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USP 232 Oral Impurities – Limits for the amounts of elemental impurities in drug products are specified by USP. USP Method 561 is now available for the Determination of the US Food and Drug Administration's (FDA) regulation of residual pesticide contaminants for the pharmacological industry.

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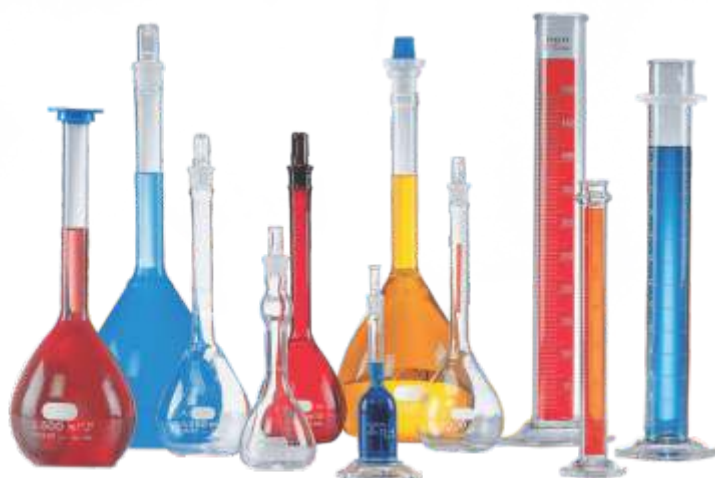


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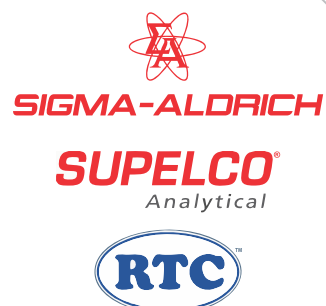


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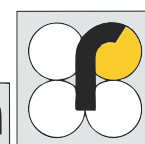
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