## Quality Control Laboratory

| Certificate of Analysis |  |  |  |
| :---: | :---: | :---: | :---: |
| Product: Magnesium Chloride Hexahydrate BP2017 |  |  |  |
| Batc Qua Pac Ref | $\begin{aligned} & \text { No.: MCH97-104 } \\ & \text { tity: } 1000 \mathrm{~kg} \\ & \text { ing: } 25 \mathrm{~kg} \\ & 3 \mathrm{P} 2017 \end{aligned}$ | Lab No.: QC718 Mfg. date: 20/12/97 Exp. date: 20/12/99 |  |
| No. | Chemical Analysis | Specifications | Results |
| 01 | Characteristics | Colourless crystals, hygroscopic | A white, crystalline powder |
| 02 | Identification A,B,C | According to BP requirements | Passes the test |
| 03 | Solubility | Very soluble in water, freely <br> soluble in ethanol ( 96 per cent) | Conforms |
| 04 | Appearance of solution | Solution S is clear and colourless | The Solution was clear \& colourless |
| 05 | Acidity or alkalinity | Max. 0.3 ml of 0.01 M NaOH or | The solution consumed 0.12 ml of |
|  |  | HCl | HCl 0.01M |
| 06 | Bromides | Max. 500 ppm | < 500 ppm |
| 07 | Arsenic | Max. 2 ppm | <2 ppm |
| 08 | Sulfates | Max. 100 ppm | < 100 ppm |
| 09 | Iron | Max. 10 ppm | < 10 ppm |
| 10 | Heavy metals | Max. 10 ppm | < 10 ppm |
| 11 | Calcium | Max. 0.1\% | < $0.1 \%$ |
| 12 | Water | 51-55\% | 52.8\% |
| 13 | Assay (as $6 \mathrm{H}_{2} \mathrm{O}$ ) | 98.0-101.0 \% | 99.9\% |
| Date of Sampling:21/12/97 |  | - Approved | $\square \quad$ Rejected |
| Date of Analysis:22/12/97 |  |  | Rel |
|  | Analyst: <br> tied | Q.C.Manager: Sorkhmanel | Authorized by: hasolporan |

We hereby certify that this product has been prepared under GMP regulation and tested according \& conform to the requirements of BP2017. The raw materials, manufacturing process and product do not contain any of the solvents listed in residual solvents $B P<5.4>$


TABRIZ PHARMA Tabriz Chemical and Pharmaceutical Ind.

