Only one step from tailored oncological therapies

Pawel Krawczyk

The personalization of oncological therapy is becoming a reality thanks to the development of molecular biology techniques (e.g. next generation sequencing, liquid biopsy) and the discovery of new molecularly targeted therapies and immunotherapeutics. Genetic laboratories and clinical centers, which form international networks as well as global, early phase clinical trials make it possible to match treatments to a particular patient. The benefit of such a procedure is currently difficult to estimate, but it seems that another path for the development of oncology is not possible.

Key words: oncology, molecularly targeted therapies, immunotherapies, gene sequencing, liquid biopsy

Introduction

Tailored oncological treatment based on molecularly targeted therapies and immunotherapies play increasing role in the multidisciplinary approach to cancer patient care. Treatment personalization extends the scope of therapy and extends the survival of patients. An example of lung cancer treatment personalization is our 57-year-old male patient with advanced lung adenocarcinoma and activating EGFR gene mutations, as well as PD-L1 expression. As the methods of treatment, we used segmentectomy (diagnostic manner), hemipelvectomy of metastases in the right hip joint with postsurgical radiotherapy, surgical cytoreduction, and six lines of systemic treatment which were based on standard cytostatics and novel personalized agents. The presence of the coexistence of rare mutations in exon 18 of the EGFR gene (G719X and G709X) in tumor cells allowed the use of the first-generation (erlotinib) of tyrosine kinase inhibitor (TKI) of EGFR (epidermal growth factor inhibitor). The appearance of the Thr790Met mutation in exon 20 of the EGFR gene, in circulating tumor DNA (ct-DNA) from peripheral blood, during the progression to erlotinib allowed the use of the third generation of EGFR-TKI (osimertinib). The presence of PD-L1 expression on tumor cells justified the attempt to apply immunotherapy (nivolumab — anti-PD-1 antibody). The patient also received a tyrosine kinase inhibitor of VEGFR, FGFR and PDGFR (nintedanib) in combination with chemotherapy. As a result, we have achieved long-term overall survival with long progression-free periods during individual therapy [1].

Of course, such a therapeutic approach could be regarded as unattainable for larger groups of patients. Personalized therapies are available only in developed countries with access to modern genetic methods of cancer diagnosis and to clinical trials with novel drugs. Today, personalized therapies could be used in a small percentage of patients with a genetic predisposition to the treatment. Moreover, cancer cells can develop resistance to even the best molecularly targeted therapies and immunotherapies. Tumors are also molecularly heterogeneous, so it is difficult to find for our patients appropriate and permanently effective personalized therapy. However, the development of new molecular methods (especially next generation sequencing and liquid biopsy) and the discovery of new drugs investigated in clinical trials raise the hope that precise oncology has arrived.

Next generation sequencing and liquid biopsy in qualification to personalized treatment in cancer patients

Next generation sequencing (NGS) is a new DNA sequencing technology which has revolutionized genomic research. Using NGS, the whole genome of cancer cells
(whole genome sequencing, WGS) can be sequenced within a single day, and the price of the examination has dropped so much that WGS could be available to even individual patients. In contrast, the previous Sanger direct technology required over a decade in order to map the human genome. NGS is increasingly translated into routine clinical practice, especially in oncology and in pediatrics. NGS could be used to sequence whole genome or specific genome areas, including coding sequences (whole exome sequencing, WES) and individual genes (e.g. oncogenes with “druggable” mutations). There are a number of different NGS platforms using different sequencing technologies. In a very simplistic way, all NGS technologies use three stages [2, 3]:

1. Library preparation with DNA fragmentation and in vitro adaptor ligation.
2. Clonal amplification using an emulsion or bridge PCR (polymerase chain reaction).
3. Cyclic sequencing of millions of small fragments of DNA in parallel using pyrosequencing, sequencing by ligation or sequencing by synthesis.

WES or sequencing of selected genes are most commonly used in the personalization of oncological treatment. The sequencing of frequently mutated genes in tumor cells in particular malignancies, is the most meaningful, cost-effective and rapid method to qualify to molecularly targeted drugs. These mutations should cause changes in proteins involved in intracellular pathways which could be blocked by molecularly targeted drugs. WES may be a useful method to evaluate the number of mutations in tumors (tumor mutation burden) in qualification to immunotherapy (a large number of mutations results in the increased number of tumor-specific antigens and increased efficacy of immunotherapy) [2, 3].

Several diagnostic companies propose the sequencing of selected genes to qualify individual patients for molecularly targeted therapies (e.g. FoundationOne, OncoDNA, ONCOMPASS, Caris, Guardant 360). NGS is not only offered to detect gene mutations (indels, substitutions, single nucleotide transcriptional variants (SNVs), but also for gene copy number variations (CNV) and chromosome translocation (rearrangement) analysis or epigenetic analysis (promoter genes methylation). In offers, NGS analysis is supplemented by simpler genetic methods, e.g. immunohistochemistry (IHC) for assessment of protein expression, fluorescence in situ hybridization (FISH) for chromosomal abnormality analysis, and real-time PCR technique for single mutation analysis. The patients’ report should consist not only of the result of genetic tests, but also of the the analysis of the availability of personalized treatment (chemotherapy, molecularly targeted therapies, immunotherapy). Such a report takes into account therapies registered for treatment of particular cancers, therapies registered for other indications (potentially useable due to the occurrence of genetic alterations), and drugs in clinical trials (often available in distant locations) [2, 3].

The possibility of genetic diagnostics and qualification for molecularly targeted therapies and immunotherapy has considerably expanded through the introduction of analysis of genetic materials from peripheral blood (liquid biopsy), with highly sensitive molecular methods (e.g. some NGS techniques, droplet digital PCR and BEAMing PCR). Circulating tumor DNA (ctDNA) is now being extensively studied, as it is a noninvasive “real-time” biomarker that can provide diagnostic and prognostic information before, during treatment and at progression. These include DNA mutations, epigenetic alterations and other forms of tumor-specific abnormalities such as microsatellite instability (MSI) and loss of heterozygosity (LOH). The major challenge of genetic analysis of material from liquid biopsy is assay sensitivity. Frequent EGFR gene mutations (deletions in exon 19 and Leu858Arg substitution in exon 21) are detected in liquid biopsies of non-small cell lung cancer (NSCLC) patients with a sensitivity of 60–75%. Therefore, the tumor tissue or cells are the most important material for examination of these mutations during qualification of NSCLC patients to EGFR-TKI therapy. However, the use of liquid biopsy for detection of EGFR mutations, responsible for EGFR-TKI resistance during disease progression (e.g. Thr790Met substitution in exon 20), extends the ability of patient qualification to third generation EGFR-TKI (osimertinib). Moreover, analysis of genetic materials from liquid biopsy allows determination of the moment of molecular progression (the appearance of genetic material of cancer cells in peripheral blood). In addition, examination of genetic mutations in ctDNA allows the selection of appropriate molecularly targeted compounds. A good example is the treatment of NSCLC patients with ALK gene rearrangement using various ALK inhibitors (crizotinib, ceritinib, alectinib, lorlatinib). Recently identified ALK gene mutations are responsible for differences in the sensitivity of ALK inhibitors. Therefore, identification of the appearance of ALK mutations during ALK inhibitor treatment allows qualification to appropriate next generation drugs and selection of the appropriate treatment sequence for these drugs [4–6].

The concept of an oncology center network with access to molecular diagnosis and clinical trials with new molecularly targeted therapies

The National Comprehensive Cancer Network developed clinical practice guidelines in oncology for non-small-cell lung cancer diagnosis and treatment. These guidelines include recommendations to perform tests for EGFR mutation and ALK rearrangement detection, and also examination of BRAF V600E mutation, MET amplification, MET exon 14 skipping mutation, RET rearrangements and HER2 mu-

43
Selection of individualized treatments based on genetic analysis of paired tumor and normal tissue was performed in the global WINTHER study. In 200 patients biopsies of metastatic lesions and matched normal tissue were executed in order to obtain material for whole genome sequencing. Patients with molecular predictive factors received matched molecular targeted therapies including drugs in the opened 1st phase clinical trials. In patients without such predictive factors, oncologists used therapeutic choice based on predictive drug efficacy scoring [11].

The SHIVA clinical trials enrolled patients with recurrent and refractory to standard treatment solid tumors. Molecular profiling using the NGS technique is performed in tumor biopsy. After centralized bioinformatics analysis of the genetic results, the molecular biology board makes the treatment decision. Eligible patients are randomized to molecularly targeted therapies or to treatment at the physician’s choice. Treatment cross over is possible at the time of disease progression [12, 13].

The National Cancer Institute — MATCH Clinical Trial explores the possibilities of the introduction to routine practice of novel treatment methods based on molecular profiles of patients’ tumors. About 6000 patients with solid tumors (including rare tumors, lymphomas and myeloma) that no longer respond to standard treatment are screened with a tumor biopsy. Changes in 143 genes are diagnosed using the NGS technique. If a patient’s tumors have genetic abnormalities that match the one targeted by a drug used in clinical trial, the patients are eligible to join the treatment portion of NCI-MATCH. 30 different therapies that target 30 different genetic abnormalities are available for cancer patients taking part in this clinical trial [3, 14].

**Conclusion**

Tremendous progress in molecular biology and the development of new molecularly targeted drugs allow treatment personalization for very narrow, genetically selected groups of cancer patients. The nature of clinical trials is evolving. In the previous decade, large clinical trials of the 3rd phase have been dominant, enrolling hundreds and even thousands of patients. Currently, early phase clinical trials enroll only individual patients with a genetic predisposition to an individual treatment method. Therefore, the effectiveness of the investigated drugs is discussed on the basis of individual case reports. Patients actively search for such clinical trials, traveling around the world for treatment. Often, such an attitude prolongs their life just enough to make even one of their dreams come true.

**Conflict of interest:** none declared

Prof. Paweł Krawczyk MD, PhD  
Medical University of Lublin  
Department of Pneumonology, Oncology and Allergology
Based on the presentation at the V Annual Conference of the Nowotwory Journal of Oncology, ‘Oncological Debates,’ held in Warszawa, 7–8th April 2017

References