

[View on rigaku.com](https://www.rigaku.com)

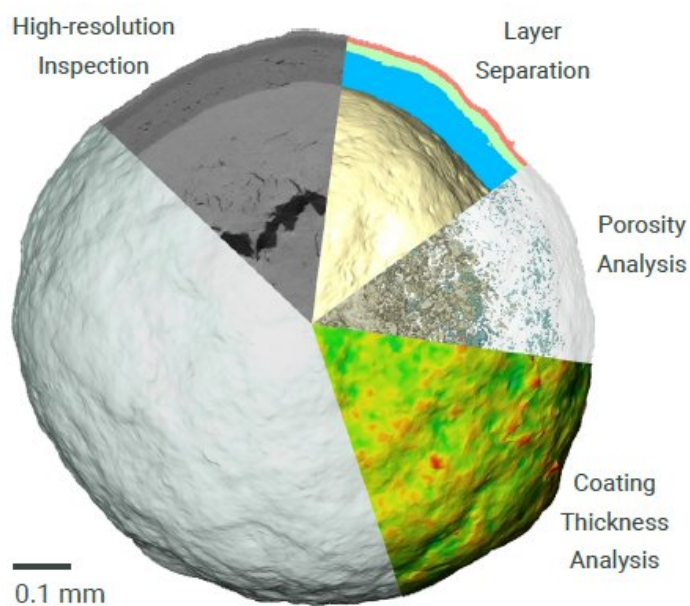
# XCT2102 - Submicron CT for the pharmaceutical industry

## Introduction

X-ray submicron computed tomography (submicron CT) is one of the most powerful methods for 3D visualization and inspection of any type of sample or product. This non-destructive method provides sufficient resolution and contrast to evaluate any microstructural features, with the ability to resolve structures even below one micron. Moreover, this method requires minimal/no sample preparation, eliminating complicated tasks such as embedding, coating or thin slicing required with other high-resolution methods. The Rigaku nano3DX represents state-of-the-art laboratory-based nanoscale X-ray imaging. This device, when used with deep learning methods, is an unmatched tool for pharmaceutical applications from R&D to production and inspection.

## Instrument

The nano3DX is a true X-ray microscope (XRM) with the ability to measure relatively large samples at very high resolution. This is accomplished by using a high-powered rotating anode X-ray source and a high-resolution sCMOS X-ray camera. The rotating anode provides for fast data acquisition and the ability to switch anode materials easily to optimize the data acquisition.

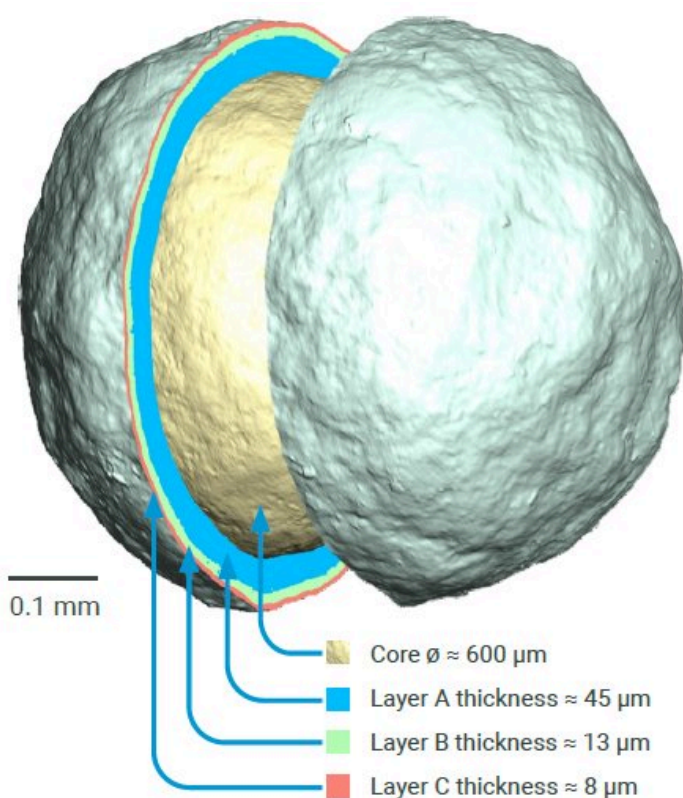


**Figure 1:** nano3DX particle analysis possibilities.

## CT data and results

The microstructure of solid dosage forms of pharmaceutical products is a critical factor that impacts disintegration and dissolution rates. As such, microstructure will also play a key role in bioequivalence and therapeutic equivalence. Being able to image the microstructure of a solid dosage form allows optimization of production and formulation procedures to achieve a robust dissolution response. If an out-of-specification dissolution result is later observed, analysis of the solid dosage form's internal structure and microstructure can yield a wealth of insights not accessible through traditional analytical approaches, and help resolve mission critical investigations.

New product development can be a highly time-consuming and expensive task. Using Rigaku's nano3DX, this process can be accelerated, providing immediate feedback on a product's internal structure when any discrepancies between expected and actual attributes can be identified.



**Figure 2:** Coating's morphological analysis—in the internal structure of the analyzed particle, four specific layers were resolvable: core layer, A= immediate drug layer, B= modified drug layer, C= cellulose acetate layer (ordered according to their thickness).

### nano3DX provides insight into key factors such as:

#### Product morphological analysis and structural analysis

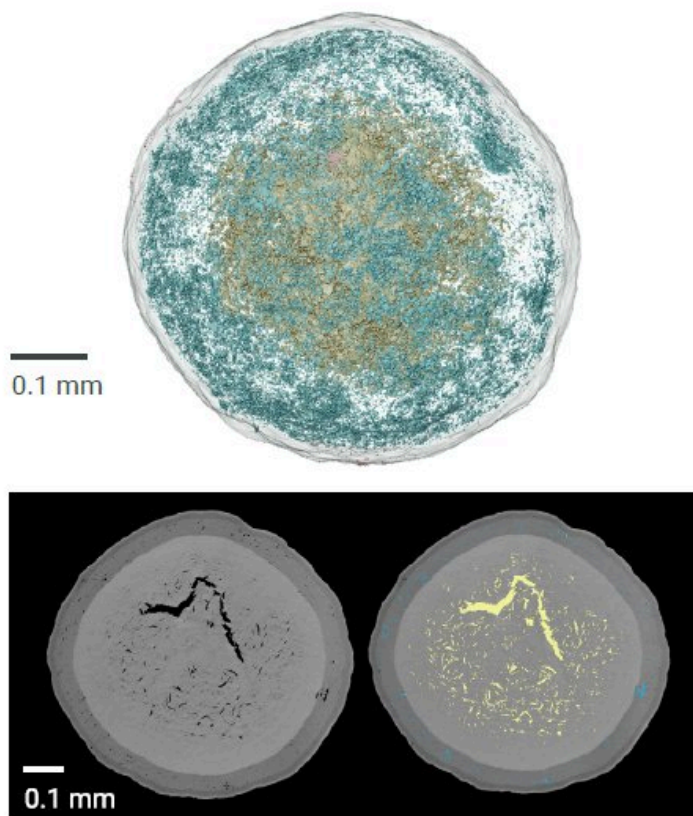
- Coatings – thickness analysis
- Porosity – pores/porous network analysis

- Crystallinity – crystalline phases analysis
- Aggregates detection
- Distribution of API analysis
- Dissolution process evaluation

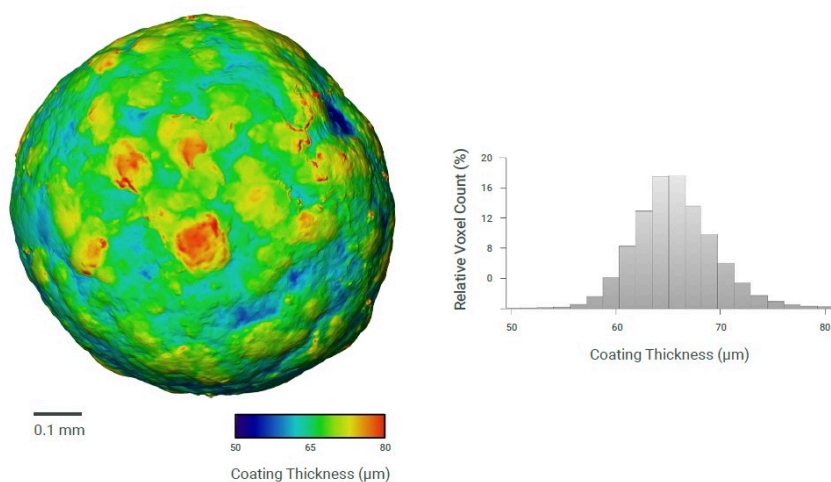
### Optimization of manufacturing process

#### – Defects Detection

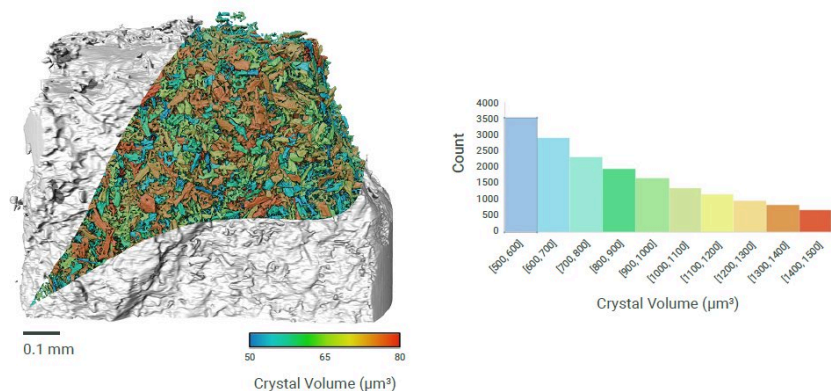
- Cracks
- Coatings thickness homogeneity
- Contamination



**Figure 3:** Porosity analysis—in the internal structure of the analyzed particle, a high number of pores were detected, which is reflected by the total porosity of 7.43%. Color coding was used according to distance from particle center: Pores in the core layer are yellow and pores in the outer layers are blue.



**Figure 4:** Coating thickness homogeneity analysis—the thickness of the outer coating layer (cellulose acetate layer) of the analyzed particle varied from 50 to 80  $\mu\text{m}$ , reflecting low manufacturing quality.



**Figure 5:** Crystallinity analysis—in the internal structure of the analyzed powder, the crystalline phases were resolvable due to high contrast and resolution. The volume of those phases varied from 500 to 1500  $\mu\text{m}^3$ .



CT analysis performed by [CEITEC](#) – CT Lab

---

## Related products



### **nano3DX**

Ultrahigh resolution nanotomography using parallel beam geometry