

## Stats Section- ABP Specs

*The outline below is taken directly from the ABP Content Outline for Pediatric Endocrinology. These are the Specs used until 10/1/2023 (more detail than the most recent version)*

**Note: Slide numbers are approximate**

### Core Knowledge in Scholarly Activities

#### A. Principles of Use of Biostatistics in Research

1. Types of variables
  - a. Distinguish types of variables (eg, continuous, categorical, ordinal, nominal)  
*Slides 3-5*
  - b. Understand how the type of variable (eg, continuous, categorical, nominal) affects the choice of statistical test *Slides 21-26*
2. Distribution of data *Slides 6-12*
  - a. Understand how distribution of data affects the choice of statistical test
  - b. Differentiate normal from skewed distribution of data
  - c. Understand the appropriate use of the mean, median, and mode
  - d. Understand the appropriate use of standard deviation
  - e. Understand the appropriate use of standard error of the mean
3. Hypothesis testing *Slides 13-15*
  - a. Distinguish the null hypothesis from an alternative hypothesis
  - b. Interpret the results of hypothesis testing
4. Statistical tests *Slides 15-25*
  - a. Understand when to use and how to interpret the chi square test
  - b. Understand when to use and how to interpret tests comparing continuous variables between two groups (eg, t test, Mann Whitney U)
  - c. Understand when to use and how to interpret tests comparing continuous variables between three or more groups (eg, ANOVA, Kruskal-Wallis)
  - d. Understand when to use paired tests
  - e. Understand the appropriate use of parametric versus nonparametric tests
  - f. Interpret a p value
  - g. Interpret a p value when multiple comparisons have been made *Slide 41*
  - h. Interpret a confidence interval
  - i. Identify a type I error
  - j. Identify a type II error
5. Measurement of association and effect *Slides 27-37*
  - a. Understand how to interpret relative risk and absolute risk
  - b. Understand how to interpret odds ratio
  - c. Understand how to interpret number needed to treat or harm
  - d. Understand how to interpret hazard ratio
  - e. Understand when to use and how to interpret correlation coefficient *Slide 25*
6. Regression
  - a. Understand when to use and how to interpret regression analysis (eg, linear, logistic) *Slide 26*
  - b. Understand when to use and how to interpret survival analysis (eg, Kaplan Meier).  
*Slides 39-40*
7. Diagnostic tests *Slides 42-50*
  - a. Recognize the importance of an independent "gold standard" in evaluating a diagnostic test
  - b. Interpret sensitivity and specificity
  - c. Interpret positive and negative predictive values

- d. Understand how disease prevalence affects the positive and negative predictive value of a test
  - e. Interpret a receiver operating characteristic curve
- 8. Systematic reviews and meta-analysis [Slide 51](#)
  - a. Understand the purpose of a systematic review
  - b. Understand the advantages of adding a meta-analysis to a systematic review
  - c. Interpret the results of a meta-analysis
- B. Principles of Epidemiology and Clinical Research Design
  - 1. Assessment of study design, performance and analysis (internal validity)
    - a. Recognize and understand the strengths and limitations of a cohort study, case control study, and randomized controlled clinical trial [Slides 53-56](#)
    - b. Recognize the use and limitations of surrogate endpoints [Slide 59](#)
    - c. Understand the use of intent-to-treat analysis [Slide 60](#)
    - d. Understand how sample size affects the power of a study [Slide 16](#)
  - 2. Assessment of generalizability (external validity) [Slide 69](#)
    - a. Understand how nonrepresentative samples can bias results
    - b. Assess how the data source (eg, diaries, billing data, discharge diagnostic code) may affect study results
  - 3. Bias and confounding [Slides 57-58, 60, 66-68](#)
    - a. Identify common strategies in study design to avoid or reduce bias
    - b. Identify common strategies in study design to avoid or reduce confounding
  - 4. Causation
    - a. Understand the difference between association and causation [Slide 61](#)
  - 5. Incidence and prevalence [Slide 70](#)
    - a. Distinguish disease incidence from disease prevalence
  - 6. Screening
    - a. Understand factors that affect the rationale for screening for a condition or disease (eg, prevalence, test accuracy, risk benefit, disease burden, presence of a presymptomatic state)
  - 7. Cost benefit, cost effectiveness, and outcomes [Slide 76](#)
    - a. Interpret cost-effectiveness ratios
    - b. Distinguish costs from charges
    - c. Understand quality-adjusted life years
  - 8. Measurement [Slides 71-75](#)
    - a. Understand the types of validity that relate to measurement (eg, face, construct, criterion, predictive, content)
    - b. Distinguish accuracy from precision
    - c. Understand when to use and how to interpret a kappa coefficient
- C. Ethics in Research
  - 1. Professionalism and misconduct in research [Slide 78-79](#)
    - a. Identify and manage potential conflicts of interest in the funding, design, and/or execution of a research study
    - b. Identify various forms of research misconduct (eg, plagiarism, fabrication, falsification)
    - c. Know how, and to whom, to report concerns of research misconduct
  - 2. Principles of research with human subjects [Slides 80-84](#)
    - a. Understand and contrast the functions of an Institutional Review Board and a Data Safety Monitoring Board
    - b. Recognize the types of protections in designing research that might be afforded to children and other vulnerable populations
    - c. Understand the federal regulatory definitions regarding which activities are considered research and what constitutes human subjects research

- d. Understand the federal regulatory definition of minimal risk and apply this to research involving children
  - e. Understand the ethical considerations of study design (eg, placebo, harm of intervention, deception, flawed design)
- 3. Principles of consent and assent *Slides 85-86*
  - a. Understand what constitutes informed consent in research
  - b. Distinguish between consent and assent in research involving children
- D. Quality Improvement
  - 1. Design of a Project *Slides 88-90*
    - a. Understand various models of quality improvement and recognize that all utilize a data-informed, iterative process using tests of change to achieve a stated aim
    - b. Understand that the aim of any quality improvement project should be specific, measurable, achievable, realistic, and time-limited
    - c. Understand strategies to optimize identification of key drivers and interventions to achieve a specific aim
    - d. Understand tools to facilitate completion of quality improvement work, including key driver diagrams and process maps
    - e. Understand each phase of a Plan-Do-Study-Act (PDSA) cycle
  - 2. Data and Measurement *Slides 91-100*
    - a. Differentiate between process, outcome, and balancing measures
    - b. Interpret a run chart and identify shifts, trends, and outliers in data
    - c. Differentiate between a run chart and a control chart
    - d. Differentiate between common cause and special cause variation