

1 *Invited Speaker Abstract*

2 **Investigating to optimal ratio between drug and co-**
3 **former in co-amorphous systems**

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7 **Abstract:** Most new low molecular weight chemical entities in pharmaceutical developments suffer
8 from a low aqueous solubility, making oral delivery challenging. Despite the numerous formulation
9 efforts that can be investigated, research especially on amorphous drugs and formulations appears
10 to be a useful approach. Whilst few drugs can be converted to an amorphous form on their own,
11 due to a too physical stability, the use of amorphous solid dispersions, i.e. the dissolution of drug
12 molecules into (amorphous) polymers is increasingly used. However, certain shortcoming of these
13 polymers based amorphous solid dispersions, such as a low drug load and a usually high
14 hygroscopicity; still necessitate the investigation of alternative approaches. One such approach is
15 the use of co-amorphous systems, i.e. to combination of initially crystalline low molecular weight
16 drugs and excipients. Usually here a 1:1 molar ratio is used, but this may not be the optimal mixing
17 ratio. In this presentation, work on investigating to optimal ratio between drug and co-former will
18 be presented and critically discussed.

19 **Keywords:** Poor solubility, amorphous, co-amorphous



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