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**U.S. FOOD & DRUG  
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## **BIOMARKER QUALIFICATION PROGRAM EDUCATIONAL MODULE SERIES—MODULE 1**

# **BIOMARKER TERMINOLOGY: SPEAKING THE SAME LANGUAGE**

**Shashi Amur, Ph.D.**

Scientific Lead, Biomarker Qualification Program, Office of Translational Sciences, Center for Drug Evaluation and Research, FDA



# BEST Resource: Harmonizing Terminology



- Created by the NIH-FDA Biomarker Working Group
- A glossary of terminology and uses of biomarkers and endpoints in basic biomedical research, medical product development, and clinical care

Available at: <http://www.ncbi.nlm.nih.gov/books/NBK326791/>



# BIOMARKER TERMINOLOGY

- **Definition of a Biomarker**
- **Types of Biomarkers**
- **Categories of Biomarkers**
- **Biomarker Validation**
- **Biomarker Qualification**
- **Context of Use**



# WHAT IS A BIOMARKER?

A defined characteristic that is measured as an indicator of normal biological processes, pathogenic processes, or responses to an exposure or intervention, including therapeutic interventions.

**Types:** Molecular, histologic, radiographic, and physiologic characteristics are types of biomarkers.

Examples:

- Blood glucose (molecular)
- Tumor size (radiographic)
- Blood pressure (physiologic)



# DIFFERENT CATEGORIES OF BIOMARKERS





# PROGNOSTIC BIOMARKERS



Can be used to select patients with greater likelihood of having a disease-related endpoint event or a substantial worsening in condition in clinical trials.

For example: **Total kidney volume**, to select patients with autosomal dominant polycystic kidney disease at high risk for progressive decline in renal function for inclusion in interventional clinical trials.



# SURROGATE ENDPOINT

An endpoint that is used in clinical trials as a substitute for a direct measure of how a patient feels, functions, or survives.

## Validated Surrogate Endpoint

Supported by a clear mechanistic rationale and clinical data providing strong evidence that an effect on the surrogate predicts a clinical benefit; therefore, such endpoints can be used to support traditional approval without the need for additional efficacy information.

**Example: Hemoglobin A1C reduction in diabetes clinical trials**

## Reasonably Likely Surrogate Endpoint

Supported by clear mechanistic and/or epidemiologic rationale but with insufficient clinical data to show that it is a validated surrogate endpoint; such endpoints can be used for accelerated approval for drugs or expedited access for medical devices.

**Example: Radiographic evidence of tumor shrinkage in some cancer types**

## Candidate Surrogate Endpoint

A surrogate under evaluation for its ability to predict clinical benefit.



# ANALYTICAL AND CLINICAL VALIDATION

**Analytical Validation: Ensures specificity, accuracy, precision, and other characteristics of biomarker test or device**

Establishing that the performance characteristics of a test, tool, or instrument are acceptable in terms of its sensitivity, specificity, accuracy, precision, and other relevant performance characteristics using a specified technical protocol (which may include specimen collection, handling, and storage procedures).

**Clinical Validation: Ensures the test or device performs as intended**

Establishing that the test, tool, or instrument acceptably identifies, measures, or predicts the concept of interest.

Concept: In a regulatory context, the concept is the aspect of an individual's experience or clinical, biological, physical, or functional state that the assessment is intended to capture (or reflect).





# BIOMARKER QUALIFICATION

A conclusion, based on a formal regulatory process, that within the stated context of use, a biomarker can be relied upon to have a specific interpretation and application in medical product development and regulatory review.

A biomarker, once qualified for a particular context of use, will be publicly available and can be applied in any drug development program for the qualified context of use.



## CONTEXT OF USE

A statement that fully and clearly describes the way the biomarker is to be used and the drug development-related purpose of the use.





# BIOMARKER TERMINOLOGY

By speaking the same “biomarker language,” we can enhance medical product development and may be able to get new treatments to patients sooner.

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