# PC-375 Cyanide Bright Zinc Process

# **Properties**

- Liquid brightener
- Produces brilliant deposits at low, medium, and high current densities
- For both, barrel and rack operations
- Excellent throwing and covering power
- Plated parts can easily be chromated
- Due to the variable metal and cyanide content of the solution, it may be used over a wide range of operating conditions
- Its brilliant deposits and its economic use make it the outstanding cyanide zinc brightener system

# **Application**

### Make-up concentrations:

Zinc oxide 44 g/l Sodium cyanide 95 g/l Sodium hydroxide 75 g/l Brightener PC-375R 4 ml/l Purifier 3 ml/l

### **Desired values:**

 Zinc Metal
 35 g/l (30-40 g/l)

 Sodium cyanide
 95 g/l (75-110 g/l)

 Sodium hydroxide
 75 g/l (60-80 g/l)

 Sodium carbonate
 max. 70 g/l

 Ratio NaCN:Zn
 3.0 (2.5-3.5)

**Make-up**: Dissolve sodium hydroxide and sodium cyanide in 1/3 of the required amount of water; considerable heat will be evolved. Add zinc oxide and stirr until dissolved. Cool to room temperature. Add the purifier, fill the tank with water to its final volume and filtrate continuously overnight. Add the required amount of PC-375R and stirr the electrolyte.

Temperature: 25 °C (20-45 °C)

(at lower temperatures current densities should be lower, at higher temperatures brightness and throwing power is reduced)

Cathodic current density:

Barrel 0.2-2.0 A/dm2 Rack 0.2-4.0 A/dm2 **Current efficiency**: 60-80 %

**Deposition rate**: 10-14 μm per Ah/dm2

Anodes: pure zinc 99.99 % according to DIN 1706 or steel anodes to

control the zinc concentration

Agitation: cathode agitation (rack or barrel movement) recommended, no

air agitation (to avoid cyanide mist and carbonate increase) Tank material: plastic or steel with plastic coating

Filtration: recommended Heating: not necessary

necessary for lines with high load on small volumes and/or Cooling:

recommended to freeze out sodium carbonate

Exhaustor: required for worker's protection

Maintenance: Analyse zinc, sodium cyanide, sodium hydroxide and sodium carbonate regulary. Zinc is controlled by changing the anodic current density or by using steel anodes. Add sodium cyanide and sodium hydroxide according analysis. Add 0.2 I purifier per 10 kg added NaOH. Freeze out excess sodium carbonate. Adjust PC-375R with the aid of Hull cell tests.

**Consumption**: PC-375R is consumed by drag-out as well as electrochemically, by anodic oxidation and cathodic build-in

The following values can give a range for the consumption (per 10,000 Ah):

PC-375R 1-3 I

# **Effects of the Electrolyte Components**

#### Zinc

Increase of zinc increases the burning limit, but reduces the throwing power. A lack of zinc produces burnings.

### Sodium cyanide

Excess cyanide reduces brightness, thus more PC-375R is needed. Lack of cyanide produces a more brittle zinc layer. The high cyanide electrolyte type is less sensitive against impurities.

### **Sodium hydroxide**

Excess hydroxide speeds up zinc dissolution too much. Lack of hydroxide reduces the burning limit, and leads to passive anodes.

#### Sodium carbonate

Excess carbonate reduces brightness, thus more PC-375R is needed. Further, it is responsible for passive anodes leading to a bad current distribution in the electrolyte and a too low zinc dissolution rate.

### **Contaminating metals**

like Cu, Pb, Cd, Sn, Ni (...) deteriorate brightness and appearance of the zinc layer and should be removed with the purifier. Chromium(VI) reduces current efficiency and coverage in the low current density area. It affects the chromatability and appearance and must be reduced to Cr(III) with sodium dithionite.

#### Additive PC-375R

Excess of PC-375R causes a spotted dull zinc deposition in the low to medium current density area, current efficiency is reduced and in extreme cases, blistering may occur. Lack of additive results into lack of brightness and throwing power.

# **Analysis Cyanide Zinc**

### **Sample Preparation**

Take the sample at a homogeneously mixed position and let it cool down to room temperature. If dull, allow to settle and decant or filtrate.

## <u>Zinc</u>

**Reagents**: 0.1 N EDTA, buffering solution (100 g/l NaOH + 240 ml/l 98 % acetic acid), indicator 1 % xylenol orange blended with KNO3

**Process**: Pipette 5 ml into a 250 ml Erlenmeyer flask, add 100 ml demineralized water, 20 ml buffering solution and a spatula tip of indicator and titrate with 0.1 N EDTA from red to yellow.

**Calculation:** consumption in ml  $\cdot$  1.3074 = g/l zinc

### Sodium cyanide

**Reagents**: 0.1 N silver nitrate solution, 10% NaOH solution, 2 % potassium iodide solution

**Process**: Pipette 5 ml into a 250 ml Erlenmeyer flask, add 100 ml demineralized water, 10 ml 10 % NaOH solution, 6 drops potassium iodide solution and titrate with 0.1 N silver nitrate solution until lasting turbidity.

**Calculation**: consumption in ml  $\cdot$  1.98 = g/l sodium cyanide

### **Sodium hydroxide**

Reagents: 1 N sulfuric acid, indicator sat. alcoholic solution of tropaeolin 0

**Process**: Pipette 5 ml into a 250 ml Erlenmeyer flask, add 100 ml demineralized water, 5 drops indicator and titrate with 1 N sulfuric acid from orange to light yellow.

Calculation: consumption in ml  $\cdot$  7.98 = g/l sodium hydroxide

### **Sodium carbonate**

Reagents: 5 % barium nitrate solution, 1 N hydrochloric acid, 1 N sodium hydroxide solution, indicator 0.04 % methyl orange solution

**Process**: Pipette 10 ml into a 250 ml Erlenmeyer flask, add 50 ml demin. water and boil the solution. Add 75 ml barium nitrate solution. After settle down of the precipitate, filtrate with a fine grained filter paper and wash with hot demin. water. Put the filter into a 250 ml Erlenmeyer flask, add 100 ml demin. water, 30 ml 1 N hydrochloric acid and boil the solution shortly. After cooling down, add 3 drops indicator and titrate excess hydrochloric acid with 1 N sodium hydroxide from red to orange-yellow.

**Calculation**:  $(30 - \text{consumption in ml}) \cdot 5.3 = g/l \text{ sodium carbonate}$ 

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