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Making Cancer History®

Importance of Clinical Trials

Filip Janku

Investigational Cancer Therapeutics (Phase I Clinical Trials Program) MD Anderson Cancer Center Houston, TX



Why Do We Need Clinical Trials

- Clinical trials answer two important questions
 - Does the new treatment work?
 - Is the new treatment safe?

How Do We Do Clinical Trials

- Clinical trials are usually carried out in "phases"
 - Phase I: What is the safe dose?
 - Phase II: Does the treatment work?
 - Phase III: Is the treatment better than existing options?
- Timeline: 10-15 years

Clinical Trials: Pros

• For mankind

- Increase in knowledge about particular disease and its therapy
- Development of new therapies
- Prove of efficacy or lack of thereof
- Identification of potential significant side effects

• For individual patient

- Access to new therapies, which are not commercially available
- Expansion of therapeutic options
- Standardized protocol-driven therapy
- Some studies in cancer patients suggested that patients on clinical trials tend to have better outcomes compared to patients treated outside of trials

Clinical Trials: Cons

- For mankind
 - None

• For individual patient

- Need to meet all qualifying criteria, which are usually not flexible
- Less flexible and often more intense schedule
- Travel, financial consequences and time commitment
- Possible risk of unknown/unexpected side effects

Why Should I Consider Clinical Trials

 Results of clinical trials are important not only for developing new therapies

 Clinical trials can provide necessary evidence to convince payers to reimburse new and effective therapies

Clinical testing is necessary tool to make the progress happen

Strategies for Clinical Trials in ECD

- Prognosis and outcomes have dramatically improved; however, overall there is still room for improvement
- We have relatively limited therapeutic armamentarium
- We have limited resources (patients, finances) and large number of questions, which need to be answered

Strategies for Clinical Trials in ECD

• **Phase I**: Access for ECD patients to these studies, which are often limited to conventional cancers

• Phase II:

- "Basket studies": clinical trials for patients with any cancer or histiocytosis with certain unifying feature (e.g. vemurafenib in patients with *BRAF* mutation)
- ECD specific phase II studies: because of limited number of patients this approach should be reserved for promising therapies with high likelihood of FDA/EMA approval
- Phase III: not feasible in ECD

Where Can I Learn About Clinical Trials?

- ECD Global Alliance Website
- Clinicaltrials.gov

• Care Centers

Examples of Clinical Trials for ECD Patients

- *BRAF* mutation positive
 - NIH: dabrafenib/trametinib
 - Multicenter: my pathway (vemurafenib)
 - Multicenter: dabrafenib/trametinib
 - Multicenter: BVD-523
 - Multicenter: PLX8394
- BRAF mutation negative
 - Multicenter: BVD-523 (if MAP2K1 mutation present)
 - Multicenter: PLX8394
 - Single Center: everolimus/anakinra

Take Home Message

Clinical Trials are Part of Standard of Care in ECD