



GFR Decline as an Endpoint for Clinical Trials in CKD

A Scientific Workshop Co-sponsored by the National Kidney Foundation
and the US Food and Drug Administration

AGENDA FOR WEBINAR - MONDAY, DECEMBER 3

Welcome and Introductions (15 minutes)

Andrew Levey, Kerry Willis, and Norman Stockbridge

Plenary Session 1 – Perspectives (105 minutes)

Goals of the Conference

Andrew Levey

Regulatory Perspective

Aliza Thompson

Clinical Perspective

Edmund Lewis, Alfred Cheung, and Dick de Zeeuw

Overall Analytical Plan

Josef Coresh

Plenary Session 2 - Observational Study Data (60 minutes)

Josef Coresh

Plenary Session 3 – Clinical Trial Data (60 minutes)

Lesley Inker

Plenary Session 4 – Simulation Studies (60 minutes)

Tom Greene

The Proposal – Surrogate Endpoints, Trial Designs, and Caveats (60 minutes)

Andrew Levey