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Dear Customer:

Please be advised that the following products 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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The information contained in this document is offered as a service to our customers and is subject to change (Innophos undertakes no obligation to notify customers of any such change unless Innophos has previously agreed in writing to do so). Customers are solely responsible for determining the suitability of Innophos' products for any contemplated markets, uses, and/or applications, and for ensuring that all such uses and applications (including customer's labeling of its products) comply with applicable law.