

Cooling Water Lime Softening Plant Optimisation

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Abstract—Recirculating cooling water is required in a power plant to condense the exhaust steam from the turbine and extract the latent heat as part of the Rankine cycle steam process. Conditioning of the recirculating cooling water is essential to maintain the cooling system free of scaling, fouling and corroding the heat transfer surfaces. A good conditioning treatment for cooling water chemistry control simultaneously reduces scaling, biological fouling, deposition and corrosion on the cooling systems. The current interventions for the control of these concerns has increased the costs accompanying the operation of a cooling water circuit. This lead to the power plant experiencing cooling changes and eventually affected plant performance. A decision by the power plant to optimise the side stream treatment control was initiated. Plant analysis, evaluation was conducted, lime system to the sedimentation plant including other chemicals for the clarifications process were optimised. Finally, a model was developed to assist in plant monitoring and data collection.

The assessment conducted before the optimisation revealed that cooling water flow rates to the plant were inconstant leading to incorrect chemicals being dosed. This resulted in the sedimentation process being ineffective and overall chemistry performance deteriorating. To address the above, flow measurements were verified and chemical dosages using the jar test and lime silo bag test to the plant were conducted. The corrected measurements were incorporated in to the model, giving an indication to the operating team to make corrections and adjustment to the lime side stream treatment keeping the operation of the plant at an optimal level. During the optimisation a pH increase was observed with clarifier 2 responding quicker than clarifier 1 where the test was conducted.

Optimisation of the cooling water side treatment to control cooling water chemistry will have a positive impact on the power plant performance especially the cooling systems in a power plant.

Keywords—Cooling water chemistry conditioning, lime softening, optimisation, side stream treatment

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I. BACKGROUND

Lime plant optimisation project was introduced as a result of plant failures and downtime occurring at the lime handling plant. Lime is dosed in the concentrated cooling water clarifiers to control the alkalinity and hardness of the concentrated cooling water. The unavailability of the lime handling and preparation plant for longer durations is not a desired practice. Lime plant unavailability compromises the long-term plant health in the concentrated recirculating cooling water system. Maintenance and operational challenges are the reason for lime handling plant outages. These challenges are associated with frequent blockages in the lime silo, chemicals and spares unavailability amongst other things.

II. INTRODUCTION

Process integration is a systematic technique for reducing water consumption and wastewater generation [1]. This technique involves water regeneration, recycling and reuse [2], with the consideration of chemical concentrations, water flowrates, pH, conductivity, organic carbons and others that may prevent water reuse or recycling within the water network [3].

Chemical treatment is the most common method used to regulate the cooling water chemistry in order to prevent deterioration in plant performance [4-8]. A lime plant is commonly used for cooling water pre-treatment in wet cooled power plants [9]. In the lime softening process chemicals are added to the cooling water so that the scale forming components are reduced [10]. The lime is used to remove alkalinity and hardness salts through precipitation of calcium carbonate and magnesium hydroxide in the clarifiers [11].

Lime reacts with mainly calcium, magnesium and bicarbonates to form scale in the clarifiers, which is then removed by desludging [12]. These reactions take place at about pH 10 [13]. If insufficient lime is added (pH lower than 10), the reaction will be incomplete and not all the bicarbonate (HCO_3^-) will be removed. If lime is over-dosed (pH higher than 10), then the excess will be carried into the cooling water system where the scale forming reactions will occur [13]. Precipitation of heavy metals out of solution is achieved at optimum pH of 10.2 (McDonald, 2006) where almost all the bicarbonates in a water solution are converted to carbonates (CO_3^{2-}). Hanekom (2008) indicated in his study that the scale that forms in the cooling water system will generally scale the warmest point in the system i.e.

the condenser [14].

III. MATERIALS AND METHODS

A. The chemistry

Alkalinity and pH measurement were used to evaluate the performance of the plant. Alkalinity (P, M and OH), defined as the acid absorbing property of water [15] is related to the optimum pH. P and M refer to phenolphthalein and methyl orange indicators respectively whereas OH measures the hydroxides. P and M alkalinity relationship was used to assess the performance of the plant as indicated by Table I. The optimum pH is achieved when $2P=M$ where only carbonates are available in solution [13].

TABLE I: ALKALINITY RELATIONSHIP

Titration results	OH alkalinity	CO ₃ ²⁻ Alkalinity	HCO ₃ ⁻ Alkalinity
P-Alk	0	0	=Total Alk
P-Alk =Total Alk	=Total Alk	0	0
P-Alk is <1/2 Total Alk	0	2*P-Alk	=Total Alk -(2*P-Alk)
P-Alk is 1/2 Total Alk	0	=Total Alk (Optimum pH)	0
P-Alk is >1/2 Total Alk	(2*P-Alk)-Total Alk	=(Total Alk -P-Alk)*2	0

B. Dosing system

Lime is stored in a lime silo at the water treatment plant. The mass of the lime delivered to the plant from the silo during the injection process is unknown due to the level transmitter that is faulty and measuring inaccurate levels. This resulted in the unexpected depletion of the lime. The commercial process is prolonged, leading to operating the plant without any lime dosages. To address the challenge during optimisation, lime silo bag tests were conducted. The tests physically measured the mass of lime that is collected in a bag at fixed time. Table II shows a typical average mass of lime recorded from the lime silo at different percentage openings.

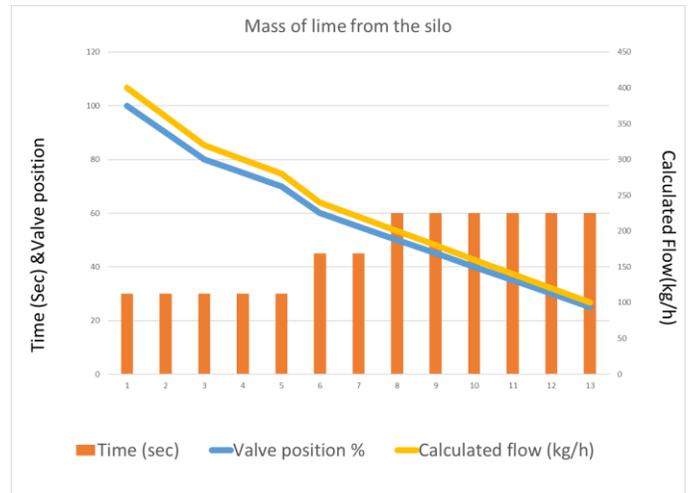


Fig.1: mass of lime from the silo

The data logged in Fig.1 were used to measure the lime dosed to the plant and predict the low levels in the silo. As lime levels drop in the silo, the commercial process can commence to ensure sufficient time for the chemical order to be placed. Chemical unavailability and lime shortages will be reduced if not completely prevented.

C. Chemical dosages

Concentrated cooling water (CCW) and dirty water dam (DWD) recovery water samples were collected from the power plant. Lime was dosed with a blend of a flocculent (poly) to enhance the clarification and sedimentation process [16]. Jar test [17] analysis recorded in Fig 2 and 3 was conducted in the laboratory to obtain the optimum dosage ratio between lime and the flocculent. Recirculating cooling water flow was kept constant, whereas different concentration of lime and flocculent (poly) were varied to observe the optimum dosage, for turbidity removal.

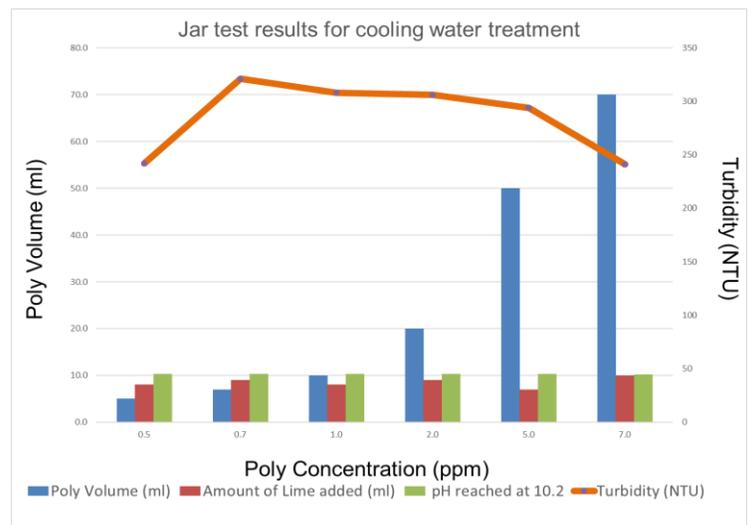


Fig.2: lime and poly Jar test measurements for cw treatment

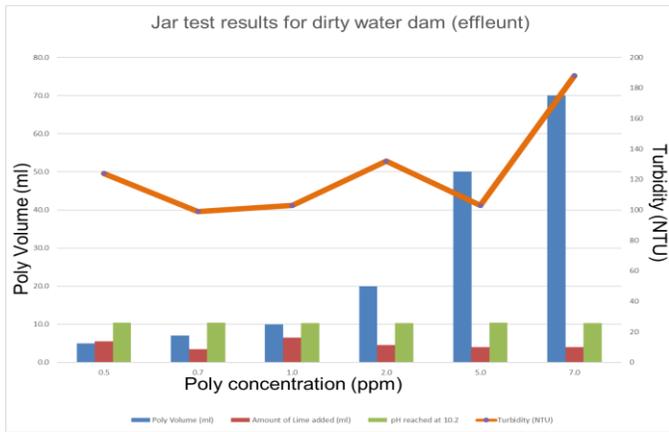


Fig.3: lime and poly Jar test measurements for dwd

The laboratory jar test [18] was used to determine the amount of chemicals that must be dosed to meet the optimum pH of 10.2 in the plant.

D. Plant flowrates

A mass balance of the plant was conducted before the plant can be optimised. The information available from the power plant initial design phase has transformed as new technology immersed, therefore plant current conditions needed to be verified and updated. In addition, the ageing power plants do not have adequate flow meters at critical parts of the plant, and some are obsolete and redundant. A portable flow transmitter [15] was used to measure and acquire the correct flow rates around the plant. Fig.3 shows the flow and mass balance around the concentrated cooling water system.

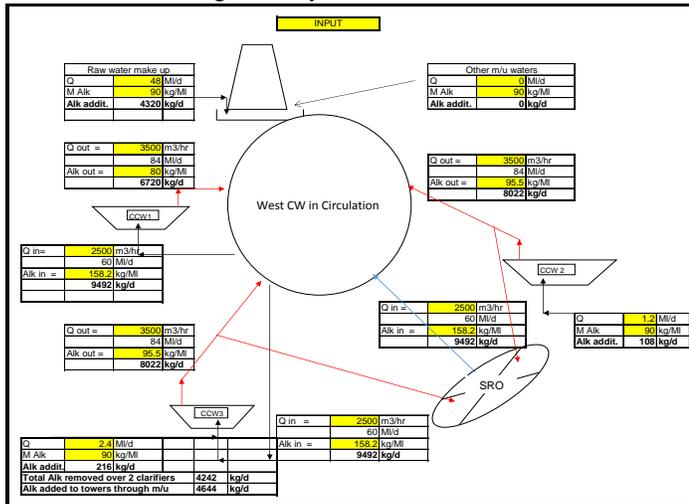


Fig.4: ccw circuit with side stream water balance

E. Data modeling

The data collected from the plant and results from the laboratory were modelled to ensure the optimum management and control of concentrated cooling water treatment. Table IV focuses on the clarifier flow input. The design CCW flow rate is 10% of the CCW system. The required CCW to be treated is controlled by populating the main feed flowrate in the model. A

previous study indicated that circular clarifier optimum flow through velocity should be between 0.7m/s and 1.4 m/s . The calculated figure on the model should be monitored and verified.

TABLE IV: CCW CLARIFIER FEED FLOW

Conditions		Key: Legends	
Date		Operator Inputs	
Time		Model outputs	
		Operator to check	
Clarifier Identity	1	SELECT	
No. of Clarifiers per 10% CW	1	SELECT	
Recommended Lime dose	50	ppm	
Recommended Poly dose	1	ppm	

Clarifier Inlet Water		
Main CW feed flow(10%)	2013.06	m ³ /h
Total Clarifier Feed	2013.06	m ³ /h
Overflow Rate(OFR)	30.38	m ³ /m ² .d
Flow-through Velocity	1.27	m/h
Upflow velocity	0.0004	m/s
Retention in Clarifier	4.74	h

Table V and VI extracted from the model indicate the chemical dosage required for the plant during optimisation as per the jar test laboratory results.

TABLE V: CCW CLARIFIER LIME DOSING RATE

Lime dosing		Lime Dosage
Lime type	Hydrated lime	
Concentration	64% Ca(OH) ₂	
Rotary feed opening	8% 100%=1250 kg/h	
Feed, Ca(OH) ₂	66 kg/h	
Motive water feed	5049 l/h	
Slurry concentration	13.0286 g/l	
Slurry concentration in ppm	13028.6 mg/l	
Dilution water supply	21091 l/h	
Concentration in dilution tank	5033 mg/l	
Clarifier feed valve position	20% 100%=100 m ³ /h	
Lime Clarifier Feed	20000 l/h	
Concentration per clarifier	50.0 ppm	

The dosing of lime is controlled by using a pH transmitter located on the outlet of each clarifier to maintain the pH levels between 10.2 and 10.4 [19]. M-alkalinity is used as a measure of efficiency in the clarifier and an indication of the concentrations levels of calcium bicarbonate salts in the CCW.

TABLE VI: CCW CLARIFIER POLY DOSING RATE

Poly/Coagulant Aid		
Poly type	Polvelectrolyte	
Chemical name	Sudfloc TM	
Concentration	100%	ppm
Pump dosing	7%	100%=1000l/h
Feed flow	68.4	l/h
Poly Makeup Water		l/h
Poly Dilution Water	9798	l/h
Poly Clarifier Feed	9471	l/h
Clarifier Poly Dosage	34.0	ppm

Manual Calibration	17%
	8.5
	4.222
Poly Dosage	

The requirement for the lime and poly dosage is calculated according to the dilution chemistry Equation 1. Dosing rates for lime and poly calculations are shown in Table V and VI on the model.

$$C_1V_1 = C_2V_2 \tag{Eq. 1}$$

Where:

C_1 = original concentration of the solution, before it is watered down or diluted.

C_2 = final concentration of the solution, after dilution.

V_1 = volume about to be diluted

V_2 = final volume after dilution

Two concentrated cooling water clarifiers were considered for the optimisation test, clarifier 1 and 2. Lime was mixed in to the mixing tank from the silo and transferred as slurry in to the dilution tank. From the dilution tank lime was pumped to clarifier 1 and 2 center cone where it is mixed with 10% concentrated cooling water. Samples were collected from the center cone and the clarifier launder for performance management.

IV. RESULTS AND DISCUSSIONS

The lime side treatment softening in a power plant is designed to treat 10% of the stations concentrated cooling water to ensure that the cooling water meets chemistry specification standards after treatment [10]. As the makeup water chemistry changes and other water management regulations take effect, the chemistry changes gradually. This resulted in ineffective treating of the CCW. The lime optimisation trial conducted at the power plant intended to improve and advice on the treatment of the side stream softening. The study was conducted between September to December 2019. A portable pH meter was used in the plant to assist in making lime dosing adjustment until the optimum pH is achieved. Fig 5 shows the behavior from clarifier 1 center cone pH after the optimisation process commenced. The measured pH from the beginning of the trial was below the optimum pH although in the afternoon an increase in pH was observed. The same trend to confirm clarifier1 behavior was observed from Fig 6. Clarifier 1 treated water pH slowly increased. In Fig 7 the pH increased to above the optimum pH, the same pattern was observed as in Fig 8. This was a clear indication that the lime slurry was slowly going through and the observed change was in less than 2 hours.

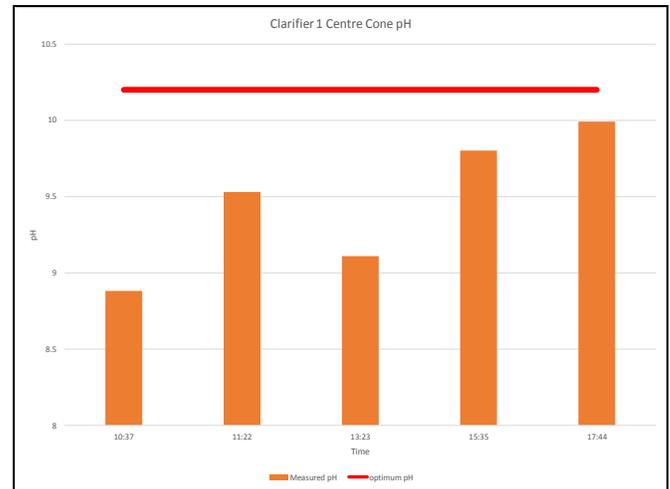


Fig.5: clarifier 1 center cone pH

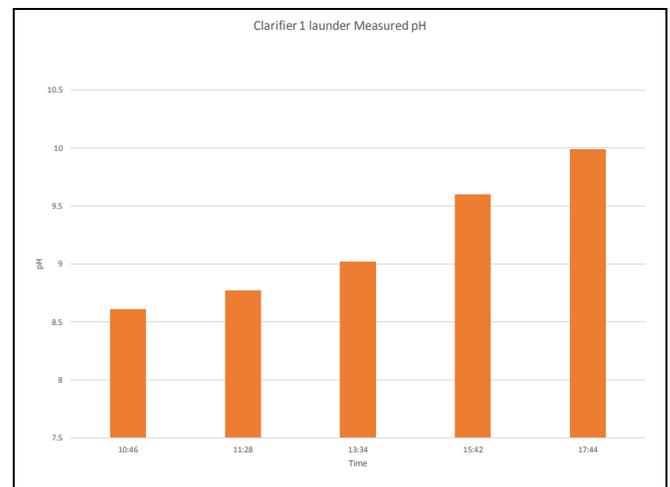


Fig.6: clarifier 1 treated water pH

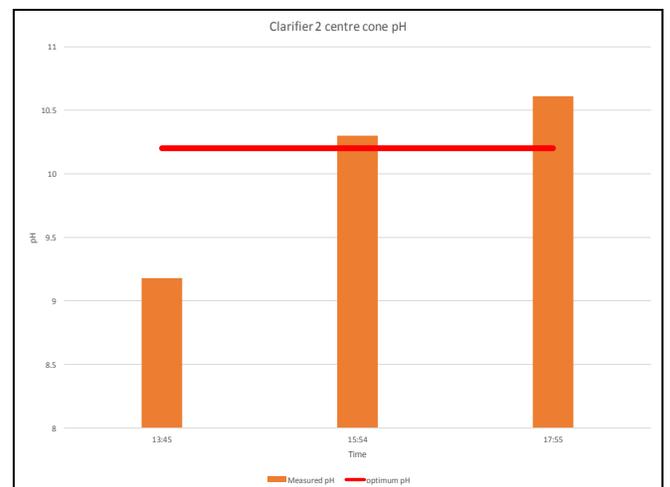


Fig.7: clarifier 2 center cone pH

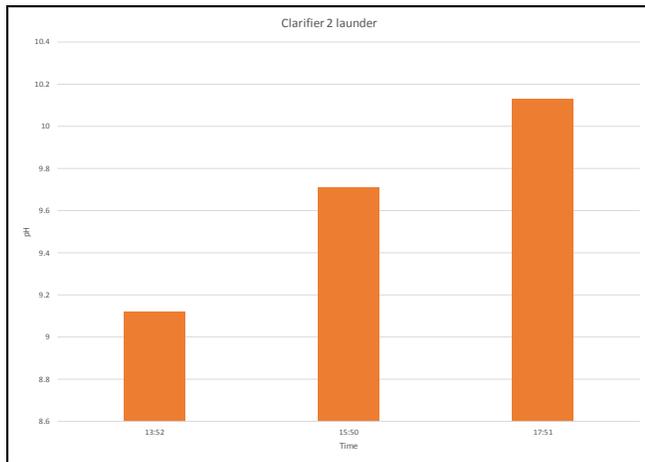


Fig.8: clarifier 1 treated water pH

Two sets of samples were taken to the laboratory for P and M alkalinity analysis. Table VII shows the optimisation results from the laboratory. Though the laboratory results do not indicate an achieved optimum pH from clarifier 2, the chemistry performance for clarifier 2 is showing an improvement. An increase in pH to just above 10 can be seen from clarifier 2 center cone. This confirms the trend observed in Fig 7. The expectation was for the alkalinity relationship to be $2P=M$, and it is unfortunate that during the test, optimum conditions were not met. This was due to the challenges experience in the power plant and the test had to be placed on hold.

TABLE VII: OPTIMISATION LABORATORY RESULTS

Optimisation laboratory results					
First set					
Sample Point	pH	P-alk (ppm)	M alk	2P	Alkalinity Relationship
Clarifier 1					
Centre cone	9.217	30.8	212.5	61.6	$2P < M$
Clarifier 2					
Centre cone	9.17	31.3	249	62.6	$2P < M$
Launder 1	9.214	32.8	208.9	65.6	$2P < M$
Second set					
Clarifier 1					
Centre cone	9.659	47	173.7	94	$2P < M$
Clarifier 2					
Centre cone	10.056	72.2	219.2	144.4	$2P < M$
Launder 1	9.514	46.7	198.9	93.4	$2P < M$
Launder 2	9.634	54.4	226.4	108.8	$2P < M$

V. CONCLUSIONS

The results from the optimisation trial show that lime was under dosed. Clarifier 1 and 2 average pH results on the center cone are 9.4 and 10 respectively. The steady increase in pH does indicates that lime was being injected in to the clarifiers. From the results it is evident that the optimum pH was never reached. This is due to the limited time that was given to conduct the trial. Great results could have been obtained, however serious challenges were experienced with regards to plant, chemicals and spares availability.

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