

# RIAG Ni 122

## Bright Nickel barrel plating process

The **RIAG Ni 122** barrel nickel electroplating process offers the following advantages:

### Properties

- Excellent brightness
- Light coloured white deposits
- Excellent metal deposit thickness distribution
- Good ductility
- Excellent brightness throwing power
- Ideal for zinc die-casting plating

### Make up

	Range	Optimum
Nickel sulphate (NiSO <sub>4</sub> x 6 H <sub>2</sub> O)	180 – 250 g/L	200 g/L
Nickel chloride (NiCl <sub>2</sub> x 6 H <sub>2</sub> O)	50 – 100 g/L	60 g/L
Boric acid (H <sub>3</sub> BO <sub>3</sub> )	40 – 50 g/L	45 g/L
<b>RIAG Ni 132 Make up</b>	10 – 15 mL/L	12 mL/L
<b>RIAG Ni 135 Carrier</b>	3 – 6 mL/L	4 mL/L
<b>RIAG Ni 122 Brightener</b>	0.1 – 0.5 mL/L	0.3 mL/L
<b>RIAG Ni 138 Tenside M</b>	1 – 3 mL/L	2 mL/L
pH-Value	3.8 – 4.5	4.2

### Operating Values

Nickel (Ni <sup>2+</sup> )	50 – 75 g/L	60 g/L
Chloride (Cl <sup>-</sup> )	15 – 21 g/L	18 g/L
Boric acid (H <sub>3</sub> BO <sub>3</sub> )	40 – 50 g/L	45 g/L

## Make up

A separate tank is filled with deionised water to 2/3 of the final volume.

The water is then heated to at least 60 °C after which the chemicals are added and the tank is filled to final volume with deionised water. To remove contaminants 0.5 mL/L Hydrogen peroxide is added and the solution is stirred for at least one hour. This is followed by addition of 3 – 5 g/L **RIAG Carb SF** activated carbon and mixing for another 30 minutes. After settling, preferably overnight, the electrolyte needs to be transferred to the working tank by filtration. Finally, the correct undiluted volumes of **RIAG Ni 132 Make up**, **RIAG Ni 135 Carrier**, **RIAG Ni 122 Brightener** and **RIAG Ni 138 Tenside M** are added whilst stirring.

## Operating Parameters

Temperature	55 °C (50 – 60 °C)
pH-value	4.2 (3.8 – 4.5)
Cathodic current density	0.1 – 2.0 A/dm <sup>2</sup>
Anodic current density	< 3.0 A/dm <sup>2</sup>
Current efficiency	< 100 %
Deposit rate	at 1 A/dm <sup>2</sup> ca. 0.2 µm/min
Anodes	Minimum purity 99.7 % Ni. We recommend polypropylene anode bags
Agitation	Essential: Barrel Rotation, Filter Pump
Tanks	Plastic or lined steel
Filtration	It is important to use continuous filtration and we recommend including activated carbon filtration as well. The filtration rate should be two to three times electrolyte volume per hour.
Heating	Immersion heaters, but thermostatic control is essential
Cooling	not required
Fume extraction	Recommended
Maintenance	Nickel sulphate, nickel chloride and boric acid should be analysed and corrected regularly. Additions of <b>RIAG Ni 122 Brightener</b> , <b>RIAG Ni 133 Carrier</b> and <b>RIAG Ni 135 Carrier</b> are made via ampere-hour consumption.  Metallic contamination can be removed by frequent selective plating-out at 0.1 – 0.3 A/dm <sup>2</sup> . The filter pump should be on with the filter outlet directed at the panels. This will ensure thorough electrolyte circulation and essential agitation at the same time.

pH-value setup	To lower the pH chem. pure Sulphuric acid (10 %) is added. To raise the pH only Nickel carbonate must be used. Ammonia or ammonia compounds must <u>not</u> be added. The brightener consumption will increase if the pH is above 4.5 while reduced levelling is noticed at a pH lower than 4.0.	
Additive consumption	The additives are consumed during electrolytic reactions as well a drag-out losses and the use per 10 kWh can therefore vary.	
	<b>RIAG Ni 122 Brightener</b>	1.0 – 2.0 L/10 kWh
	<b>RIAG Ni 132 Make up</b>	0.3 – 0.7 L/10 kWh
	<b>RIAG Ni 135 Carrier</b>	0.3 – 0.7 L/10 kWh
	<b>RIAG Ni 138 Tenside M</b>	0.2 – 0.5 L/10 kWh

## Function of electrolyte components

### RIAG Ni 122 Brightener

For uniform bright coatings it is essential to add the **RIAG Ni 122 Brightener** in small doses and according to the operating instructions. Smaller more frequent additions are important for optimal brightness and deposit quality. Regular pH-control and correction (4.2) reduce the **RIAG Ni 122 Brightener** consumption as well.

### RIAG Ni 135 Carrier

The **RIAG Ni 135 Carrier** is responsible for uniform bright coatings over a wide current density range (especially for low current density) and can be determined analytically.

### RIAG Ni 132 Make up / RIAG Ni 133 Carrier

The **RIAG Ni 132 Make up / RIAG Ni 133 Carrier** affects deposit ductility and the recommended concentration should not be reduced. It can also be determined analytically.

### RIAG Ni 138 Tenside M (for mechanically agitated solutions)

Reduces the surface tension and prevents pitting.

### RIAG Ni 146 Support

The additive **RIAG Ni 146 Support** is used to remove dark plating in low current density areas that cannot be removed by selective plating-out. However, the maximum **RIAG Ni 146 Support** addition must be limited to 1 mL/L. Higher dosing reduces levelling and deposit brightness. **RIAG Ni 146 Support** can also be used to compensate for brightener overdosing (0.2 – 0.5 mL/L). To avoid overdosing of **RIAG Ni 146 Support** additions should be made in small volumes.

## RIAG Ni 143 Purifier

Zinc-die casting processing in rack or barrel mode often leads to zinc and copper electrolyte contamination. This can be treated by additions of 0.1 – 0.5 mL/L **RIAG Ni 143 Purifier**. The volume **RIAG Ni 143 Purifier** to be added depends on contamination levels but overdosing results in loss of deposit brightness as well as levelling and must be avoided.

## Activated Carbon

Continuous filtration over activated carbon is recommended. This can be done via a by-pass whereby the carbon will remove organic contaminants such as oils and breakdown products. For this we recommend our **RIAG Carb SF** dust-free product which has an active surface area of 1500 m<sup>2</sup> /g. The additional brightener consumption should not exceed 5 %.

## RIAG Ni 147 Oxidant

Iron contamination (pitting) can be removed effectively by additions of **RIAG Ni 147 Oxidant**. The maximum concentration of 0.5 g/L should not be exceeded. The salt is first dissolved in hot water and the iron is removed via the filter.

## Environmental considerations and product safety

All concentrates, rinse waters and waste solution must be treated and discharged in accordance with local effluent control regulations. Information can be gleaned from the material safety data sheets. Chemicals shall not be stored below 10 °C.

## Liability

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## Analysis (Analytical Methods)

Sample preparation:

The sample must be taken from a well-mixed point and allowed to cool down to 25 °C.

### Boric acid

Reagents: Sodium hydroxide solution 0.1 mol/L  
Bromcresol purple (1 % in Ethanol)  
Mannitol

Procedure: 10 mL electrolyte are transferred via pipette into a 100 mL measuring flask and filled up to the mark with deionised water followed by mixing well.  
10 mL of this mixture is given into a  
250 mL beaker via pipette followed by  
100 mL deionised water addition.  
2 – 3 g Mannitol is added followed by addition of 10 drops Bromcresol purple.  
0.1 mol/L Titration with Sodium hydroxide from yellow to green, to dark green and finally to blue-violet.

Calculation: Boric acid (g/L) = consumption of mL NaOH x 6.18

### Nickel chloride

Reagents: Silver nitrate solution 0.1 mol/L  
Potassium dichromate solution 5 %

Procedure: 5 mL electrolyte are transferred into a  
250 mL glass beaker and diluted with  
50 mL deionised water.  
10 drops of Potassium dichromate solution are added.  
Titration with 0.1 mol/L Silver nitrate solution from white yellow to a light brown end point.

Calculation: Nickel chloride (g/L) = consumption of mL AgNO<sub>3</sub> x 2.380

Chloride (g/L) = consumption of mL AgNO<sub>3</sub> x 0.709

## Nickel

Reagents: Buffer solution pH 10  
Na<sub>2</sub>EDTA 0.1 mol/L  
Murexide (Sodium chloride 1: 100)

Procedure: 10 mL electrolyte are transferred via pipette into a  
100 mL measuring flask and filled- up to the mark with deionised  
water and mixed well  
10 mL of this mixture is given into a 250 mL glass beaker by  
pipette followed by  
15 mL Buffer solution addition  
100 mL deionised water and  
1 spatula tip of Murexide are added The sample colour should then be  
deep yellow  
Titrate immediately with Na<sub>2</sub>EDTA 0.1 mol/L to a blue-  
end- point

Calculation: Nickel (g/L) = consumption of mL Na<sub>2</sub>EDTA 0.1 mol/L x 5.869

Nickel sulphate (g/L) = [A – (B x 0.247)] x 4.48

A = Nickel concentration in g/L

B = Nickel chloride conc. in g/L