standard, F	Measured, F	Difference F	Standard, F	Measured, F	Difference F	Standard, F	Measured, F	Difference F	Difference F
Probe	3-1	38.0	38.6	-0.6	212.4	212.4	0.0	340.8	338.8	2.0	0.47
Probe	3-2	38.0	38.0	0.0	211.6	213.4	-1.8	333.8	332.4	1.4	-0.13
Probe	3-3	37.0	37.8	-0.8	211.6	212.2	-0.6	334.8	332.6	2.2	0.27
Probe	3-4	39.0	39.4	-0.4	211.8	212.2	-0.4	336.2	334.8	1.4	0.20
Probe	4-1	38.0	37.8	0.2	211.2	210.6	0.6	343.8	341.6	2.2	1.00
Probe	4-2	39.0	38.6	0.4	212.2	211.6	0.6	326.4	322.8	3.6	1.53
Probe	4-3	39.0	38.4	0.6	211.4	212.8	-1.4	334.2	331.4	2.8	0.67
Probe	4-4	38.0	37.8	0.2	211.6	212.6	-1.0	335.0	331.6	3.4	0.87
Probe	4-5	38.0	38.6	-0.6	211.6	212	-0.4	347.8	345.6	2.2	0.40
Probe	4-6	38.0	38.8	-0.8	211.6	213.8	-2.2	350.8	348.2	2.6	-0.13
Probe	4-7	37.0	38.4	-1.4	211.4	211.8	-0.4	337.2	341.2	-4.0	-1.93
Probe	5-2	39.0	38.6	0.4	211.8	213.4	-1.6	330.8	330.4	0.4	-0.27
Probe	5-3	38.0	39.4	-1.4	211.8	213.2	-1.4	330.2	328.8	1.4	-0.47
Probe	5-4	38.0	38.0	0.0	212	211.8	0.2	329.2	327.0	2.2	0.80
Probe	5-5	37.0	38.0	-1.0	211.8	212.2	-0.4	328.0	327.8	0.2	-0.40
Probe	5-6	38.0	38.4	-0.4	212	212.8	-0.8	324.4	323.0	1.4	0.07
Probe	5-7	39.0	38.8	0.2	212	213	-1.0	326.2	324.2	2.0	0.40
Probe	5-8	37.0	37.6	-0.6	212	212.8	-0.8	328.0	328.0	0.0	-0.47
Probe	5-9	37.0	39.6	-2.6	211.4	212.8	-1.4	330.4	327.8	2.6	-0.47
Probe	7-1	38.0	38.4	-0.4	212	210	2.0	329.0	326.6	2.4	1.33
Probe	7-2	37.0	37.6	-0.6	211.8	212.8	-1.0	330.6	328.8	1.8	0.07
Probe	7-3	38.5	39.2	-0.7	212.2	211	1.2	327.6	327.6	0.0	0.17
Probe	10-1	39.0	38.8	0.2	212	211	1.0	325.4	324.2	1.2	0.80
Probe	10-2	37.0	38.4	-1.4	212	211.4	0.6	328.0	326.6	1.4	0.20
Probe	10-3	39.0	40.4	-1.4	211.8	211.2	0.6	328.2	326.2	2.0	0.40
Free Standing Pitot	11-S	76.2	78.6	-2.4	196.4	201.2	-4.8	367.4	366.2	1.2	-2.00
	10-S	39.0	38.6	0.4	212	212.6	-0.6	327.0	326.8	0.2	-0.00
	F1	39.0	38.4	0.6	212	212	0.0	341.8	342.2	-0.4	0.07
	F3	39.0	38.4	0.6	212	213.6	-1.6	328.8	327.8	1.0	0.00
	F4	38.0	37.8	0.2	212.6	211	1.6	340.2	338.6	1.6	1.13
	F5	62.8	62.6	0.2	198.2	199	-0.8	368.4	367.8	0.6	-0.00
	F23	37.0	37.0	0.0	212	213	-1.0	341.4	343.6	-2.2	-1.07
	F40	39.0	38.2	0.8	212	212.8	-0.8	355.4	355.6	-0.2	-0.07
	F51	38.0	37.8	0.2	213	213	0.0	330.4	330.2	0.2	0.13
	F83	38.0	38.6	-0.6	212.2	213	-0.8	338.2	338.0	0.2	-0.40
	F84	38.0	38.2	-0.2	210.8	213.6	-2.8	355.6	351.4	4.2	0.40
	F85	38.0	37.6	0.4	212	212.4	-0.4	355.4	352.2	3.2	1.07
	B1	36.0	34.6	1.4	212	213.4	-1.4	367.0	371.2	-4.2	-1.40
	B2	36.0	36.0	0.0	212	210.6	1.4	371.0	366.8	4.2	1.87
	B3	81.0	78.2	2.8	211.8	216.4	-4.6	402.8	401.2	1.6	-0.07
	B4	36.0	36.8	-0.8	212	209.6	2.4	367.0	364.2	2.8	1.47
	B5	81.0	78.8	2.2	212	209.4	2.6	388.6	379.6	9.0	4.60
	B6	36.0	37.2	-1.2	212	211.4	0.6	367.0	369.6	-2.6	-1.07
	B7	37.0	37.6	-0.6	211.6	208.4	3.2	389.6	396.0	-6.4	-1.27
	B8	38.0	34.6	3.4	212	214.6	-2.6	366.8	375.4	-8.6	-2.60
	B9	36.0	35.0	1.0	212	212.8	-0.8	370.0	369.6	0.4	0.20
	B10	38.0	40.2	-2.2	212	209.6	2.4	369.4	370.8	-1.4	-0.40
	B11	38.0	39.8	-1.8	212	208.6	3.4	369.4	362.2	7.2	2.93
	B12	40.8	41.0	-0.2	212	213.2	-1.2	369.0	370.2	-1.2	-0.87
	B13	37.0	33.6	3.4	212	212.6	-0.6	371.4	374.2	-2.8	-0.00
	B14	36.0	36.4	-0.4	212	212.8	-0.8	371.4	371.6	-0.2	-0.47
	B15	38.0	37.0	1.0	212	212.6	-0.6	371.4	370.6	0.8	0.40
AVERAGE		40.7	40.8	-0.1	211.3	211.7	-0.4	347.7	346.8	0.9	0.2
		-0.02%			-0.05%			0.11%			
Hivot Dial Gauges											
	9169	62.2	67	-4.8			0.0			0.0	-4.80
	9142	62.4	64	-1.6	212.2	211	1.2			0.0	-0.20
	D-5	61.6	60	1.6			0.0			0.0	1.60
	D-2	61.6	60	1.6	211.6	210	1.6	365.8	360.0	5.8	3.00
	D-7	61.4	58	3.4			0.0			0.0	3.40
	D-9	61.6	60	1.6	211.6	210	1.6	365.0	362.0	3.0	2.07
	D-12	64.4	70	-5.6	212.4	200	12.4			0.0	3.40
	D-13	64.6	70	-5.4	212.8	212	0.8			0.0	-2.30
	D-14	61.4	58	3.4			0.0			0.0	3.40
Standard Used		Fluke 5895570									

## DRIFT CORRECTION DOCUMENTATION

### EPA Drift Equations:

#### **Method 3a : Oxygen and Carbon Dioxide**

$$C_{gas} = \frac{(C_{ma} - C_{oa}) * (C - C_m)}{(C_m - C_o)} + C_{ma} \quad (\text{Eq. 3a-1})$$

#### **Method 6c : Sulfur Dioxide**

$$C_{gas} = \frac{C_{ma} * (C - C_o)}{(C_m - C_o)} \quad (\text{Eq. 6c-1})$$

;  $C_{oa} = 0$

#### **Method 7e : Nitrogen Oxides**

Section 8, Method 7e; "Follow Section 8 of Method 6C (Eq. 6c-1)."

#### **Method 10 : Carbon Monoxide**

The EPA does not currently address Gas Filter Correlation Instruments therefore there are no current standards.

#### **Method 25a : Total Gaseous Organic Concentration (TGOC)**

This method does not mention correcting for drift although there are established limits.

### Horizon Engineering Drift Correction Equations:

$$C_{gas} = \frac{(C_{id} - Z_x) * (C_{ma} - C_{oa})}{(S_x - Z_x)} \quad S_x = \frac{(C_{mf} - C_{mi}) * (T_x - T_{ci})}{(T_{cf} - T_{ci})} + C_{mi}$$

$$Z_x = \frac{(C_{of} - C_{oi}) * (T_x - T_{ci})}{(T_{cf} - T_{ci})} + C_{oi} \quad T_x = \frac{(T_{te} - T_{ts})}{2} + T_{ts}$$

EPA Definition

Horizon  
Engineering

<b>C<sub>gas</sub></b>	Effluent gas concentration, dry basis	<b>C<sub>gas</sub></b>
<b>C<sub>ma</sub></b>	Actual upscale calibration gas concentration	<b>C<sub>ma</sub></b>
<b>C<sub>oa</sub></b>	Actual zero/low calibration gas concentration	<b>C<sub>oa</sub></b>
<b>C<sub>m</sub></b>	Average of initial and final system upscale calibration bias responses	
	Initial system upscale calibration bias response	<b>C<sub>mi</sub></b>
	Final system upscale calibration bias response	<b>C<sub>mf</sub></b>
<b>C<sub>o</sub></b>	Average of initial and final system zero/low calibration bias responses	
	Initial system zero/low calibration bias response	<b>C<sub>oi</sub></b>
	Final system zero/low calibration bias response	<b>C<sub>of</sub></b>
<b>C</b>	Average gas concentration indicated by gas analyzer, dry basis	<b>C<sub>id</sub></b>
	Starting test time	<b>T<sub>ts</sub></b>
	Ending test time	<b>T<sub>te</sub></b>
	Initial system bias calibration response time	<b>T<sub>ci</sub></b>
	Final system bias calibration response time	<b>T<sub>cf</sub></b>
	Mid-point of test time or gas sampling interval to be analyzed	<b>T<sub>x</sub></b>
	Approximate upscale response at mid-point test time	<b>S<sub>x</sub></b>
	Approximate zero/low response at mid-point test time	<b>Z<sub>x</sub></b>

#### **Notes of Exception:**

- 1] TGOC is first recorded on a wet basis then corrected to a dry basis.
- 2] The TGOC instruments used have some historic data on instrument response to different hydrocarbons. For Propane the response is 1 to 1 molecule while Methane is 1.037 to 1 molecule. We correct for the instruments "over response" to the methane.



# Test Method for Determination of Dry Kiln VOC Emissions

April 5, 1996

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## 1.0 INTRODUCTION

Lumber dry kilns have been identified by the EPA and other environmental agencies as a source of Volatile Organic Compounds (VOCs). The green lumber contains VOCs, which are emitted during the drying process. In order to measure the emissions from dry kilns, it is recommended to apply a test method incorporating EPA Method 25A. However, it is not practical to use the standard EPA Method 25A for dry kilns, because of the following conditions:

- a.) Lumber drying can take over 100 hours to process one load.
- b.) Most dry kilns have multiple vents and often have significant leakage around the loading doors.
- c.) The venting process is periodic. The vents open to release moisture and VOCs in an irregular pattern.

The multiple vent configuration of most dry kilns and the periodic venting makes it difficult to measure the exhaust flow rate. The leakage from doors and other gaps is not measurable and therefore will produce inaccurate results. In addition, tests would need to be repeated for every species of wood the plant dries.

This method applies EPA Method 25A in a controlled environment, where a sample of the lumber is dried in a laboratory dryer and the VOC emissions are measured. The measured quantity of emissions can then be applied to determine accurate emission factors for the actual process by mathematical methods.

## 1.1 PRINCIPAL

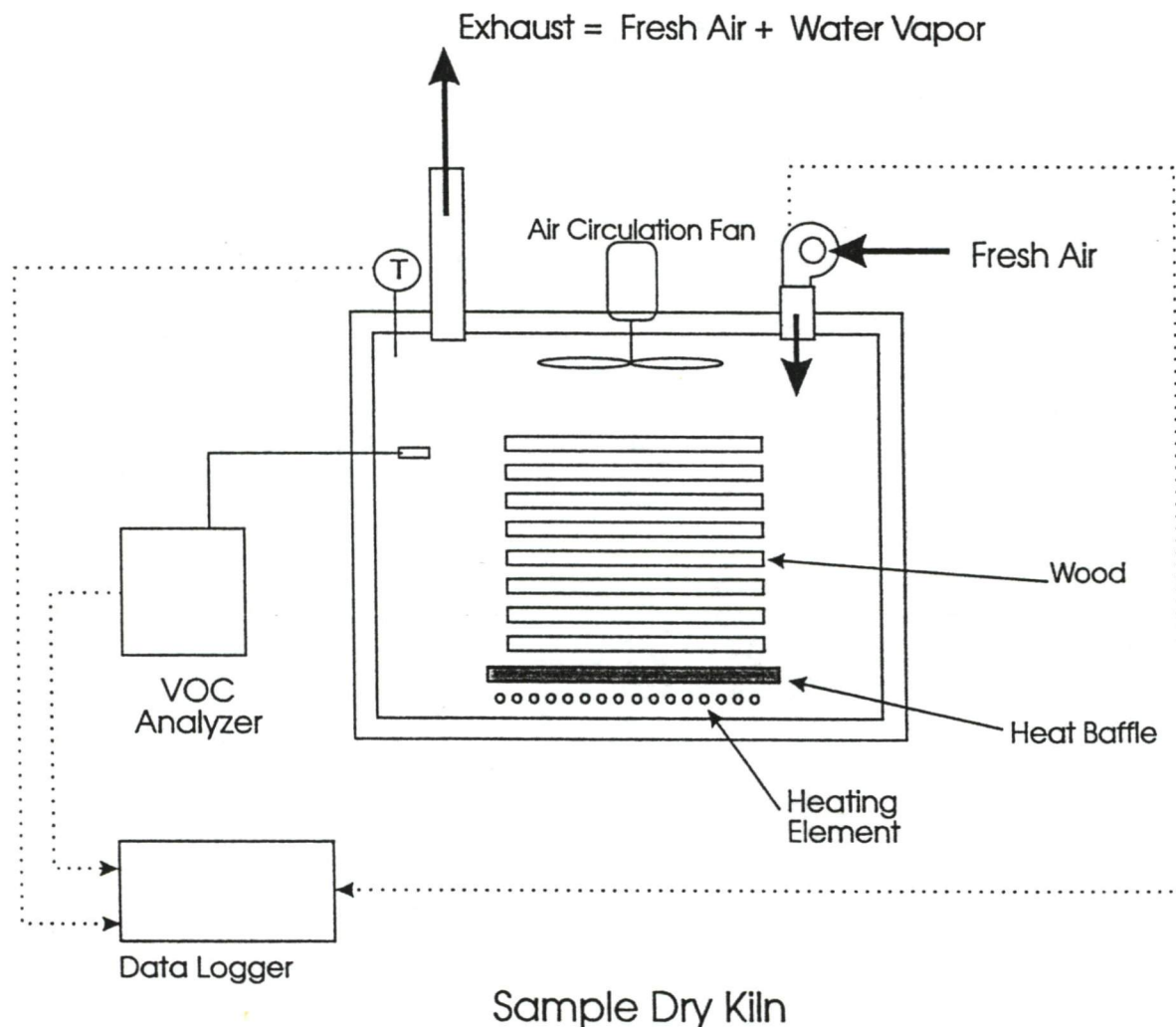
The method for VOC measurements is based on simulated drying conditions in a laboratory size lumber dry kiln that operates in a controlled environment and can dry approximately 10 to 15 board foot of lumber.

The method is considered to be a worst case analysis, where the highest temperature for a typical drying cycle is applied to the sample at all times. The humidity is not controlled. The maximum temperature is to be that used at the actual kiln site. This is normally about 200°F. This method allows any sample drying time, but normal times of approximately 36 to 48 hours will result in a stable (dry) test load weight. Testing times can be extended if the test load weight is not stable. VOC concentrations from the test kiln are not expected to reach zero near the end of the drying cycle.



The VOC analyzer indicates concentration on a wet basis. To correct the concentration to a dry basis it is not necessary to continuously measure the moisture content of the sample stream even though the moisture varies over the drying cycle. An average moisture content for VOC analyzer correction is calculated at any time based on the dry air volume delivered to the kiln and the amount of moisture evaporated from the sample load. As long as the air flow rate to the kiln is greater than that extracted by the analyzer, moisture and air escaping from the oven through cracks are not a problem.

## 1.2 SYSTEM SCHEMATIC





### 1.3 APPLICABILITY AND SENSITIVITY OF RESULTS

From the laboratory test results emission factors can be calculated for a typical drying schedule. Separate emission factors can be calculated for each wood species to any percentage dryness.

### 1.4 TEST KILN APPARATUS

**Test Kiln:** Industrial drying oven, convection type, with sealed doors and openings, of a size sufficient to hold the test load with adequate air circulation space around the test load. The kiln shall be equipped with the following instrumentation:

**Load Cell and Platform:** The entire test load shall be sensed by the load cell on a continuous basis. The load cell suspension system shall be designed to minimize binding. The platform should allow the use of non-organic sticker boards to separate the test load boards in a manner similar to actual drying conditions.

**Temperature Sensor:** A continuous record of the kiln temperature shall be maintained.

**VOC Sample Probe:** A stainless steel or glass probe to gather sample for the analyzer. The probe outlet should be kept to a minimum length and insulated to prevent condensation before the heated sample line connection.

**Air Inlet:** To be placed in a location where the air becomes mixed quickly with oven internal air.

**Air Outlet:** A pressure relief line to allow excess exhaust air to vent. This line should be heated or kept sloped down to prevent accumulation of condensed water vapor that could block the exhaust stream.

**Other Test System Equipment Necessary:**

**Total Hydrocarbon Analyzer System:** Heated total hydrocarbon analyzer and sample line, constructed, operated, and calibrated according to EPA Method 25A.

**Inlet Air System:** A system of providing a constant, measured, hydrocarbon free air to the system. The air should either be dried or have its temperature and humidity measure so that moisture in the stream can be quantified.

Data Logging System: A system to provide a continuous record of the recorded parameters throughout the testing period. Data is to be recorded at intervals no longer than two minutes apart.

## 2.0 SAMPLE COLLECTION PROCEDURE

Depending on the species and on the location of the board within the log, the VOC content will vary. It is recommended that the collected samples represent a cross section of the log from which the board were cut.

Resin rich soft woods often have localized pitch concentration. These so-called pitch pockets can release significantly more VOC than the average board. Sample boards with pitch pockets should not be selected for the test batch of lumber.

Each species of lumber must be tested separately in order to determine species specific VOC release. Therefore all sample boards for a specific test must be of the same species.

The selected boards must be cut into sample boards between 18" and 24" long (all samples boards should be of approximately the same length).

The board thickness and the width of the boards must represent the average dry kiln load.

The samples must be collected immediately after the log is sawed into boards (within 8 hours).

At least 6 separate boards must be used to compile the sample load.

The composite sample load must be at least 10 board foot based on U.S. Lumber Scale.

Each board must be marked with the date of collection, a batch number and a board number (example - Mar 20/96 - 1/3). This means that the piece came from the first of the six selected boards and is the third piece of the same board. It is best to use pencil for marking. Marking pens may add VOCs to the board.

After the sample board are collected, prepare a data sheet with the following information:

- a.) Company Name
- Address
- Telephone Number



### Contact Person

- b.) Date of sample preparation.  
Responsible person collecting the sample.  
Signature of the responsible person.
- c.) Species of the lumber.
- d.) Total number of pieces shipped and the total board feet in the sample batch.
- e.) Dry kiln identification in which this lumber is normally dried.  
Identify more than one kiln, if appropriate.
- f.) Identify each sample piece as shown in the following example:

<u>Sample #</u>	<u>Nominal Size</u>	<u>Length</u>
1/3	8/4" by 6"	18" (plus or minus 1/8")

- g.) Provide the normal drying schedule for this lumber and the maximum drying temperature.
- h.) Provide the final moisture content for this lumber.

Immediately after collecting the samples the entire package of sample boards must be shrink-wrapped or enclosed in a plastic bag and sealed with tape to avoid moisture and VOC loss.

## 2.1 SAMPLE SHIPPING PROCEDURE

The samples should be packaged in a box to avoid damage of the vapor seal during shipping. To ensure arrival at the laboratory within 48 hours of the date the samples were cut and wrapped, select a carrier that can deliver within the specified time.

## 2.2 PREPARATION AND SET-UP BEFORE TESTING

The testing laboratory must be prepared to perform the test within 96 hours after the samples were collected. Samples should be refrigerated in the shipping materials until the testing is started.

The VOC analyzer must be calibrated following EPA Method 25A. The load cell must be calibrated with known weights. The oven should be preheated for several hours at a temperature slightly above the anticipated test maximum to avoid condensation.

After the preparation, place the lumber in the sample dry kiln and start the VOC sampling device. After the drying cycle has been started, the sample kiln door must be latched and may not be opened during the entire drying process.

The lumber in the sample dry kiln must be dried to the maximum temperature at which the lumber is normally dried at the plant site. Test kiln temperature may be increased at intervals, however, to avoid very high humidity in the chamber.

The heating system and internal air circulation system for the dry kiln must be operating continuously during the drying process.

## **2.3 DATA COLLECTION**

During the drying cycle the following information shall be collected and recorded.

- a.) VOC concentration, in ppmvC, inside the sample dry kiln once every two minutes.
- b.) The temperature in the sample dry kiln.
- c.) The in-flow of fresh air into the sample dry kiln in scfh. The flow rate shall not be less than 10 scfh and not more than 100 scfh for every 10 board foot of lumber in the sample kiln. The meter temperature and the relative humidity of the in-flow air should be recorded.
- d.) The weight of the lumber once every two minutes.
- e.) The total drying time in hours and minutes shall be recorded.

## **2.4 TERMINATING THE DRYING CYCLE**

The lumber will be dried until the weight of the wood has become stable to less than +/- 0.25 lb over a 12 hour period. Some variation in weight can be expected due to inlet air humidity changes.

Final calibrations checks should be conducted on the VOC analyzer as outlined in EPA Method 25A. A post check on the weighing system must also be performed.



### 3.0 DATA EVALUATION THEORY

The air in-flow rate and the total air flow data for the entire cycle will be the summarized meter reading in cubic feet. The air in-flow corrected to a dry standard (dscf) will be the same as the out-flow dscf. This will be the volume used in the pounds of VOC calculation.

The water vapor volume will be calculated from the total water loss of the sample plus the water introduced in the in-flow air. From the total water vapor volume and the total dry air volume a percentage moisture can be calculated for any time during the test cycle.

With the results of VOC concentration in ppmvC (wet basis), the percentage moisture, and the volumetric flow in dscf, the total VOC release in lbC can be calculated for any lumber moisture content.

From the result in lb of VOC for the test sample, an emission factor in lb of VOC per 1000 board feet of lumber can be calculated.

### 3.1 EQUATIONS TO DETERMINE EXHAUST FLOW

The actual exhaust flow from the sample dry kiln is the sum of the air flow plus the water vapor flow from the evaporated water in the wood. However, this is not used in the emission factor calculation.

#### a.) Air in-flow in dscf

$$V_{sd} = Y V_m T(\text{std}) P_b \text{ mfg}(2) / P(\text{std}-1) T_m(\text{abs})$$

$V_m$  = meter reading volume in actual cft

$Y$  = gas meter correction factor

$T(\text{std})$  = standard temperature, 527.67°R

$T_m(\text{abs})$  = meter temperature in degree Rankin.

$P_b$  = pressure in inch Hg at test site.

$P(\text{std}-1)$  = standard pressure, 29.92129 inHg

$\text{mfg}(2)$  = mole fraction of dry meter air

#### b.) Mole fraction of dry meter air

$$\text{mfg}(2) = 1 - B_{ws}(2)/100$$

$$Bws(2) = RH Vp / Pb(2)$$

Bws(2) = percent moisture of in-flow air

RH = relative humidity of in-flow air

Vp = vapor pressure of moisture content of in-flow air

Pb(2) = barometric pressure in kPa

**c.) Vapor pressure of moisture content of in-flow air**

$$Vp = \exp(A + B T_m + C/T_m + D/T_m^2)$$

$$A = 18.6866$$

$$B = -0.00243724$$

$$C = -4509.47$$

$$D = -149541.0$$

\*in this equation  $T_m$  is in  $^{\circ}C + 273.15$

### 3.2 EQUATION TO DETERMINE EXHAUST MOISTURE

**a.) Mole fraction of dry gas**

$$mfg(1) = 1 - Bws(1)/100$$

Bws = percent moisture of exhaust

**b.) Percent moisture**

$$Bws(1) = 100 Vw(std) / Vw(std) + Vm(std)$$

Vw(std) = volume of water vapor, scf

Vm(std) = volume of dry gas, scf

**c.) Volume of water vapor**

$$Vw(std) = 0.04707 W / 0.99823 + Vw(std)_{in} + Vw(std)_{initial}$$

W = weight loss of wood, grams

$Vw(std)_{in}$  = volume of water vapor in the in-flow gas, scf

$Vw(std)_{initial}$  = volume of water vapor in over at start of test

### 3.3 VOC CONCENTRATION



**a.) VOC concentration corrected**

$\text{VOC}(\text{cor}) = \text{VOC}(\text{dry})$  corrected for drift per EPA Method 25A

**b.) VOC dry calculation**

$\text{VOC}(\text{dry}) = \text{VOC}(\text{wet}) / \text{mfg}(1)$

$\text{VOC}(\text{wet}) =$  average from analyzer in ppm

$\text{mfg}(1) =$  mole fraction of dry air in oven

### **3.4 TOTAL SAMPLE VOC IN POUNDS**

$\text{Mgas} = \text{VOC}(\text{cor}) \text{ MW Pstd}(2) \text{ Vsd} / 1000000 \text{ R T}(\text{std})$

$\text{VOC}(\text{cor}) =$  ppm dry, corrected for drift

$\text{MW} =$  molecular weight of carbon, 12.01 lbm / lbmol

$\text{Pstd}(2) = 2116.22 \text{ lbf} / \text{ft}^2$

$\text{Vsd} =$  volume of sample (section 3.1)

$\text{R} = 1545.33 \text{ ft lbf} / \text{lbmol } ^\circ\text{R}$

$\text{T}(\text{std}) =$  absolute standard temp., 527.67  $^\circ\text{R}$

### **3.5 VOC EMISSION FACTOR**

It is recommended to express the VOC emission factor is in Lbs. of VOC per 1000 board foot of lumber based on U.S. lumber scale. For other lumber scales the numbers must be corrected.

**a.) Emission factor in Lbs./1000 BF (U.S.)**

$\text{EF} = \text{Mgas} / (\text{BF}_{\text{sample}}) * 1000$  (in Lb / 1000 BF U.S.)

$\text{BF} =$  Total board foot of lumber dried in the sample kiln in U.S. lumber scale.