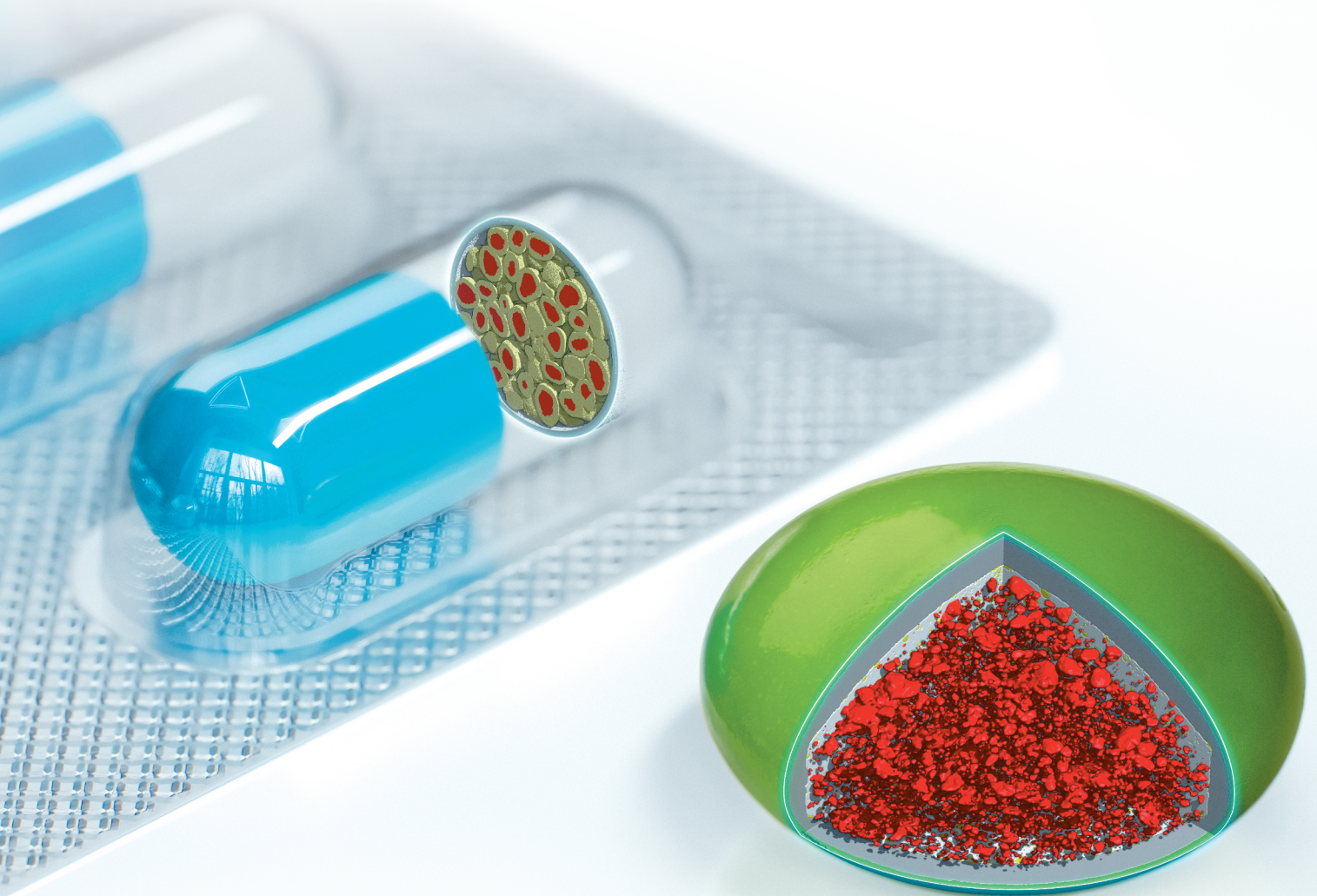


# GEO DICT

The Digital Material Laboratory

DIGITAL SOLUTIONS  
FOR  
PHARMACEUTICAL  
FORMULATION



## THE MOTIVATION

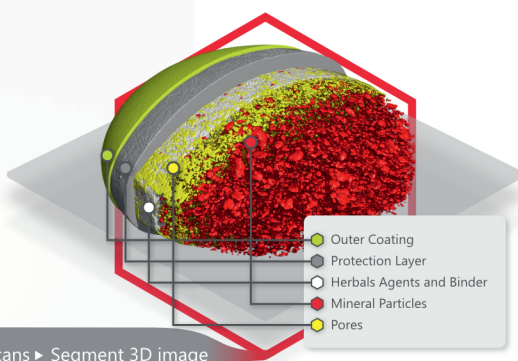
- Trial-and error in formulation is **costly, slow, and hard to scale.**
- Lack of microstructure-property insights limits robust design.
- Digital tools and workflows are needed for **faster, transparent, and reproducible development.**

## OUR SOLUTION

- **GeoDict software:** platform for digital material development.
- Combines **image-based modeling, digital twins, and multiphysics simulations.**
- Supports scientist and engineers in **all stages of solid dosage and delivery design.**

## YOUR BENEFIT

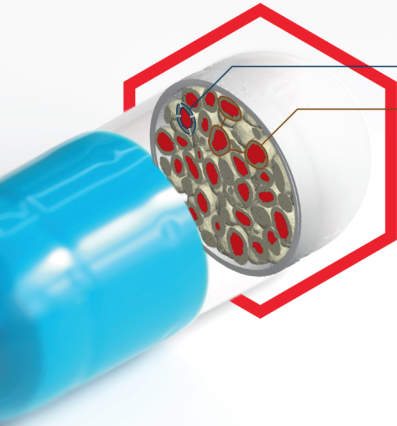
- Accelerate **formulation development** with predictive simulations.
- Improve robustness and ensure **consistent product quality.**
- Enable **QbD, scale-up, and regulatory submissions** by digitally identifying safe operating ranges for formulations and manufacturing.



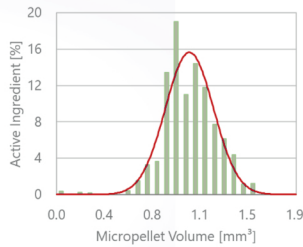
Collect and process 3D scans ▶ Segment 3D image

## DIGITALIZATION

- Segment and convert  $\mu$ CT scans into 3D digital structures of powders, tablets, pellets, capsules... with image filters and AI tools
- These digital structures are the basis for analysis and multiphysics simulations.

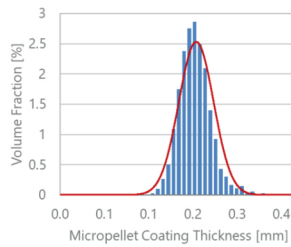


Micropellet Size Distribution



■ Histogram ■ Gauss fit

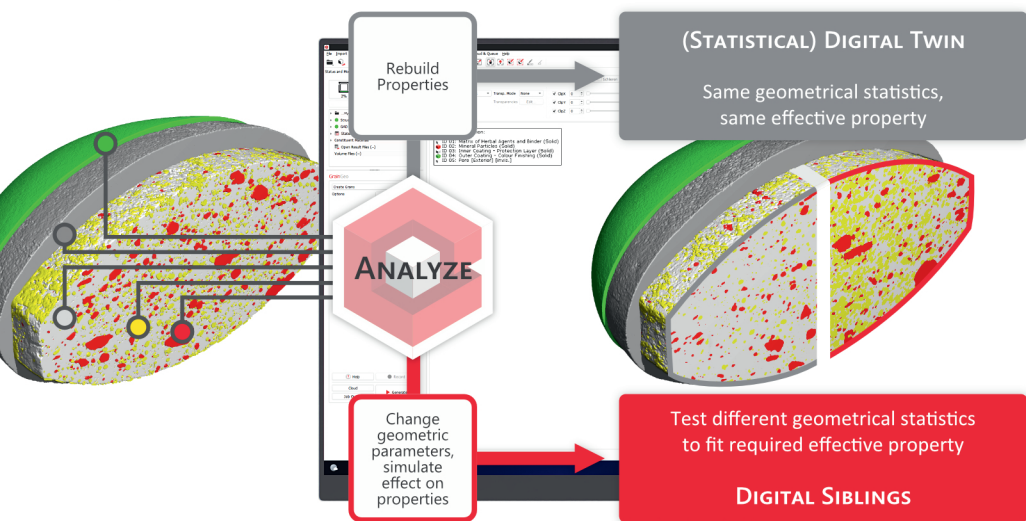
Coating Thickness



■ Histogram ■ Gauss fit

## NON-DESTRUCTIVE ANALYSIS & QUALITY CONTROL

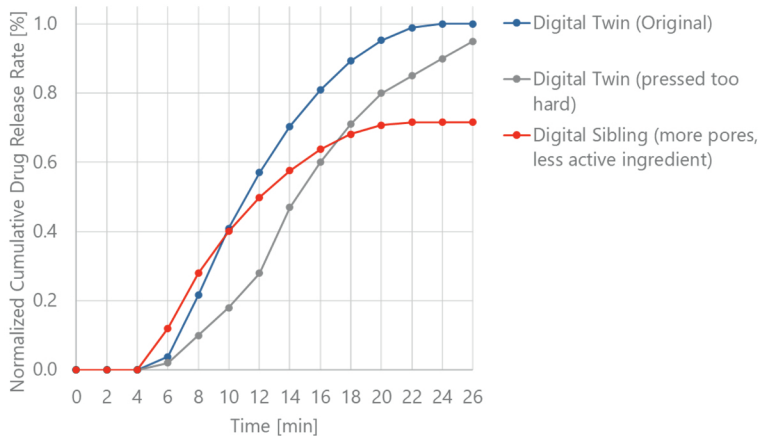
- Extract grain & pore size distribution, packing density, porosity, tortuosity, and coating thickness directly from 3D scans.
- Perform digital testing of stability and dissolution without destroying samples.
- Enable simulation-based analytics for reliable quality control.



## DIGITAL TWINS & DIGITAL SIBLINGS

- Build digital twins of powders, granules, tables, capsules, and more in full 3D detail from parameters extracted from digital analysis.
- Generate digital siblings by systematically varying particle morphology, excipient ratios, compaction force, or coatings uniformity.
- Conduct sensitivity analysis and virtual scale-up prior to manufacturing.

Results from Dissolution Simulation - Cumulative Drug Release Rate



## PROPERTY PREDICTION

- Run multiphysics simulations to predict packing, compaction, mechanical robustness, friability, permeability, release, and dissolution kinetics.
- Directly link microstructure to critical quality attributes.
- Support QbD strategies, regulatory submissions, and high reproducible, accelerated formulation development with physic-based predictions.