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Dear Customer:

Please be advised that the following products theoretically comply with the requirements for Residual Solvents in excipients per the current editions of United States Pharmacopeia (USP) General Chapter <467> Residual Solvents, General Chapter 5.4 of the European Pharmacopoeia, and the International Conference of Harmonization (ICH) Q3c Residual Solvent (Organic Volatile Impurities) Guideline. These products are not manufactured with Class 1, Class 2 or Class 3 solvents listed in the aforementioned sources. Based on knowledge of the manufacturing processes and raw materials, other organic solvents are not likely to be present. Therefore, Innophos does not routinely test these products for the presence of residual solvents.

A-TAB® Dicalcium Phosphate Anhydrous (USP, Ph. Eur., & JP Grades)

CALIPHARM® A Dicalcium Phosphate Anhydrous (USP, Ph. Eur., & JP Grades)

CALIPHARM® D Dicalcium Phosphate Dihydrate (USP, Ph. Eur., & JP Grades)

CALIPHARM® T Tricalcium Phosphate (NF & Ph. Eur. Grades)

DI-TAB® Dicalcium Phosphate Dihydrate (USP, Ph. Eur., & JP Grades)

NUTRA-TAB® Co-processed Tricalcium Phosphate and Guar Gum Excipient

TRI-TAB® Tricalcium Phosphate (NF & Ph. Eur. Grades)

TRICAL-WG® Tricalcium Phosphate (NF & Ph. Eur. Grades)

The Quality and Regulatory Department

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