

## **September 2010:**

### **Simplifying Drug Dissolution Testing:**

#### **Changes at the USP**

**Can a dissolution test be used for checking/establishing lot-to-lot consistency?  
Maybe not!**

**Using non-compendial vs compendial methods (apparatuses/procedures)**

**Drug Dissolution Testing: Objective vs Practices**

## **August 2010:**

**Determining blood concentration-time (C-t) profiles from in vitro dissolution results  
and product evaluation – carbamazepine.**

**Changing a dissolution method in the middle of product development stage and/or  
after?**

**“Developing IVIVC” and establishing drug concentration-time (C-t) profiles**

### **Info**

**PVT (Performance Verification Test) – Difficulties and a suggestion to address those**

## **July 2010:**

**BCS and its role in product evaluation – Are underlying assumptions appropriate?**

**One Step (Product Evaluation) Approach**

**Guidance documents and their limitations**

**Trends in FDA dissolution methods database**