

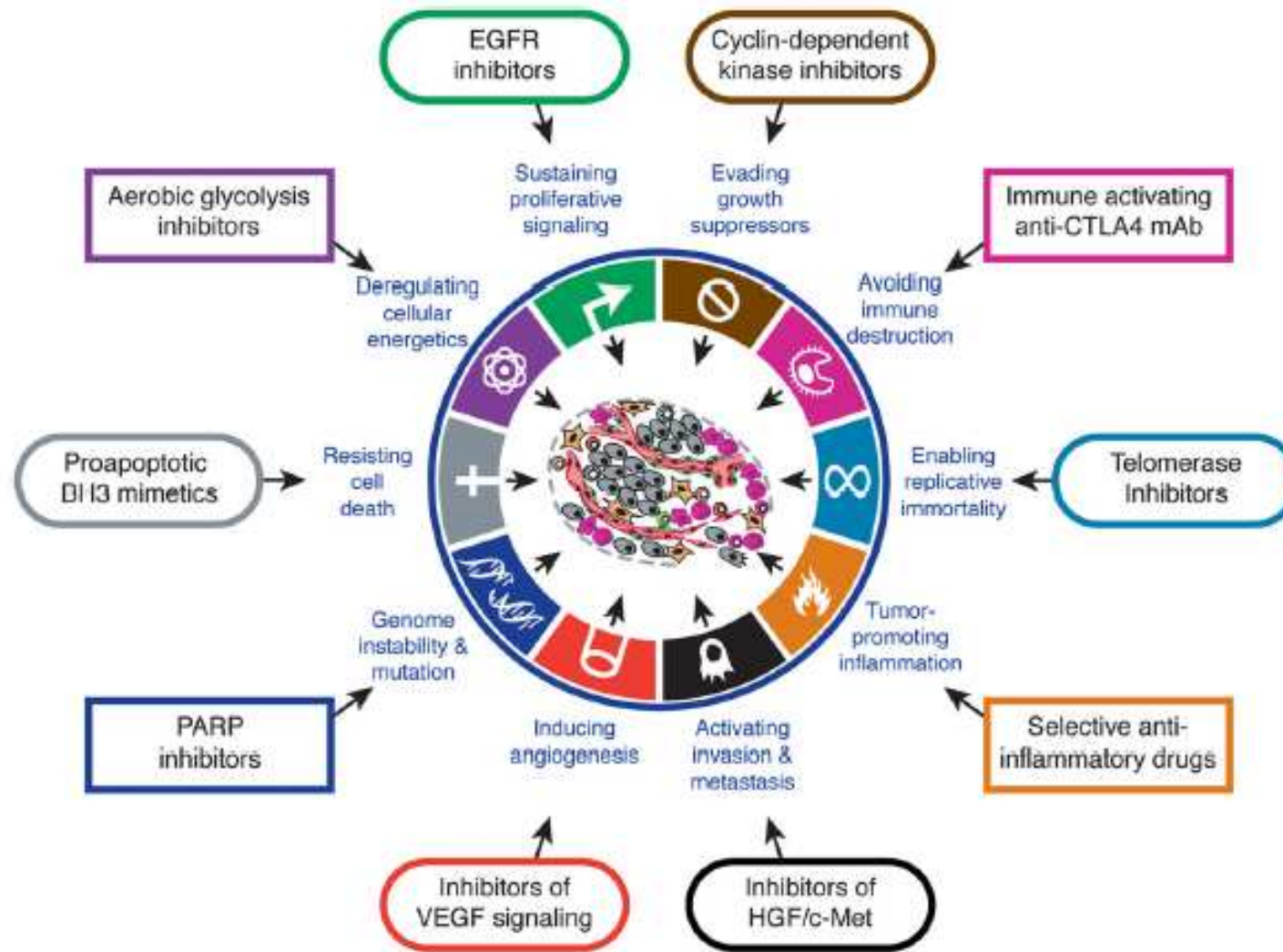
# **PERSONALIZED CANCER CARE: WHAT DOES IT MEAN?**

***THE ASPIRATION TO BASE A TREATMENT ON THE UNIQUE  
BIOLOGICAL FEATURES OF A PATIENT'S DISEASE THROUGH THE  
IDENTIFICATION OF VALIDATED BIOMARKERS OF  
RESPONSE/RESISTANCE TO THERAPY***

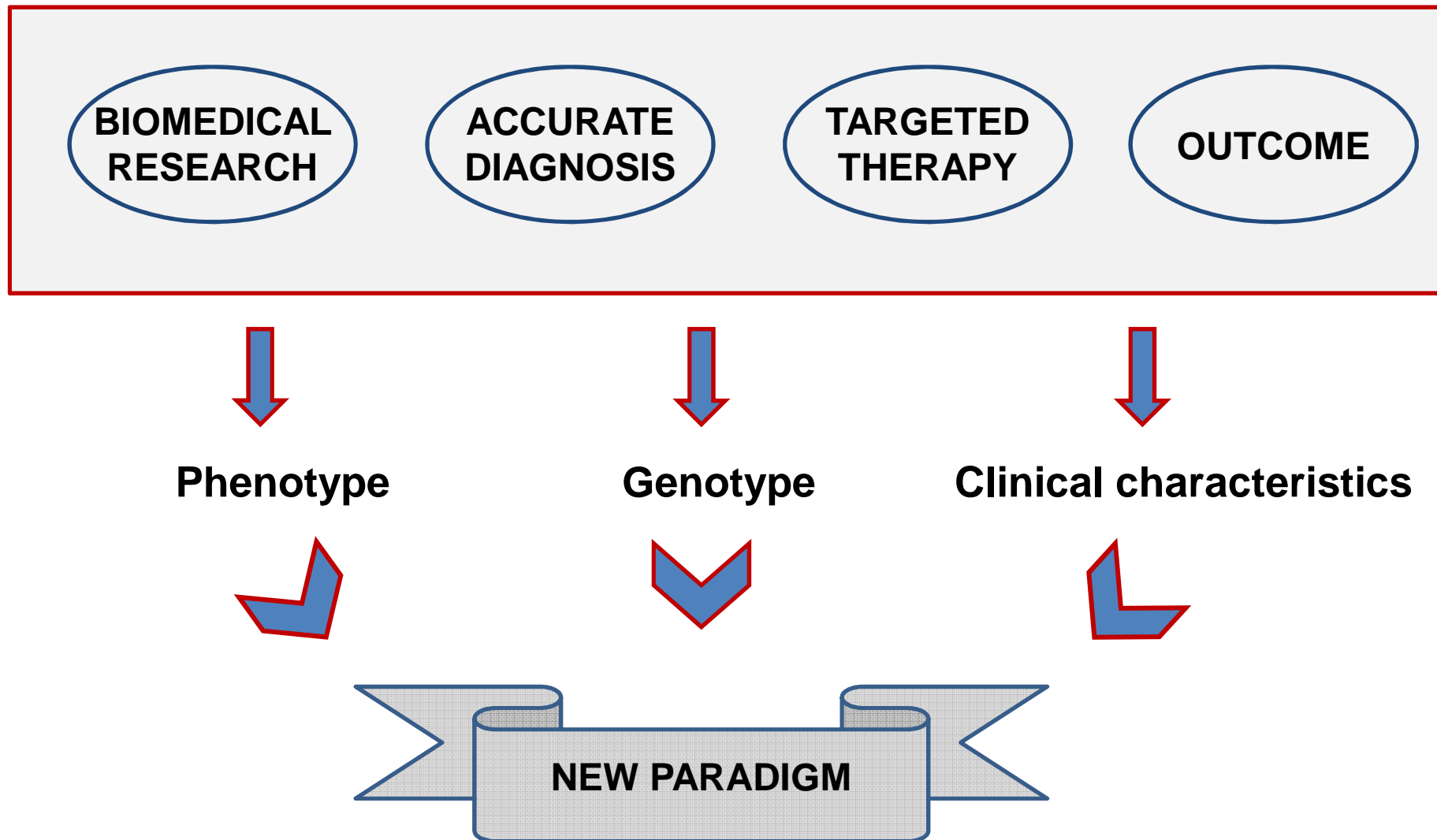
## **AIMS**

- ☐ ***TO IMPROVE DRUG EFFICACY***
- ☐ ***TO AVOID INAPPROPRIATE DRUG EXPOSURE (PRIMARY RESISTANT TUMORS)***
- ☐ ***TO MAXIMIZE QUALITY OF LIFE***
- ☐ ***TO SPARE UNNECESSARY COSTS***

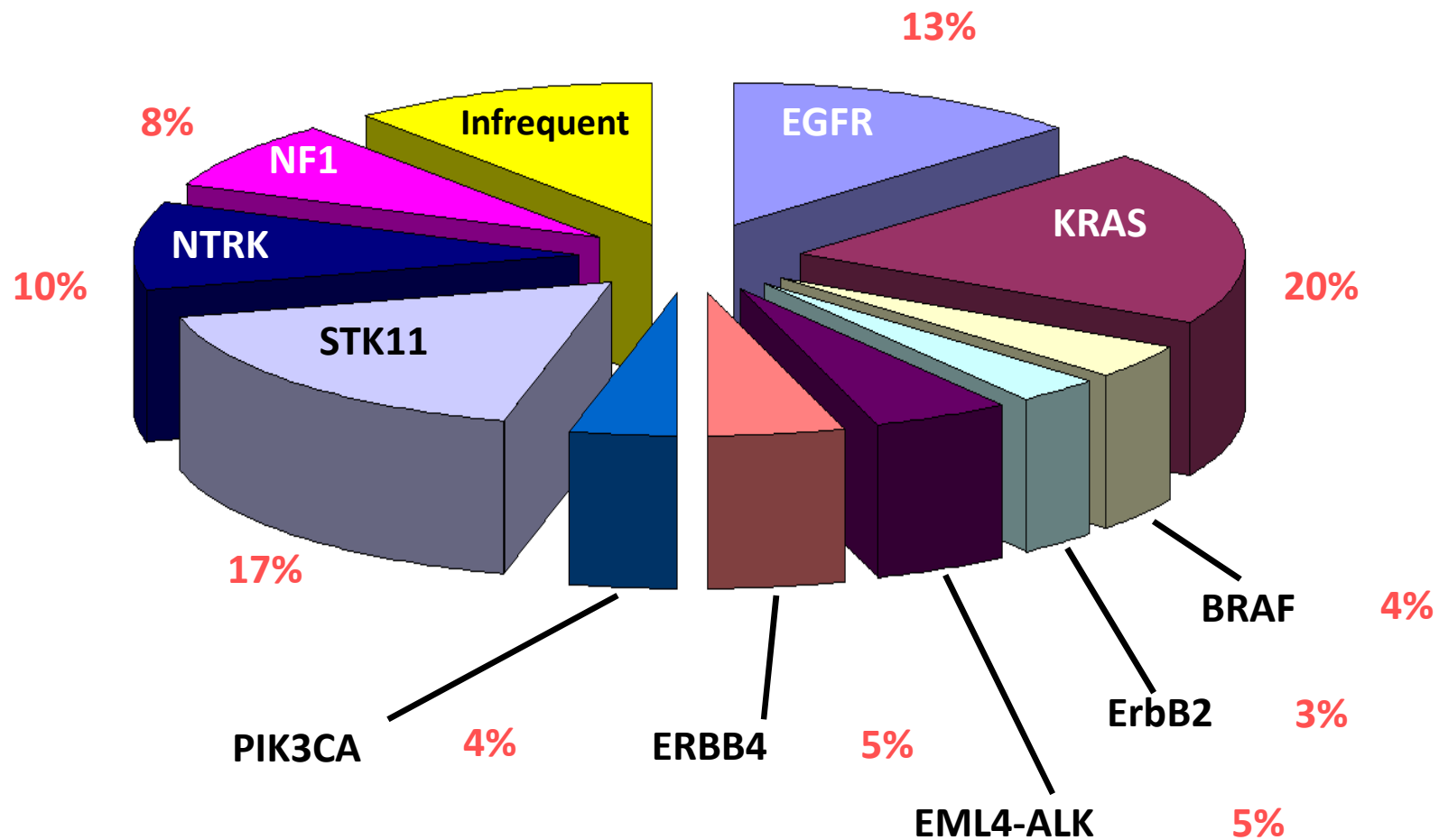
# THERAPEUTIC TARGETING OF THE HALLMARKS OF CANCER



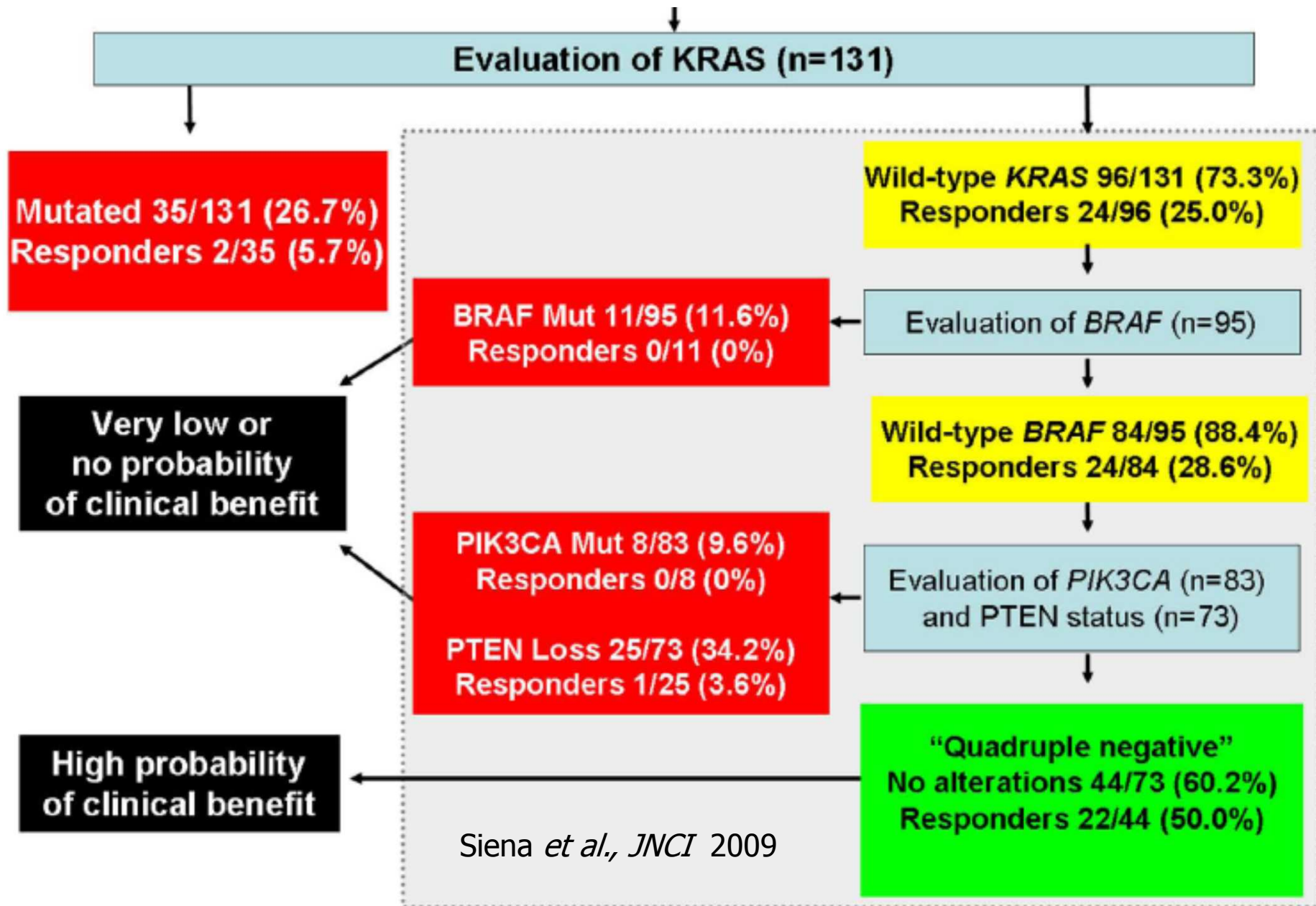
# PRECISION MEDICINE IN ONCOLOGY



# GENETIC MUTATIONS IN LUNG ADENOCARCINOMAS



## RESULTS ON 131 PATIENTS



# MULTIPLE MOLECULAR ALTERATIONS EFFECT ON RESPONSE TO anti-EGFR mAbs

Because of the occurrence of **multiple molecular alterations** within the same tumor, we investigated **a cohort of 131 patients treated with EGFR-targeted moAbs in the chemorefractory setting** by separating patients according to the actual number of molecular abnormalities within the same tumor (i.e. none vs 1 vs  $\geq 2$  alterations) among *KRAS*, *BRAF*, *PIK3CA* mutations and loss of PTEN.

The **probability of response** was:

- ✓ **50.0%** (22/44) among patients with no alterations,
  - ✓ **4.2%** (2/47) among patients with 1 alteration and
  - ✓ **0%** (0/23) for patients with  $\geq 2$  alterations,
- } ( $p < 0.0001$ )

