



Orally Inhaled Drug Product PSGs: General Considerations Using the Alternative Bioequivalence (BE) Approach In Lieu of Comparative Clinical Endpoint (CCEP) BE Study for Suspension-Based Metered Dose Inhalers

Advancing Generic Drug Development 2024: Translating Science to Approval

Day (1), Session (3): (Research to Support Guidance Development for Inhalation Drug Products)

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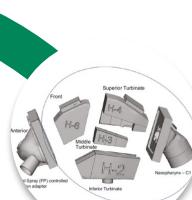
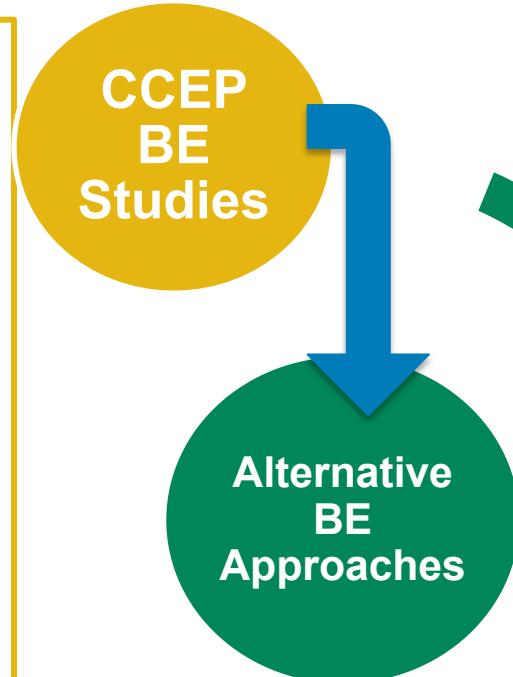
Learning Objectives

- Recognize the challenges with conducting comparative clinical endpoint (CCEP) bioequivalence (BE) studies for orally inhaled drug products (OIDPs).
- Describe the available tools, supportive FDA research, and external input for developing alternative BE approaches.
- Identify recently developed product-specific guidances (PSGs) for suspension-based metered dose inhalers (MDIs) with alternative BE approaches to the CCEP BE study and study design considerations.

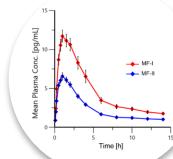
The Challenges with CCEP BE Studies



- FDA traditionally recommends CCEP BE studies as part of a BE assessment for locally-acting MDIs and DPIs.
- CCEP BE studies can pose several **challenges** for generic applicants developing an MDI or DPI.
 - *Higher variability* → lower accuracy and reproducibility
 - *Flat exposure-response* → lower sensitivity
- Ultimately, these challenges necessitate using **large numbers** of patients often over a **long study duration**.
 - *Costly*
 - *Time Consuming*



In Vitro Methods

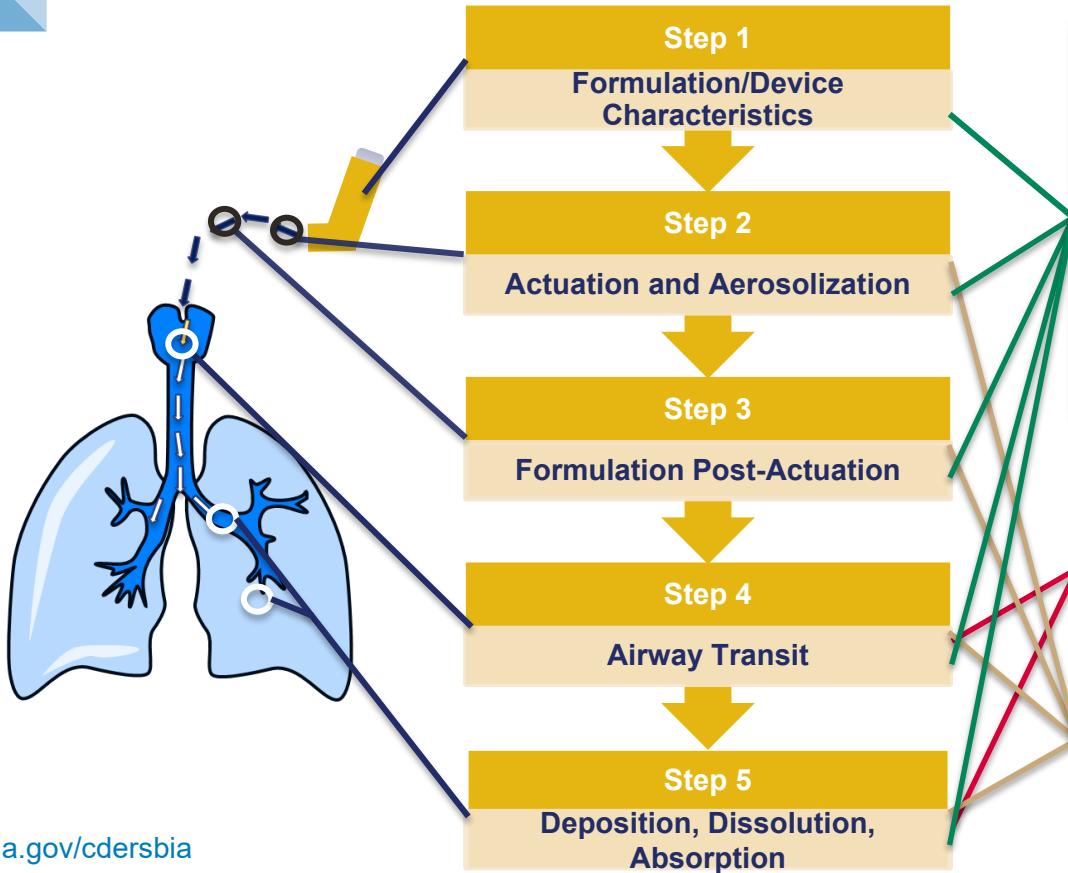


In Vivo Methods



In Silico Methods

Potential Methods for Assessing Contributing Factors to Local Drug Delivery



IN VITRO STUDY METHODS

- Realistic Aerodynamic Particle Size Distribution (APSD)
- Dissolution
- Optical Suspension Characterization
- Droplet Size Distribution by Laser Diffraction
- Morphology-assisted Raman Spectroscopy (MRS)
- Scanning Electron Microscopy (SEM)
- X-ray Tomography
- Shadowgraphic imaging/shadow motion analysis
- Phase Doppler Interferometry/Anemometry
- Particle Image Velocimetry
- Optical Photothermal Infrared Microscopy
- Atomic Force Microscopy – Infrared Microscopy
- Cell Permeability Assays

IN VIVO STUDY METHODS

- Charcoal Block PK Study
- Imaging-based Study (e.g., Scintigraphy)

IN SILICO STUDY METHODS

- Computational Fluid Dynamics (CFD)
- Regional Deposition Modeling
- Physiologically Based PK modeling (PBPK)
- Population PK Modeling

Alternative BE Approach: Solution MDIs

FDA

Product-specific guidances (PSGs) on *Beclomethasone Dipropionate Metered Inhalation Aerosol* (NDA 020911; NDA 207921), *Ipratropium Bromide Metered Inhalation Aerosol* (NDA 021527), and *Ciclesonide Metered Inhalation Aerosol* (NDA 021658)

If a generic demonstrates formulation sameness (qualitative and quantitative) and device similarity to the reference MDI, FDA recommends additional supportive studies to help ensure **equivalence at the local site of action** (i.e., lungs):

Actuation,
Aerosol
formation

Formulation
Post-
actuation

Transit
through the
airways;
Deposition,
Dissolution,
Absorption

Methods for
further
support

Characterization of Emitted Sprays (velocity profiles and evaporation rates)

- Understand emitted droplet size and evaporation process of formulation (volatiles + non-volatiles)

Morphology Imaging Comparisons (characterization of full range of residual drug particle sizes)

- Understand residual particle morphology and size distribution of emitted formulation

More Predictive APSD Testing (representative mouth-throat models and breathing profiles)

- Understand impact of patient variability

Dissolution

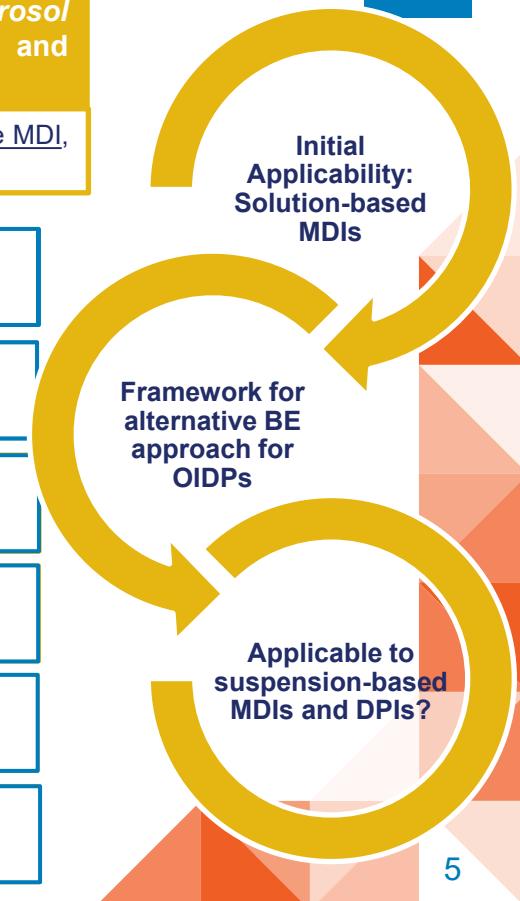
- Understanding how drug(s) dissolves at the site of action for absorption once deposited

Quantitative Methods and Modeling (e.g., PBPK, CFD studies)

- IVIVCs to bridge gap between in vitro product performance and regional drug deposition

Alternative PK BE Studies

- Understanding how PK studies may correlate to local deposition



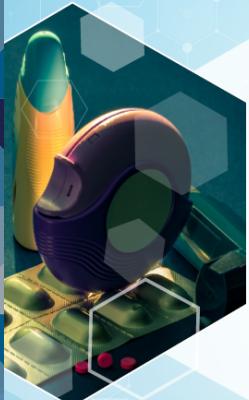
External Input Informs FDA Thinking on Alternative BE Approaches for OIDPs



Considerations for and Alternatives to Comparative Clinical Endpoint and Pharmacodynamic Bioequivalence Studies for Generic Orally Inhaled Drug Products

April 20-21, 2023
8:30 AM – 5:30 PM

In-Person and Virtual Options to Attend



The purpose of this two-day orally inhaled drug products (OIDP) workshop is to discuss the current scientific and regulatory perspectives for using *in vivo*, *in vitro*, and *in silico* studies as alternatives to comparative clinical endpoint (CCEP) and pharmacodynamic (PD) bioequivalence (BE) studies, and to explore potential designs for alternative BE approaches that can address the particular challenges associated with establishing local drug delivery equivalence for suspension-based metered dose inhalers (MDIs) and dry powder inhalers (DPIs).

Workshop Topics:

- Reviewing successes with the use of CCEP and PD BE studies to establish BE for locally acting OIDPs, and discussing relevant challenges
- Evaluating alternative BE approaches that utilize *in vitro*, *in vivo*, and *in silico* studies, instead of CCEP and PD BE studies, and discussing relevant technical and practical issues when used with different OIDPs
- Discussing the integration of multiple alternative *in vitro*, *in vivo*, and *in silico* studies to form cohesive alternative BE approaches in lieu of CCEP or PD BE studies for MDIs and DPIs

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<https://www.complexgenerics.org/education-training/considerations-for-and-alternatives-to-comparative-clinical-endpoint-and-pharmacodynamic-bioequivalence-studies-for-generic-orally-inhaled-drug-products-2/>



- Two-day workshop to discuss the Agency's *scientific understanding and regulatory perspective on alternative BE approaches* with industry representatives and academic experts.
- In-person attendees participated in small group discussions that provided FDA with valuable insight into the *industry's experiences* with alternative BE approaches and their thinking on potential approaches for complex OIDPs (suspension MDIs and DPIs).



External Input Informs FDA Thinking on Alternative BE Approaches for OIDPs



- Most **alternative approaches** are generally *applicable to both MDIs and DIs* irrespective of their formulation.
- Certain approaches are *more critical and informative*.
- Inclusion of a particular study may be *product-specific* (e.g., dependent on the drug substance properties).
- Some approaches useful for *product development* vs. others for assessing **BE**.

Useful Study Methods

- Realistic APSD
- Dissolution
- In silico methods

Potentially Useful or Confirmatory

- Particle morphology
- Charcoal-block PK study

Study Methods with Limited Utility

- Evaporation rate and velocity profile evaluation
- Pre-actuation characterization of the formulation

Implementing the Agency's Current Thinking for Suspension MDIs



- Recent suspension-based MDI PSGs: **option-based approach** for establishing BE
 - Specific study designs (e.g., supportive characterization studies or optional components) remain **product-specific**

Option 1

Formulation Sameness

- No difference in formulation (e.g., Q1/Q2 sameness to RS)*

Product Performance Equivalence

- In Vitro BE studies

Systemic Exposure Equivalence

- In Vivo PK BE Study

Local Drug Delivery Equivalence

- Alternative BE approach (In Vitro Studies, Characterization Studies, Charcoal PK BE Study, In Silico Studies)*

Device Similarity Equivalence

- Device Similarity to the RLD

Option 2

Formulation Sameness

- None**

Product Performance Equivalence

- In Vitro BE studies

Systemic Exposure Equivalence

- In Vivo PK BE Study

Local Drug Delivery Equivalence

- In Vivo CCEP BE Study**

Device Similarity Equivalence

- Device Similarity to the RLD

Implementing the Agency's Current Thinking for Suspension MDIs



BEVESPI AEROSPHERE BREZTRI AEROSPHERE

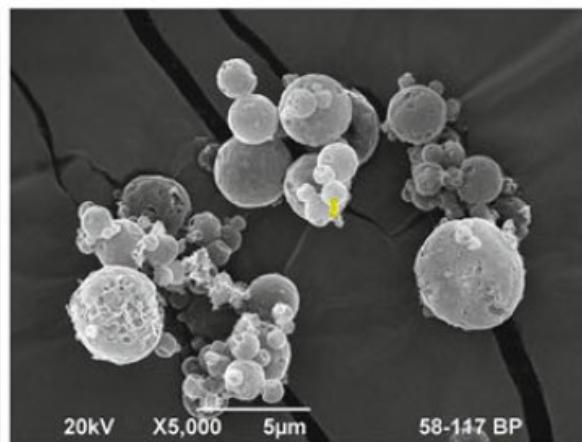
Formoterol Fumarate;
Glycopyrrolate Metered
Inhalation Aerosol



Budesonide; Formoterol Fumarate;
Glycopyrrolate Metered Inhalation
Aerosol



- ***FDA-approved suspension-based MDIs***
- **Indication:** the maintenance treatment of patients with chronic pulmonary obstructive disease (COPD).



An example of phospholipid-based porous particles utilized in several MDI products.

- **Formulation:** co-suspension formulation of drug particles and phospholipid-based porous particles in propellant.
 - **Porous particles:** 1,2-distearoyl-sn-glycero-3-phosphocholine (DSPC) and calcium chloride

Suspension MDI PSGs

Incorporating Alternative BE Approaches

Draft Suspension MDI PSGs (Feb 2024)

Formoterol Fumarate; Glycopyrrolate Inhalation Aerosol, Metered

Budesonide; Formoterol Fumarate; Glycopyrrolate Inhalation Aerosol, Metered

Draft Suspension MDI PSGs (Aug 2024)

Fluticasone Propionate Inhalation Aerosol, Metered

Fluticasone Propionate; Salmeterol Xinafoate Inhalation Aerosol, Metered

Albuterol Sulfate Inhalation Aerosol, Metered

Levalbuterol Tartrate Inhalation Aerosol, Metered

Option 1 BE Approach

- Formulation**
 - The test (T) product should contain ***no difference in inactive ingredients or other aspects of the formulation*** relative to the RS that may affect local or systemic availability (e.g., Q1/Q2 formulation sameness)
- In Vitro BE Studies**
 - SAC, APSD, spray pattern, plume geometry, priming/repriming
 - Realistic APSD (rAPSD)***
 - Dissolution****
- Comparative Characterization Studies**
 - Particle Morphology of the Emitted Dose***
- In Vivo Studies**
 - In Vivo PK BE Study
 - In Vivo PK BE study with Charcoal Block***
- Additional Information**
 - Optional Computational Modeling study
 - Device similarity to the RLD

Suspension MDI PSGs

Incorporating Alternative BE Approaches

Draft Suspension MDI PSGs (Feb 2024)

Formoterol Fumarate; Glycopyrrolate Inhalation Aerosol, Metered

Budesonide; Formoterol Fumarate; Glycopyrrolate Inhalation Aerosol, Metered

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Fluticasone Propionate Inhalation Aerosol, Metered

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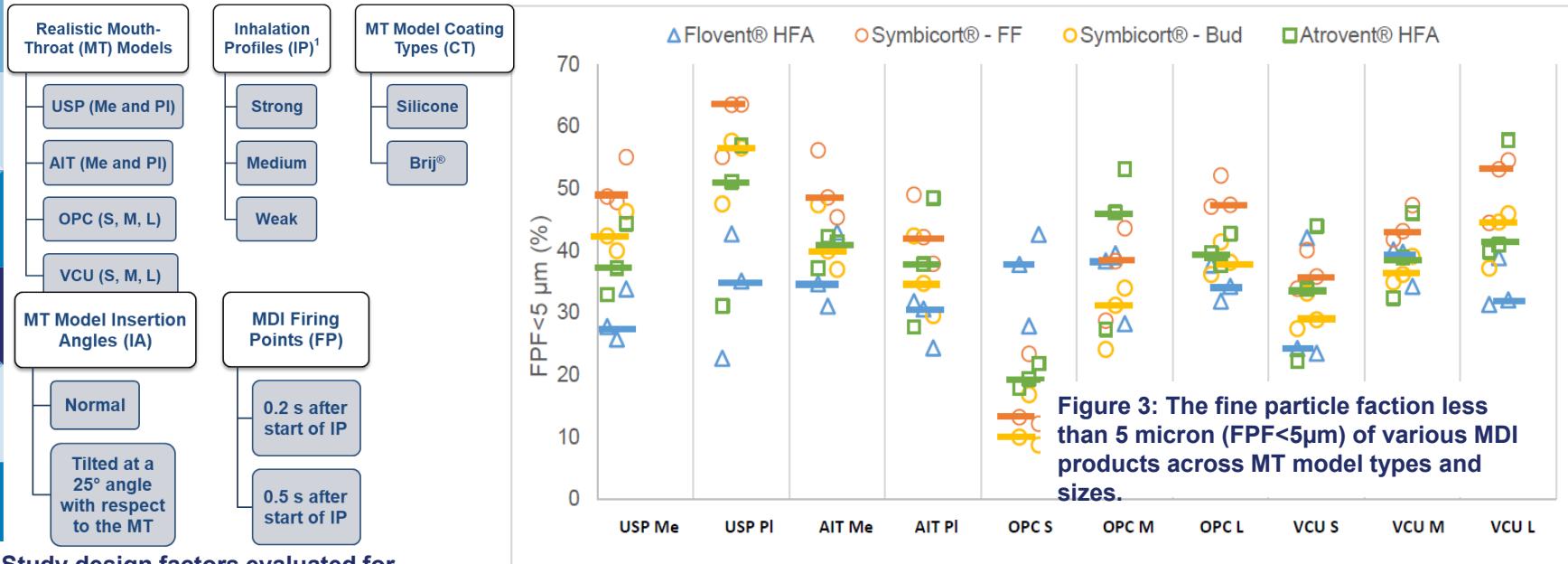
Option 2 BE Approach

- **Formulation**
 - *No recommendations provided (e.g., T product formulation can be Q1/Q2 or non-Q1/Q2 to RS formulation)*
- **In Vitro BE Studies**
 - SAC, APSD, spray pattern, plume geometry, priming/repriming
- **Comparative Characterization Studies**
 - Particle Morphology of the Emitted Dose
- **In Vivo Studies**
 - In Vivo PK BE Study
 - *CCEP BE study in subjects with asthma*
- **Additional Information**
 - Optional Computational Modeling study
 - Device similarity to the RLD

Realistic APSD Study Design Considerations



- GDUFA-Funded Research Outcomes
 - Response to the various study factors is **product-specific**.
 - **Method Development:** consider mouth-throat (MT) types and size, inhalation profiles (IPs), and other factors.



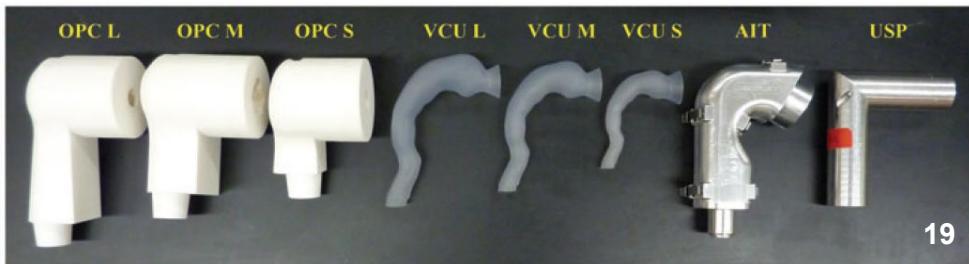
Study design factors evaluated for rAPSD with solution and suspension-based MDIs.

USP: United States Pharmacopeia; AIT: Albert Idealized Throat; OPC: Oropharyngeal Pharmacopeia Consortium; VCU: Virginia Commonwealth University

Realistic APSD Study Design Considerations

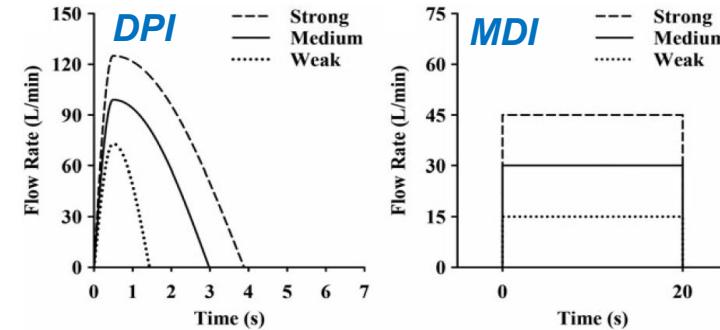
FDA

Realistic mouth-throat (MT) models

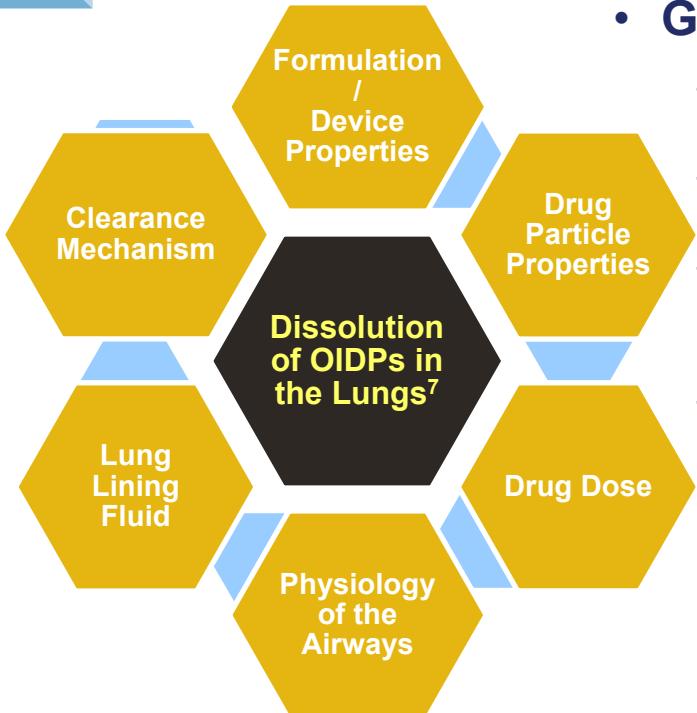


- **PSG Recommendations:**
 - **Beginning** lifestage.
 - Include different **MT sizes** and **IPs** that reasonably cover the expected inter-subject variability of the indicated patient population via **bracketing approach**.
 - **Example:** Small and large MT sizes + weak and strong IPs the cover patient population.
 - Correlate in vitro performance to in vivo lung deposition data, if available.
 - IPs obtained from patients.
 - **BE: population bioequivalence (PBE) of impactor sized mass (ISM)** for each MT model-IP combination.
 - **Alternative statistical approaches** may be used if scientifically justified.
 - Request a **Pre-ANDA meeting** to discuss **alternative approaches** to the study design and/or statistical methods.

Inhalation profiles (IPs)



Dissolution Study Design Considerations for OIDPs

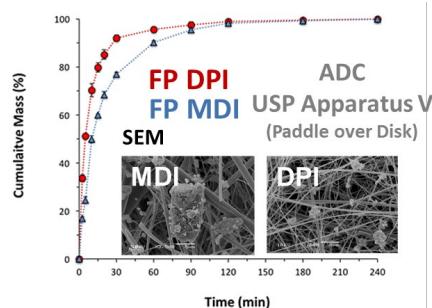


Drug dissolution in the lungs can be impacted by multiple factors.

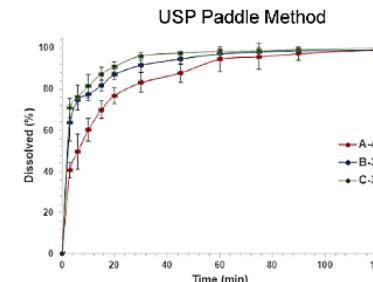
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- GDUFA-funded research

- Many contributing factors that can affect *dissolution performance* and *study sensitivity*.
- Currently no standardized method; method development is *product-specific*.
- Can develop dissolution methods that are sensitive and discriminatory to meaningful differences in *formulation* and/or *manufacturing process*.
- The need for dissolution studies is *drug-* (e.g., high/low solubility) and *product-specific*.



Dissolution of OIDPs are sensitive to differences in both dosage form (left) and particle size (right).



Dissolution Study Design Considerations for OIDPs



- PSG Recommendations:

Sample Collection

- *Beginning* Lifestage.
- Collect aerosolized dose of *similar drug mass* between T and RS products.
- Optimized and validated method (e.g., apparatus, sample collection, dose, media type and volume, stirring/agitation rate, sampling times).
- Discriminatory (e.g., differences in *deposited drug particle size*).
- BE: Comparative analysis of dissolution profiles with an appropriate statistical method (e.g., *similarity [f2] factor*).

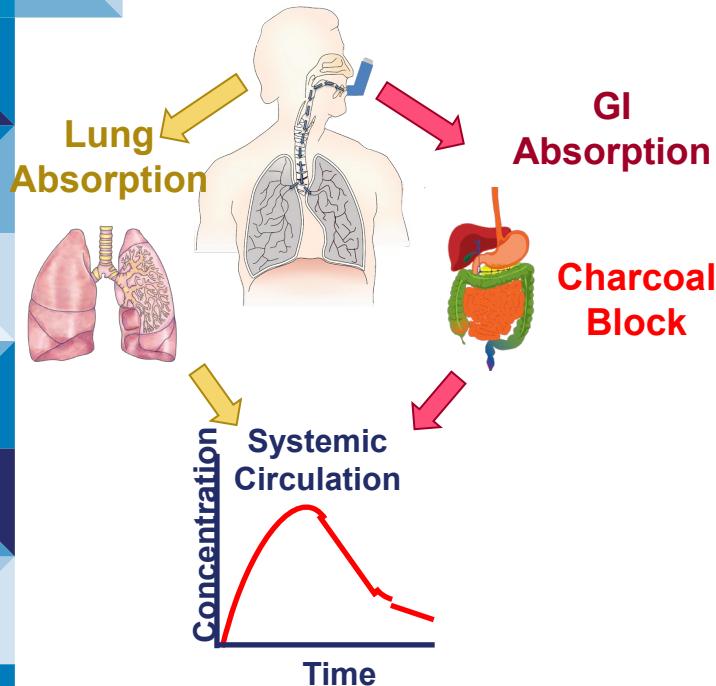
Dissolution Apparatus

Dissolution Media

Method Validation

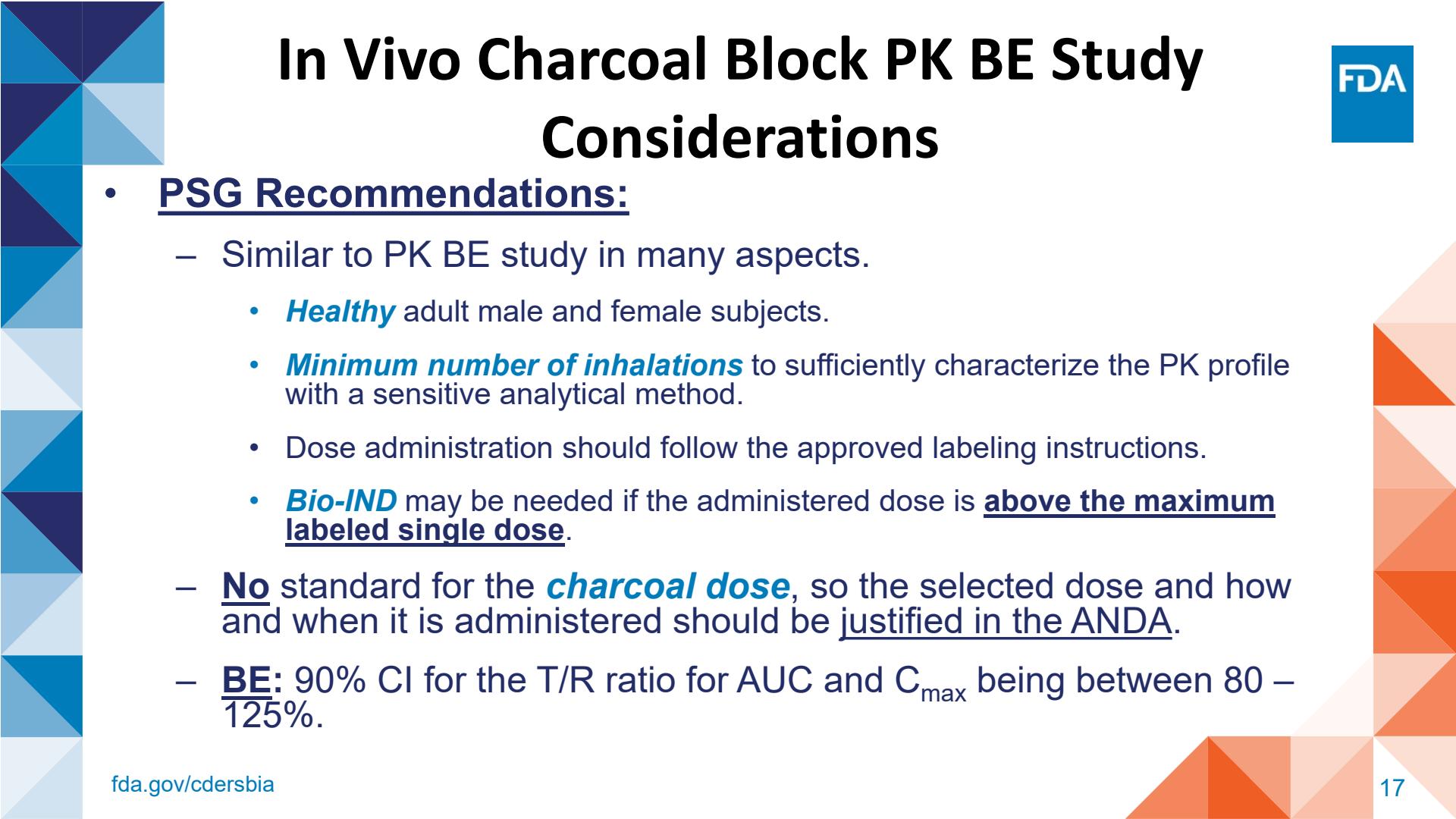
Assessment

In Vivo Charcoal Block PK BE Study Considerations



Drug absorption into the systemic circulation following dosing with certain OIDPs can occur through both lung absorption as well as gastrointestinal (GI) absorption. Dosing with charcoal can block GI absorption.

- For OIDPs, a portion of the emitted dose may be swallowed rather than inhaled and end up in the GI tract.
- For drugs with **significant gut absorption**, systemic levels may be difficult to distinguish between inhaled vs. swallowed portions.
- **Charcoal block PK studies** allow for a more direct analysis of the lung dose contribution in systemic circulation by eliminating the GI tract dose contribution.

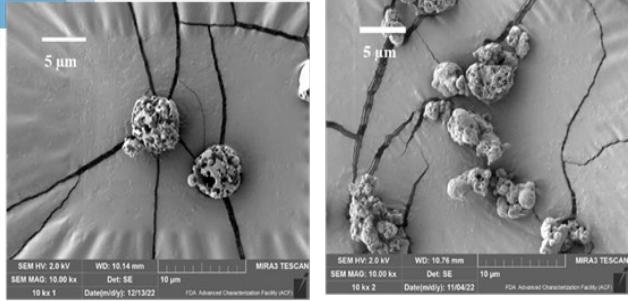


In Vivo Charcoal Block PK BE Study Considerations

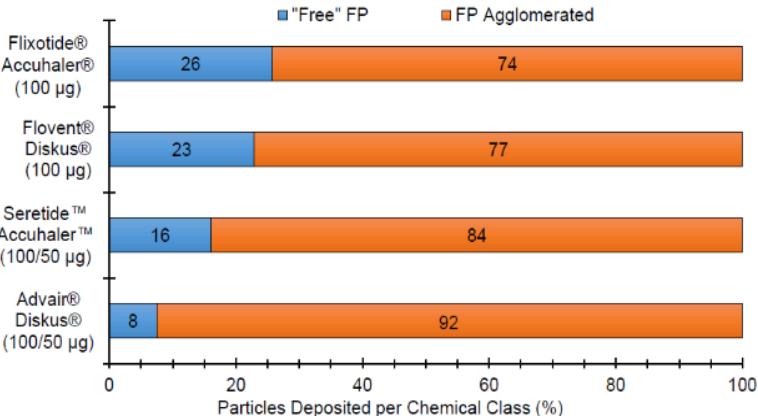


- **PSG Recommendations:**
 - Similar to PK BE study in many aspects.
 - *Healthy* adult male and female subjects.
 - *Minimum number of inhalations* to sufficiently characterize the PK profile with a sensitive analytical method.
 - Dose administration should follow the approved labeling instructions.
 - *Bio-IND* may be needed if the administered dose is above the maximum labeled single dose.
 - No standard for the *charcoal dose*, so the selected dose and how and when it is administered should be justified in the ANDA.
 - BE: 90% CI for the T/R ratio for AUC and C_{max} being between 80 – 125%.

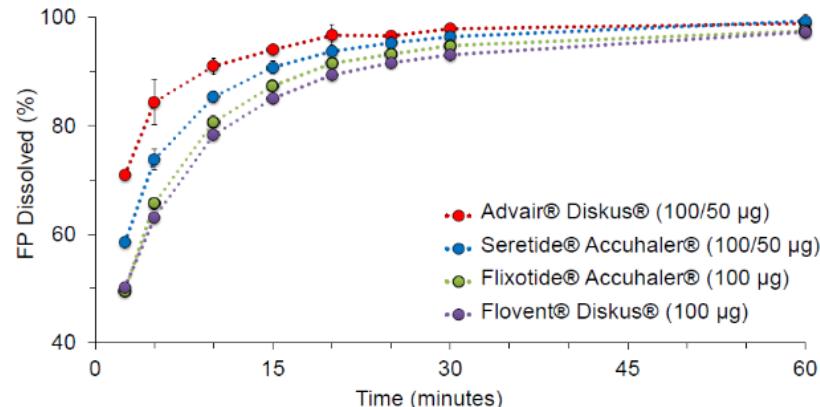
Comparative Characterization Study Considerations

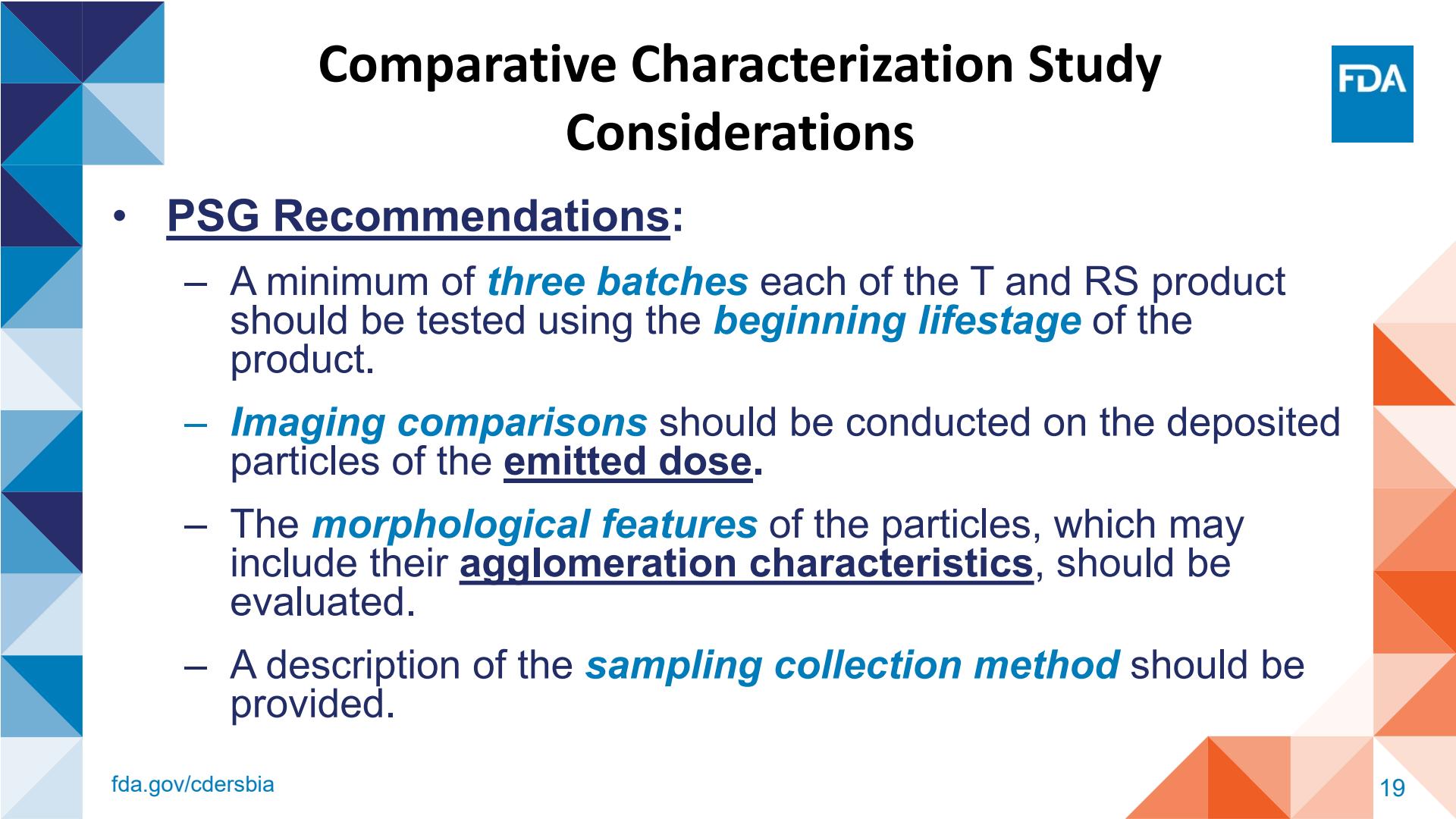


SEM images of phospholipid porous particles found in a marketed DPI (left) and MDI (right)



Microstructural differences in the deposited particle agglomerates (left) may be one potential contributing factor to performance differences, such as with dissolution performance (right).



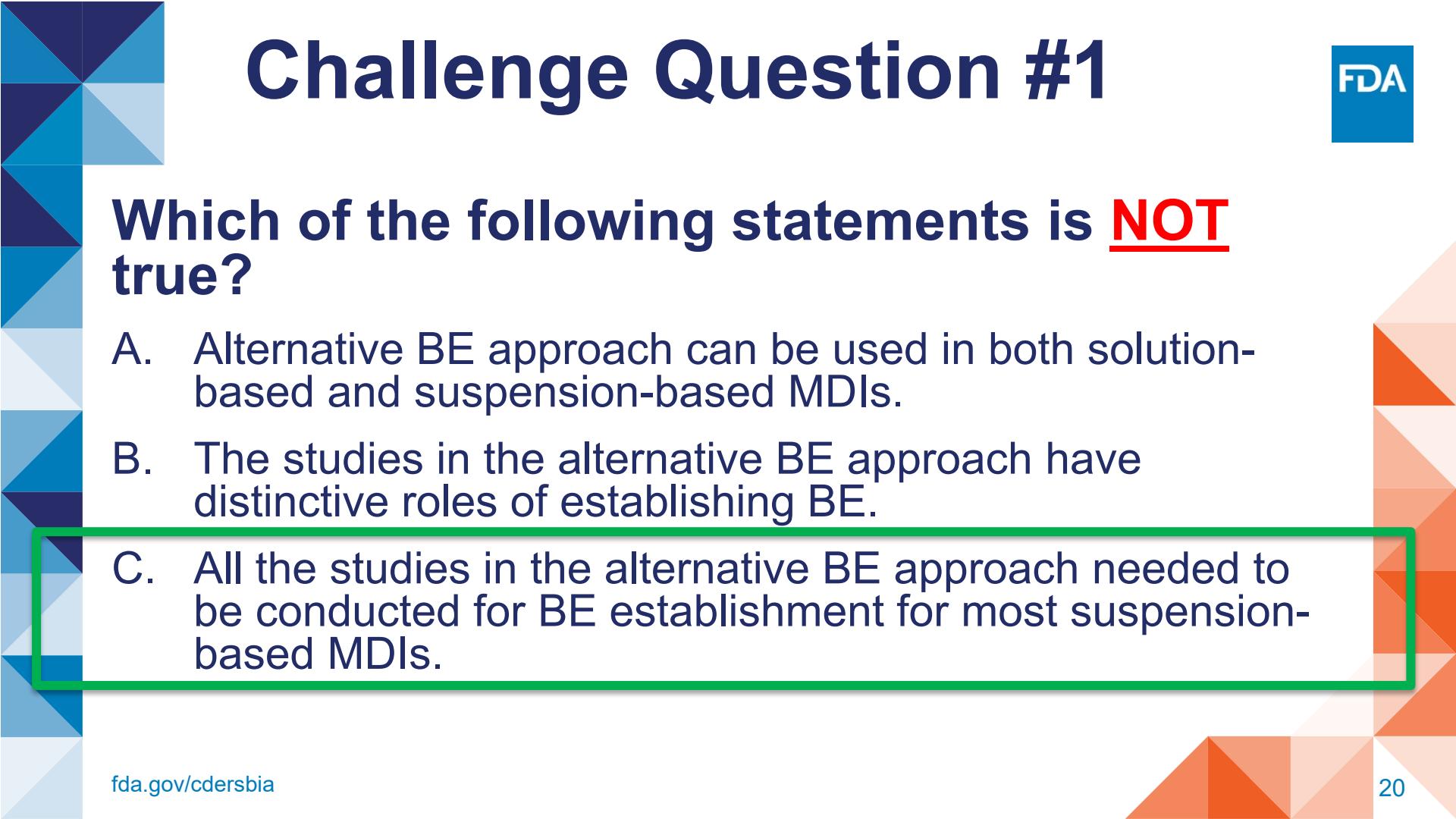


Comparative Characterization Study Considerations



- **PSG Recommendations:**

- A minimum of ***three batches*** each of the T and RS product should be tested using the ***beginning lifestage*** of the product.
- ***Imaging comparisons*** should be conducted on the deposited particles of the **emitted dose**.
- The ***morphological features*** of the particles, which may include their **agglomeration characteristics**, should be evaluated.
- A description of the ***sampling collection method*** should be provided.

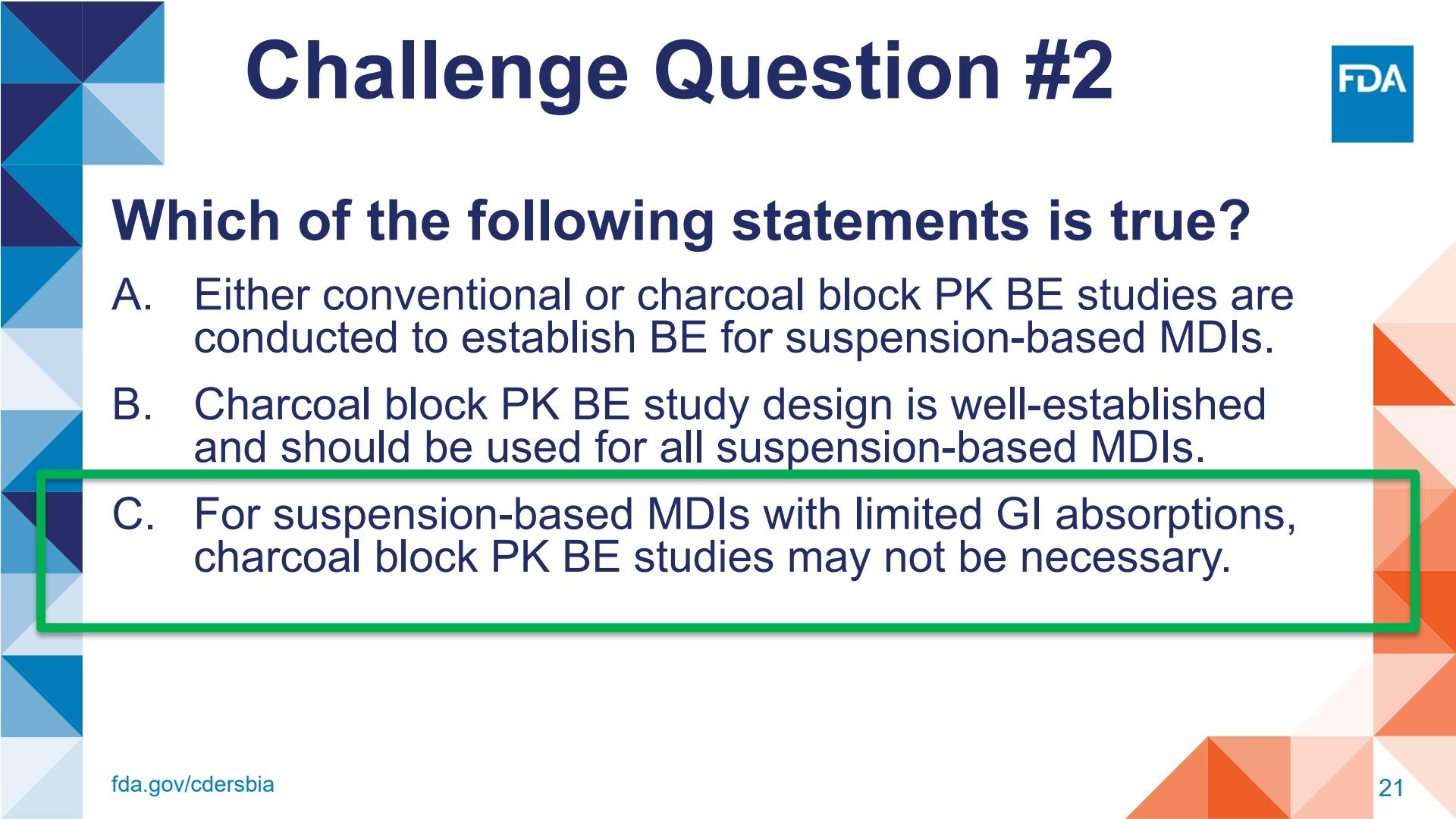


Challenge Question #1



Which of the following statements is **NOT** true?

- A. Alternative BE approach can be used in both solution-based and suspension-based MDIs.
- B. The studies in the alternative BE approach have distinctive roles of establishing BE.
- C. All the studies in the alternative BE approach needed to be conducted for BE establishment for most suspension-based MDIs.



Challenge Question #2



Which of the following statements is true?

- A. Either conventional or charcoal block PK BE studies are conducted to establish BE for suspension-based MDIs.
- B. Charcoal block PK BE study design is well-established and should be used for all suspension-based MDIs.
- C. For suspension-based MDIs with limited GI absorptions, charcoal block PK BE studies may not be necessary.

Summary



- The challenges with conducting **CCEP BE studies** can lead to higher costs and longer drug development timelines for generic developers of OIDPs.
- To address these challenges, FDA has explored ***in vitro, in vivo, and in silico study designs*** through GDUFA-funded research initiatives to identify ***alternative approaches*** that can be used in lieu of the CCEP BE study for establishing local drug delivery equivalence.
- Following completion of the **FDA-CRCG workshop** on alternative BE approaches for OIDPs in 2023, FDA has utilized the input received from industry and academic attendees to aid the development of several ***PSGs for suspension-based MDIs***.
- These ***developed PSGs*** present FDA's efforts to expand alternative BE approaches beyond just solution-based MDIs and highlight the ***additional study considerations*** needed when applying alternative BE approaches to specific drug products.



Questions?

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