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Chapter 1 Pharmaceutical Salts and Co-crystals: Retrospect and Prospects 1 R. Desiraju Acknowledgement 6 References 6 Chapter 2 Fundamental Aspects of Salts and Co-crystals 9 Andrew D. Bond 2.1 Introduction 9 2.2 Definitions: Salts and Co-crystals 10 2.3 Supramolecular Chemistry, Intermolecular Interactions and Crystal Packing 12

### **Pharmaceutical Salts and Co-crystals - GBV**

â€¢ Salts and salt co-crystals are additional solid forms pharmaceutical scientists can use to change properties of an API. â€¢ Salts and salt co-crystals can exhibit polymorphism and polymorph screening should be performed to find stable, metastable, hydrated, and solvated salt forms. â€¢ Issues, such as disproportionation and hygroscopicity ...

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pseudopolymorphs (or solvates), and co-crystals (Figure 3). Liquid drug formulations from salification are rarely found and are usually based on eutectic mixtures [24], however, salt drug generation can result in a liquid salt, known as ionic liquid.

### **Pharmaceutical Salts: Solids to Liquids by Using Ionic**

Pharmaceutical Co- Crystals, which classified co-crystals as a drug product intermediate (or as an in-process material), and replaces the 2013 guidance of the same name.

### **Regulatory Classification of Pharmaceutical Co-Crystals**

Thus, the salt is often the preferred form for isolating and purifying the drug. Historically, the number of available salts was rather limited; however, today there is a wide range of chemical entities that are recognized as being safe, which can be used in the preparation of drug products (cf Table 1) , .

## **Pharmaceutical salts: Theory, use in solid dosage forms**

An Overview of Solid Form Screening During Drug Development Ann Newman Seventh Street Development Group PO Box 526, Lafayette, IN 47902 ... This document was presented at PPXRD - Pharmaceutical Powder X-ray Diffraction Symposium Sponsored by The International Centre for Diffraction Data. ...  
â€¢Salts, cocrystals, polymorphs, dispersions, amorphous

## **An Overview of Solid Form Screening During Drug Development**

Salts can be broadly classified into two groups, covalent and ionic. Figure 1. The formation of the covalent pharmaceutical salt fluticasone propionate. Covalent-bonded salts are formed when the parent drug and the neutralisation agent are held together by the sharing of electrons.

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