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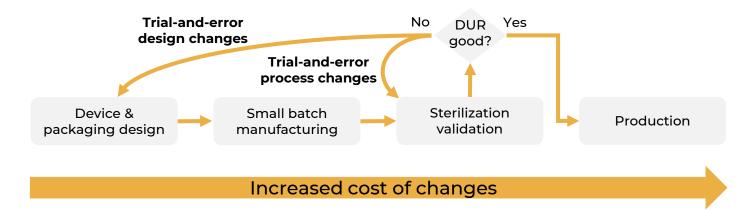
# Accelerated time to market with Design for Sterilization

#### The status quo:

## Sterilization validation can only start with manufactured, packaged devices

In a typical design process, sterilization validation is only considered after the product has been fully designed, built, tested, and readied for production.

With this approach, sterilization validation typically takes <u>many months and</u> <u>can cost hundreds of thousands of dollars.</u>

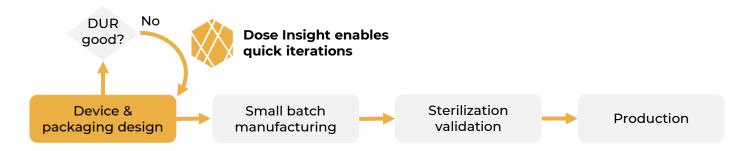


## With **Dose Insight**:

## Sterilization validation starts as soon as you have a CAD model!

Virtual dose mapping allows for design choices to be evaluated with respect to sterilization, and to systematically optimize the sterilization process.

Dose Insight's service addresses all uncertainties early in the design process, resulting in <u>significant cost savings and accelerated time to market.</u>



Increased cost of changes