

# Performance Engineering for Coal-Fired Power Plants (Advanced Coal Quality & Combustion)

by Rod Hatt

## Class Outline

Introduction of Attendees and Instructors

### Review of Basic Coal Quality

It is important to understand the results and meaning of the following coal tests. These are all covered in our “**Basic Coal Quality and Combustion Class**”

#### Proximate

##### Originally

Moisture

Ash

Volatile

Fixed Carbon

##### Short Prox

Sulfur

Heating value (HHV, LHV)

##### Uses

Rank determination - Reactivity

Ash loading

SO<sub>2</sub>

Air Dry moisture

#### Ultimate

Carbon

Hydrogen

Nitrogen

Oxygen

## **Forms of Sulfur**

Look for Pyrite, Sulfate = oxidized

## **Ash Fusion Temperatures**

This is a physical test with variable atmospheric conditions.

Best used as an indication of what combustion conditions do to slag viscosity.

High variability lab to lab.

## **Major and Minor Ash Chemistry**

More consistent lab to lab?

Try to infer slag viscosity and fouling potential.

Elemental loading

ESP Performance

## **Hardgrove Grindability Index**

Variable with low and mid ranked coals due to moisture influences.

## **Equilibrium moisture**

Coals are dustier if they fall below the equilibrium moisture level, even if it is just the surface of large pieces.

## Combustion Tune-up Procedures

### Air Flow and Balancing

Oxygen (O<sub>2</sub>) Readings

Short history of Bailey Combustion Controls

In situ Probes

Verifying O<sub>2</sub> Readings

How much leakage

Induced draft fans

Comparing O<sub>2</sub> to air flow measurements

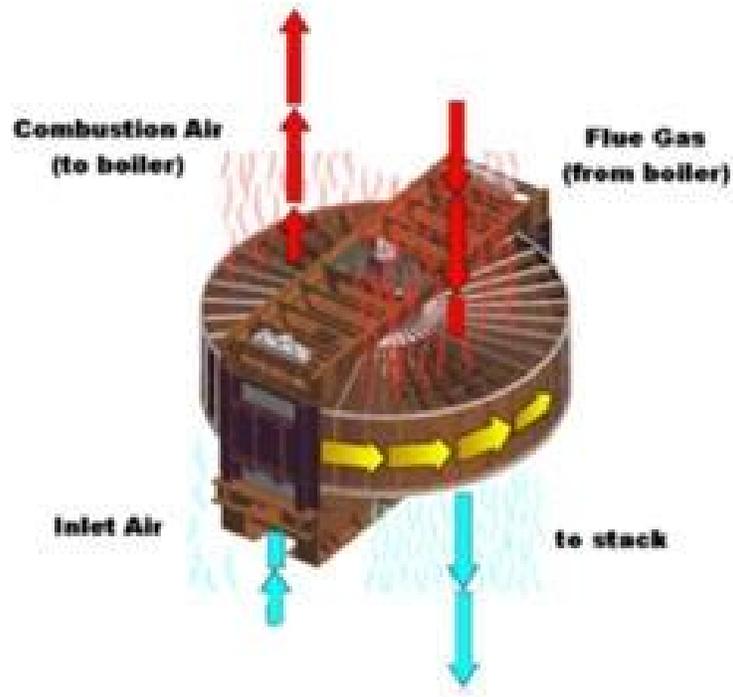
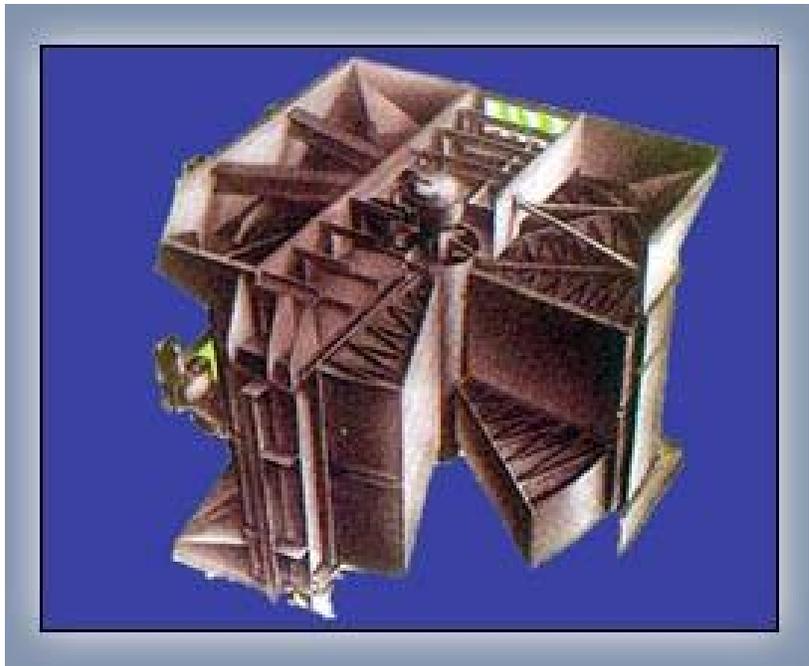
**Excess air =  $20.9/(20.9-O_2)$**

Look up design air flows

Calculate air flow and compare to measured flow

Air Heater Leakage

**=  $90 \times ((O_2 \text{ leaving} - O_2 \text{ Entering}) / (21 - O_2 \text{ Leaving}))$**   
or perform both excess air and take difference.



Other Leakage

Is the leakage before the ESP

## Balancing

O2 levels  
Temperature  
Flows  
Opacity  
NOx

## Pulverizer Performance

### Coal Sizing

Hard work

Sampling is critical

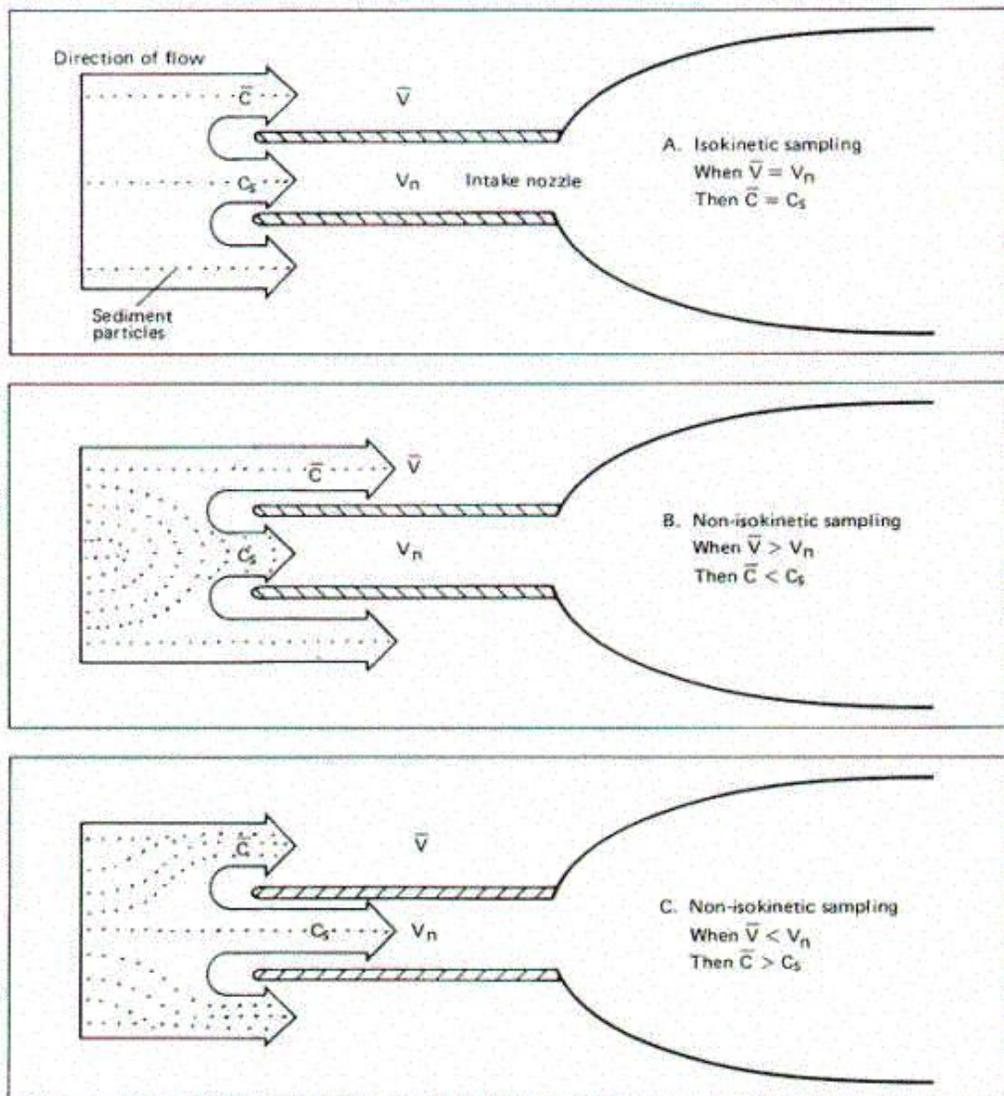
ASTM, ASME, Dick Storm, Rotoprobe

Screening, - worn screens, QCQA

Rosin Rambler

200 mesh for capacity

50 mesh for LOI



Air and Fuel Flow  
Plugged pipes  
Plugged riffles

Clean and Dirty Coal Pipe Measurements  
Error in measurements  
Both important  
Combine with iso-kinetic sampling for mass balance.

Pyrite Rejects  
These are a sample of the rocks and minerals in a coal

**Consider tuning this aspect of mill for coal cleaning**

Primary Air Flow  
Moves coal through pipes  
Minimum velocities to avoid settling out of dust  
Temperature control  
Impacts flame front

Look for coal settling out on air inlet ducts

Adjustments include:  
Roller and journal pressure, alignment, air flow,  
Classifiers, temperature, ball charge, ball size,

### LOI Testing and Combustion Verification

LOI Test  
Loss on Ignition  
Includes drying step

Carbon in Ash  
More expensive

HHV of Ash  
Done by standard additive method

Sampling Location  
ESP Hoppers  
Dust silo  
Ductwork

Sampling Methods  
Bucket  
Pipe  
Corn thief  
Street elbow  
Iso kinetic  
Grid-work of points  
EPA Method 17  
EPA Method 5

Operator Feedback  
Don't wait for test results if it is black  
Calibrated jars  
Take data and discuss operations with operator while  
sample is being collected

CO Analyses  
Use with O<sub>2</sub> and NO<sub>x</sub>

NO<sub>x</sub> Analyses

## Results Engineering

### Boiler Efficiency Testing

Output/Input

Eff. = Heat in Steam/Btus of Coal

Heat Loss Method

Start with 100% and then remove

- Moisture in Air
- Moisture in Fuel
- Moisture from Hydrogen
- Unburnt Carbon
- Heat in Flue gas
- Radiation
- Unknown and fudge

L I N E	<b>Table 12 Combustion Calculations</b> Based on quantities per 10,000 Btu fuel input				L I N E	
1	Fuel— <i>Bituminous Coal, Virginia</i>		<b>Conditions</b>		Date	a
2	Analysis As Fired		by test or specification			
3	Ultimate, % by Wt	Proximate, % by Wt	Total air	%	120	b
4	C 80.31	Moisture 2.90	Air temperature to heater	F	80	c
5	H <sub>2</sub> 4.47	Volatile 22.05	Air temperature from heater	F	350	d
6	S 1.54	Fixed carbon 68.50	Flue gas temperature leaving unit	F	280	e
7	O <sub>2</sub> 2.85	Ash 6.55	H <sub>2</sub> O per lb dry air	lb	0.013	f
8	N <sub>2</sub> 1.38	100.00				g
9	H <sub>2</sub> O 2.90		Unburned fuel loss	%	0.4	h
10	Ash 6.55		Unaccounted loss	%	1.5	i
11	100.00		Radiation loss (ABMA), Fig. 27, Chapter 4	%	0.8	j
12	Btu per lb, as fired, 14,100					k
13	<b>Quantities per 10,000 Btu Fuel Input</b>					13
14	Fuel burned, $100(100 - \text{line } 10) \div \text{line } 12 - .007 \times \text{line } h$		lb		0.66	14
15	Dry air, line b [(value from Fig. 4, Table 11 or Eq. 6) - .08 × line h]		lb		9.11	15
16	H <sub>2</sub> O in air, line 15 × line f = 9.11 × 0.013		lb		0.12	16
17	Wet gas, total, lines (14 + 15 + 16)		lb		9.89	17
18	H <sub>2</sub> O in fuel, $100(8.94 \times \text{line } 5 + \text{line } 9) \div \text{line } 12$ , or Table 11		lb		0.30	18
19	H <sub>2</sub> O in flue gas, total, line 16 + line 18		lb		0.42	19
20	H <sub>2</sub> O in flue gas, total, in percent, (line 19 ÷ line 17) × 100		%		4.31	20
21	Dry gas, total, line 17 - line 19		lb		9.47	21
22	<b>Losses per 10,000 Btu Fuel Input</b>					22
23	Unburned fuel, $10,000 \times \text{line } h \div 100$		Btu		40	23
24	Unaccounted, $10,000 \times \text{line } i \div 100$		Btu		150	24
25	Radiation, $10,000 \times \text{line } j \div 100$		Btu		80	25
26	Latent Heat, H <sub>2</sub> O in fuel, $1040 \times \text{line } 18$		Btu		312	26
27	Sensible heat, flue gas, line 17 × Btu from Fig. 1 @ line e and line 20 = 9.89 × 50		Btu		495	27
28	Total losses, lines (23 + 24 + 25 + 26 + 27)		Btu		1077	28
29	Total losses in percent, (line 28 ÷ 10,000) × 100		%		10.8	29
30	Efficiency, by difference, 100 - line 29		%		89.2	30
31	<b>Quantities per 10,000 Btu Fuel Input</b> Combustion temperature, adiabatic					31
32	Heat input from fuel		Btu		10,000	32
33	Heat input from air, lines (15 + 16) × Btu from Fig. 8 @ line d temp		Btu		612	33
34	Heat input, total, lines 32 + 33		Btu		10,612	34
35	Less latent heat loss, H <sub>2</sub> O in fuel, line 26		Btu		-312	35
36	Heat available, maximum		Btu		10,300	36
37	Less (lines 24 + 25) × 0.5*		Btu		-115	37
38	Heat available, line 36 - line 37		Btu		10,185	38
39	Heat available per lb of flue gas, line 38 ÷ line 17		Btu		1021	39
40	Adiabatic temperature, from Fig. 1 for lines 20 & 39		F		3513	40

\*Note: It is customary to reduce the maximum heat available, line 36, by from 1/3 to 1/2 of the unaccounted plus radiation losses, on the assumption that a portion of these losses occurs in the combustion zone.

## Problem areas

Hard work  
Still need coal and ash samples  
Use theory rather than measure

## Improving Boiler Efficiency

### Improving Combustion

Low LOI  
Low excess air

### Optimizing Combustion

O<sub>2</sub> verse Boiler Efficiency

### Optimizing NO<sub>x</sub>

May require higher CO and or higher LOI  
Small sizing critical

## Unit Heat Rate

### Input/Output

**Heat rate = Btus input / kilowatts output**

### Coal Sampling and Testing Results are Critical

#### Concerns and Opportunities

Piss in the bucket or don't let the sample dry out

Gross and net differ due to the auxiliary power that the plant uses itself

### Performance Diagrams

Feed water heaters and turbine cycle  
ONE SHOT ITERATION

## **Ash Deposits . Slagging verse Fouling**

Slag is Molten, Fouling is Sulfate Salts

TABLE I - Major Causes of Ash Deposits

Fuel Related	<p><b>Large pyrite particles that impact the furnace wall before they completely combust</b></p> <p><b>Clay minerals that contain significant amounts of iron, calcium, sodium or potassium causing them to have low melting temperatures</b></p> <p><b>Interaction of pyrite, clays and alkalis with alumino silicates to form low viscosity melts</b></p> <p>Extremely fine or organically bound alkalis</p>
Equipment Related	<p><b>Soot blowers not in operation or used improperly</b></p> <p><b>Poor pulverization of fuel</b></p> <p><b>Improper air to fuel ratio</b></p> <p><b>Burners damaged or improperly adjusted</b></p> <p><b>Changes in operation of boiler or other equipment</b></p>
Design Related	<p><b>Furnace size too small for fuel</b></p> <p><b>Tube material and/or spacing inadequate</b></p> <p><b>Soot blowing coverage inadequate</b></p> <p><b>No means provided to observe slag buildup</b></p>

Most Slag begins on the Furnace walls and the proceeds up the furnace

## **Basic Analytical Procedures**

### Ash Fusion Temperatures

- Spread between ID and FT
- Spread between Oxidizing and Reducing
- Spread Between Laboratories
- Impact of Combustion
- Cross Plotting between other Coal Characteristics

### Major and Minor Ash Elements

ASTM designates the reporting of ash elements in their fully oxidize form. This is due to the assumption that the elements are found in the ash, and ash is produced in an oxidizing atmosphere at high temperature. These elements may or may not be in the fully oxidizes form in actual power plant ash, and certainly not all in the oxidized form in the original coal. The challenge in interpreting ash analyses is to understand the actual minerals in the coal and the mineral transformations. An additional complication is that some elements can be bound to the actual coal material and are not in a mineral form. This is particularly true for calcium and sodium in low rank coals. The high oxygen content of these coals provides places for these elements to slick to the coal due to ground water movement.

One other element that causes confusion in ASTM analyses is sulfur. It is reported as  $SO_3$  . sulfur trioxide . which is a gas commonly found in flue gas. The  $SO_3$  reported in ASTM analyses is truly sulfur in the sulfite or sulfate form. A reaction takes place during the ashing of the coal between the calcium and the sulfur gases forming calcium sulfate. This reaction takes place at temperatures in the  $1500^{\circ}$  .  $1800^{\circ}$  F. range. The same reaction is used in fluidized bed combustors to capture sulfur using

limestone. The laboratory ashing furnace has a temperature of 1300<sup>o</sup> - 1400<sup>o</sup> F and of course the burning coal raises this locally. The SO<sub>3</sub> reported in coal ash analyses is usually much higher than that found in the fly and bottom ash, due to their formation at much higher temperatures where the calcium and sulfur gases can not react. The main fouling reaction is this same reaction and will be discussed in detail in a following section.

### **Acid Oxides or Glass Formers**

**SiO<sub>2</sub>** . Silicon dioxide - Silica . Represents elemental silicon found in the mineral Quartz, and the aluminosilicate rocks Shale, Slate, and Clays

**Al<sub>2</sub>O<sub>3</sub>** . Aluminum oxide - Alumina . Represents the aluminum found in the aluminosilicate rocks and minerals, not the mineral corundum (Al<sub>2</sub>O<sub>3</sub>) which is an industrial abrasive and also the gems rubies and sapphires

**TiO<sub>2</sub>** . Titanium dioxide . Titania . Seems to always be about one to two percent

### **Basic Oxides or Fluxing agents or Gluing Elements**

**Fe<sub>2</sub>O<sub>3</sub>** . Iron oxide . hematite . Represents the elemental iron in its fully oxidized form. Iron in coal is in pyritic and carbonate forms, iron in ash can be in elemental, reduced and oxidized form. This element is **one of the most important** in influencing slagging behavior

**CaO** . Calcium oxide . lime . represents both mineral (calcite, limestone) and organically bound calcium. The oxidation state of this element is basically fixed to CaO. There is less impact of the oxidizing conditions on the fusion temperatures of high calcium coal ash. This element is also **one of the most important** in slag and fouling deposits

**MgO** . Magnesium oxide . generally lower, (less than six percent) than the calcium, but follows calcium levels as the mineral dolomite.

**K<sub>2</sub>O** . Potassium oxide . typically found in clays and other aluminosilicates

**Na<sub>2</sub>O** . Sodium oxide . found in several forms depending on coal rank and depositional environment, Organically bound form common in low rank coals and NaCl found in coals saturated with sea or briny water. Sodium is also **one of the three important elements** in slag and fouling deposits.

**SO<sub>3</sub>** . Sulfur trioxide . actually in the solid form of sulfates combined with calcium and other basic elements. Not commonly found in high levels in fly and bottom ash. It is found in fouling deposits held together by calcium sulfate.

#### Basic Slagging factors

**B/A** . Base to acid ratio, sum of total bases divided by sum of all acid elements

$$B/A = \frac{Fe_2O_3 + CaO + MgO + K_2O + Na_2O}{SiO_2 + Al_2O_3 + TiO_2}$$

Since most of the basic elements are troublesome the B/A is an indication of the level of fluxing agents.

**Slagging Factor** . Dry sulfur times the B/A provides a result that is similar to an iron squared term  $(Fe_2O_3)^2$ , this is due to  $Fe_2O_3$  being a base and a significant portion of the sulfur being in the pyritic form,  $FeS_2$ .

**Ash and Elemental Loading** . ash and important elements reported in pounds per million Btus. We all ready use \$/MBtu to evaluate the price of coal and report sulfur dioxide  $SO_2$  levels in Lbs./MBtu. These calculations report the ash and elemental

$$\text{Lb. Ash/MBtu} = \% \text{Ash}/(\text{Btu/lb.}/10,000)$$

$$\text{Elemental loading} = \text{Ash Load} \times \text{Element \%}/100$$

Example . Which coal has more iron?

	<u>Coal 1</u>	<u>Coal 2</u>
Ash	=9.0%	14.0%
Btu/lb	=12,300	11,200
Fe <sub>2</sub> O <sub>3</sub>	=12.0%	9.0%

Coal 1

$$\text{Lb. Ash/MBtu} = 9/(12,300/10,000) = 9/1.23 = \mathbf{7.3}$$

$$\text{Lbs Fe}_2\text{O}_3 = 7.3 \times 12.0/100 = 7.3 \times 0.12 = \mathbf{0.88}$$

Coal 2

$$\text{Lb. Ash/MBtu} = 14/(11,200/10,000) = 14/1.12 = \mathbf{12.5}$$

$$\text{Lbs Fe}_2\text{O}_3 = 12.5 \times 9.0/100 = 7.3 \times 0.09 = \mathbf{1.13}$$

**Percent iron that Coal 2 has more than Coal1 =**

$$\mathbf{1.13/.88 = 28 \% \text{ more iron in Coal 2}}$$

This example illustrates the importance of data manipulation rather than just using the straight ASTM levels reported by the laboratories.

## **Advanced Methods for Slag**

### Mineral and Elemental Analyses of Coal

#### Computer Controlled Scanning Electron Microscopy (CCSEM)

Uses Energy Dispersive X-ray Analyses (EDAX) to size and quantify elemental composition.

#### Low Temperature Ashing

Microscopically identify minerals present

#### Chemical Fractionation

Provides information on Organically Bound elements

#### Elemental Analyses of float sink and/or size fractions

Can be performed by ASTM coal laboratories

Pounds of iron per million Btu

Pounds calcium, sodium, and other elements

### Ash Deposit Analyses

#### Sampling Procedure for Ash Deposits

#### Polarized Light and Scanning Electron Microscopy

Look for the glue holding the material together  
Look for signs of combustion conditions

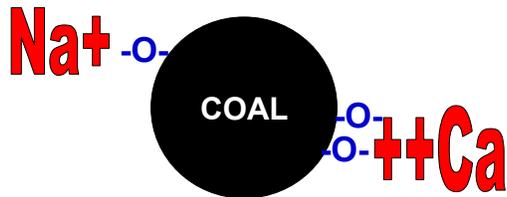
#### Forms of Iron by chemical fractionation

#### Mossbauer spectroscopy

## Fouling Deposit Formation

Organic calcium and sodium elements vaporize during combustion

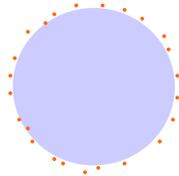
### Organically Bound Alkalis



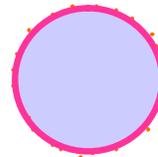
The calcium reacts with sulfur gases to form calcium sulfate.

Sodium can condense onto aluminosilicate flyash particles and melt the surface

### Sodium Condenses on Surface



### Causing a Molten Layer on Surface



## Electrostatic Precipitators

### Basic Operation Principles

High Voltage DC . Transformer Rectifier Sets

Voltage Control . Controlling the wave form and sparking rate

Sizing ESP . Plate area (fixed) and Flue gas volume  
(operational variable?)

Migration Velocity or why the particle goes to the plate

Ash Resistivity concerns

Gas Velocity . impacts ESP size and treatment time

Rapping . important equipment that can be high maintenance

Hopper Evacuation . useful for determining ash partitioning

Dust Collection and Storage

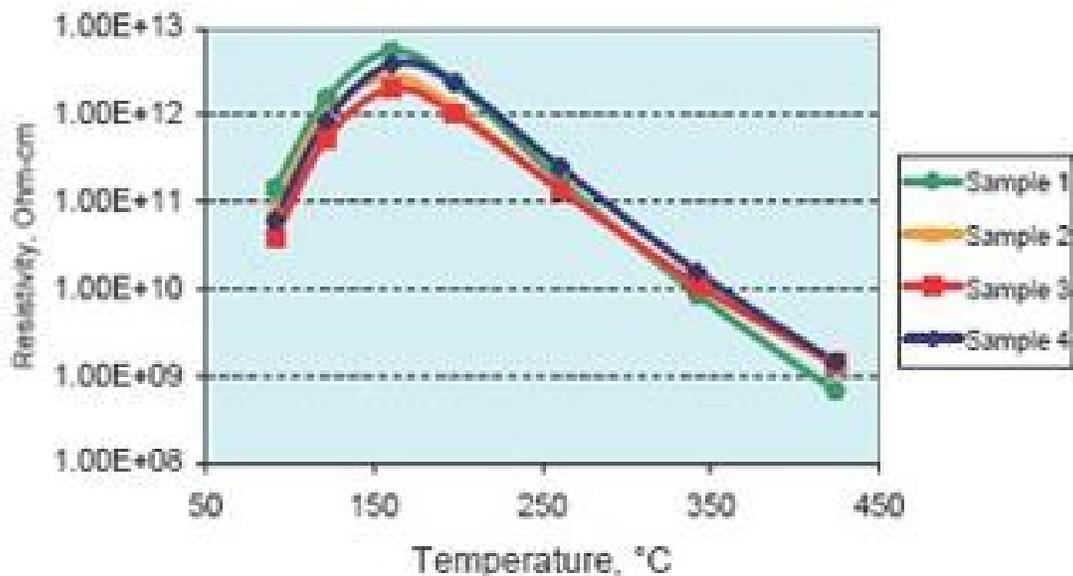


Figure 3: Variation of Resistivity vs Temperature of typical Indian fly ash samples.

## Advanced ESP Methods and Tune-up

### Combustion Influences

ESP is meant to collect ash, not carbon from incomplete combustion

Resistivity, volume of material, and potential fires all point towards spending time on combustion

Air in leakage and air heater leakage can significantly impact ESP performance

Balancing furnace and flue gas flows

What do you balance, flow, temperature, excess air?

### Calculating and Measuring Fly Ash Resistivity

Calculation Methods

Laboratory Measurements

In situ Measurements

Cup Collection Method

Point to Plane Resistivity Probe

Correlating and making sense of resistivity data

### Rappers

Adjustment and different thoughts on rapping sequencing

Inspections and Maintenance

Measuring rapper force

High Voltage controls

Spark rate theories

Ramp rate, hold and setback settings

Measuring gas flows and particular loadings

EPA Methods 5 and 17

Hotwire anemometer

ESP Inspections

Learn form your inspector

Mechanical and Electrical

Tune-up Procedures

Good Combustion

Furnace conditions and temperatures

Air Leakage

Rapper Performance

Balancing

Voltage controls