Detection of ROS1 rearrangement in non-small cell lung cancer: current and future perspectives

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Abstract: ROS1 rearrangement characterizes a small subset (1%–2%) of non-small cell lung cancer and is associated with slight/never smoking patients and adenocarcinoma histology. Identification of ROS1 rearrangement is mandatory to permit targeted therapy with specific inhibitors, demonstrating a significantly better survival when compared with conventional chemotherapy. Detection of ROS1 rearrangement is based on situ (immunohistochemistry, fluorescence in situ hybridization) and extractive non-in situ assays. While fluorescence in situ hybridization still represents the gold standard in clinical trials, this technique may fail to recognize rearrangements of ROS1 with some gene fusion partner. On the other hand, immunohistochemistry is the most cost-effective screening technique, but it seems to be characterized by low specificity. Extractive molecular assays are expensive and laborious methods, but they specifically recognize almost all ROS1 fusions using a limited amount of mRNA even from formalin-fixed, paraffin-embedded tumor tissues. This review is a discussion on the present and futuristic diagnostic scenario of ROS1 identification in lung cancer.

Keywords: lung, adenocarcinoma, ROS1, FISH, immunohistochemistry, NGS, rearrangement

Introduction

ROS1 is a gene encoding a receptor tyrosine kinase; it is closely related to ALK and LTK and identified in several human tumors, including non-small cell lung cancer (NSCLC).1–8

Recently, the US Food and Drug Administration approved the use of crizotinib, (Xalkori®, Pfizer Inc., New York, NY, USA) a specific small molecule inhibitor, in the therapy of ROS1 rearranged NSCLC.9–11

Detection of ROS1 rearrangement is then a critical step in the treatment of NSCLC and may be performed using different techniques, including immunohistochemistry (IHC), fluorescence in situ hybridization (FISH) and molecular extractive methods (e.g., reverse transcription-polymerase chain reaction [RT-PCR]).

Since ROS1 protein is absent in normal lung tissue and the prevalence of ROS1 rearrangement in NSCLC ranges from 0.5% to 2%, IHC appears a cost-effective screening assay, thus permitting rapid results with less cost.12–17

All the different methodologies adopted to identify ROS1 rearrangement have some advantages as well as limitations when compared to each other. In the current clinical practice, FISH represents the gold standard in light of its use in determining ROS1 positivity in clinical trials. However, several studies comparing the sensitivity and specificity of other techniques with FISH results have been published.18–36

While coordinated use of IHC and FISH testing does represent the routine practice in real-life laboratories, emerging molecular assays, including mRNA expression of...
the 3’ region over 5’ region of ROS1 gene (NanoString assay) and next-generation sequencing (NGS), could become an appealing and futuristic standard, permitting simultaneous tests for several “druggable” drivers using limited amount of tumor tissue or liquid biopsies.37–40

The ROS1 oncogene

ROS1 gene is located at chromosome 6q22 and encodes for a receptor tyrosine kinase belonging to the insulin receptor family (Figure 1). The rearrangement of ROS1 gene leads to a constitutively activated downstream signaling with oncogenic properties. ROS1 rearrangement was firstly detected in a glioblastoma cell line,41 but was also reported in cholangiocarcinoma, gastric adenocarcinoma, ovarian serous carcinoma, colonic adenocarcinoma, epithelioid hemangioendothelioma and spitzoid melanocytic tumors.2–7,34 The fusion gene partners of ROS1 comprise several genes, including CD74, EZR, FIG1, CCD6, KDELR2, SLC34A2, LRIG3, SLC34A–ROS1 (1q21.2) - ROS1 (6q22.1), SDC4 (6q22.1), TPM3 (1q21.2) - ROS1 (6q22.1), SDC4 (20q12) - ROS1 (6q22.1), SLC34A2 (4q15.2) - ROS1 (6q22.1), CD74 (5q32) - ROS1 (6q22.1) and FIG (GOPC) - ROS1,(del(6)(q22;q25.3)).

All rearrangements involve the 3’ region of the kinase domain of ROS1 to the 5’ region of the partner gene. ROS1 rearrangements characterize about 0.5%–2% of unselected NSCLCs.

The seminal work by Rikova et al42 first identified ROS1 and ALK fusion genes in NSCLC using a phosphoproteomic approach, characterizing tyrosine kinase signaling in tumor cell lines and samples.

Although not every single fusion transcript has been evaluated for its oncogenic potential thus far, the preservation of the entire ROS1 kinase domain, whatever the partner gene may be, should be sufficient to drive carcinogenesis.42 Once constitutively activated, ROS1 signaling mainly rests on extracellular regulated MAP kinase (ERK), phosphatidylinositol-4,5-bisphosphate 3-kinase catalytic subunit alpha (PI3K)/mechanistic target of rapamycin (mTOR) and Janus kinase (JAK)-signal transducer and activator of transcription (STAT) intracellular pathways (Figure 2).1

Of note, ROS1 and ALK are evolutionarily conserved and share >80% sequence identity within their ATP-binding sites.9,12 These latter observations are strongly sustained by the clinical activity of crizotinib in both ROS1- and ALK-driven NSCLC and allow to approach them, with special regards to their inhibition by novel compounds, in a parallel way. Nevertheless, biologic differences between the two oncogenes, still not objectified at the cellular level, ostensibly

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Figure 1 ROS1 gene is located at 6q22 and its rearrangement involves several fusion gene partners. Abbreviations: NSCLC, non-small cell lung cancer; TK, tyrosin kinase; TM, transmembrane; WT, wild type.
account for diversities observed in preclinical models and in the clinics for ROS1- and ALK-rearranged NSCLC.

The well-defined oncogenic role that ROS1 plays in lung cancer carcinogenesis, as well as the major clinical improvement generated by the ALK/MET/ROS1 inhibitor crizotinib (which, indeed, recently received the US Food and Drug Administration approval for ROS1-rearranged cases too) make the detection of ROS1 rearrangement a crucial task in the field of molecular medicine.\(^{11}\) Moreover, the relatively long survival of ROS1-positive NSCLC patients stimulates researchers and clinicians to find new therapeutic strategies to maintain an active ROS1 adduction. As evidenced in EGFR-mutated and ALK-rearranged NSCLC, overcoming resistance to first-generation targeted treatments with novel molecules is achievable and dramatically fruitful, at least in significant quotes of patients.

This review is focused on the current status of the methodologies used to detect ROS1 oncogene rearrangement in NSCLC. Presently, in situ (IHC and FISH) and extractive, non-in situ assays are available worldwide with important differences concerning advantages and limitations. In particular, a summary of the studies comparing IHC and FISH results is reported and the future scenario of ROS1 detection in routine practice along with the advent of promising non-in situ techniques is reported. A present and futurist algorithmic approach to ROS1 rearrangement in NSCLC is briefly discussed.

**Clinical implications of ROS1 detection**

The identification of ROS1 rearrangement is of crucial interest in NSCLC patients due to the therapeutic consequences it generates. Cytotoxic treatment with pemetrexed–platinum doublets shows special activity in ROS1-positive patients,\(^{43}\) as seen for lung tumors driven by either ALK or RET rearrangements.\(^{44,45}\) Nevertheless, ROS1 positivity appears to be the best biomarker of pemetrexed activity in terms of both response rate and long-term outcomes, with estimations of median progression-free survivals of around 7 months.\(^{46-48}\)

After having demonstrated significant activity in ALK-driven diseases,\(^{49}\) the tyrosine kinase inhibitor crizotinib, firstly developed as an anti-MET molecule, was shown to harbor relevant activity in ROS1-rearranged NSCLC too.\(^{9}\) As seen for the first- and second-generation EGFR inhibitors gefitinib, erlotinib and afatinib, the biologic and clinical behavior of crizotinib in ROS1- (and ALK-) positive patients configures the scenario of the disruption of oncogene addiction. Being strictly dependent on ROS1 signaling for survival, growth and progression, its pharmacologic inhibition engenders apoptosis of tumor cells, therefore translating in clinical benefit. Similar to the other oncogene-addicted models in NSCLC, tumor responses are observed in the vast majority of patients, making disease progressions rare events being often explained by diagnostic, pharmacokinetics or molecular caveats.\(^{9,10}\) Albeit ROS1 rearrangement does not seem to be an intrinsic prognostic factor, long-term disease control exerted by crizotinib is mostly relevant, almost doubling the one obtained in EGFR- and ALK-driven tumors undergoing specific treatment. According to the most recent updates, more than 19 months of progression-free survival are achievable with crizotinib in ROS1-positive cancers,\(^{50}\) compared to the 9–12 months observed with the cited oncogenes.\(^{9,51,52}\)

Nevertheless, as the intrinsic nature of advanced NSCLC implies, almost unequivocally, every ROS1-positive patient will undergo disease progression while receiving crizotinib. Already assimilated for EGFR and ALK models, beyond-progression strategies, together with local treatments in terms of oligo-progressive diseases involving or not the central nervous system,\(^{53-55}\) are ostensibly applicable in ROS1-rearranged tumors too. Sooner or later anyway, the molecular escapes that cancer cells find out to slip away crizotinib inhibition need to face novel-generation inhibitors. As seen for the mentioned oncogene-addicted NSCLC, mutations occurring in ROS1 kinase domains preclude crizotinib activity. Since their first report,\(^{56}\) a few of them have been reported to be clinically meaningful.\(^{57}\) Given the homology in their respective kinase domains, ALK and ROS1 share a spectrum of active inhibitors beyond crizotinib (e.g., ceritinib, lorlatinib, entrectinib), developed in order to overcome the resistance to the first-generation molecule. Several clinical
trials, series and single cases reported the activity of these molecules, sometimes with regard to the precise reversion of the molecular event leading to crizotinib resistance. Administration of such drugs after crizotinib exhaustion appears to be of major importance, engendering new responses and positive long-term outcomes, similar to what is observed in ALK-rearranged tumors and in EGFR-mutated ones harboring the resistance mutation T790M, undergoing osimertinib treatment.

Such evidence witnesses the dramatic relevance of the correct detection of \textit{ROS1} rearrangements in patients whose cancers lack other genetic abnormalities. The clinical benefit originating from ROS1 inhibition entails that not even a single ROS1-positive patient should be undiagnosed.

\section*{Molecular diagnostics}

The identification of \textit{ROS1} gene rearrangement is mandatory to treat the patients with \textit{ROS1}-positive NSCLC (Figure 3).

It is generally observed that \textit{ROS1} gene rearrangements more often occur in younger and never/light smokers with adenocarcinoma. In a recent study on 727 lung adenocarcinomas from patients with stage IV disease, ROS1 fusions were independently associated with female sex, younger age at diagnosis and absence of smoking history. Compared with ALK-positive adenocarcinoma, the \textit{ROS1} positive counterpart is more significantly associated with a peripheral location.

Despite all these data, the clinicopathologic features cannot be robustly used to recognize \textit{ROS1}-positive patients. In addition, since this genetic alteration occurs in about 0.5\%–2\% of all NSCLCs, a screening test is necessary in terms of cost-effectiveness.

\textit{ROS1} rearrangement is generally detected using in situ methods, namely FISH and IHC (Figure 4).

Since the clinical trials demonstrating crizotinib efficacy in \textit{ROS1}-positive patients have substantially adopted FISH testing, this method is considered the “gold standard” for determining \textit{ROS1} positivity. Nevertheless, several experiences have evidenced a fair-to-perfect agreement between FISH and IHC tests using the ROS1 rabbit primary antibody D4D6.

At the same time, validated extractive technologies (RT-PCR, NGS, nCounter platform) represent new promising
ROS1 rearrangement detection: present and future

FISH and IHC

FISH using a dual color “break-apart” probes approach is considered the “gold standard” in detecting ROS1 gene rearrangement. If tumor cells are rearranged, the two ends of ROS1 are separated and the portion containing the tyrosine kinase domain is fused with another partner to create an ROS1 fusion gene.

The probes label the 3′ (centromeric) part of the fusion breakpoint with green fluorochrome and the 5′ (telomeric) part with orange fluorochrome. The criteria for ROS1 FISH identification in NSCLC are identical to those proposed for ALK rearrangement, with two main patterns as follows:

1) the break-apart pattern (“conventional” pattern) with one fusion signal and two separated 3′ and 5′ signals and
2) an atypical pattern showing an isolated 3′ signal (usually one fusion signal and one isolated 3′ green signal without the corresponding 5′ signal; Figure 5).19

The green fluorochrome is the part containing the kinase domain of ROS1 gene.

FISH testing for ROS1 is applicable either on biopsy or in cytologic specimens, and the cut-off of rearranged signals to quote ROS1 positivity is based on detection of 15% or more among 50 neoplastic nuclei.66,67

Different probes have been used in literature to detect ROS1 rearrangement (Table 1), and it is important to note that some break-apart FISH assays for ROS1 fusion cannot detect intrachromosomal deletion, as GOPC–ROS1 rearrangement.68

IHC is an effective screening tool to detect ROS1-positive NSCLC, since either sensitivity or specificity may reach over 90% when compared with FISH results (Table 1).18–36 Indeed, screening of NSCLC (all nonsmokers and smokers with nonsquamous NSCLC) by IHC may prevent unnecessary FISH analysis in ROS1-negative cases and, thus, dramatically reduce the costs of testing.69

Data obtained from IHC are based on the use of the rabbit primary monoclonal antibody D4D6 (Cell Signaling Technology, Danvers, MA, USA) applied at different dilutions...
usually from 1:50 to 1:500) with different amplification kits and detection systems in various automated immune
stainers (Table 1).

ROS1 protein is basically absent in normal human lung tissue, but in IHC, its expression may be observed in reactive
alveolar type II pneumocytes and macrophages (Figure 6). Therefore, it is mandatory to insert a positive external con-
trol or a cell block obtained from ROS1-rearranged cell line, preferentially on the same slide of the tumor to be tested or
on a separate slide to be included in the same batch.

Pathologists should also be aware that in bone biopsies
performed in metastatic NSCLC, it is possible to observe
osteoclasts revealing a moderate-to-strong granular cyto-
plasmic staining.

Among the different scoring systems to detect ROS1
targets in metastatic NSCLC, it is possible to observe
expression intensity (negative [score 0], weak [1+], moderate
[2+] or strong staining [3+] was considered. Some authors
adopted an h-score (multiplying the percentage of positive
tumor cells and the intensity of staining from 0 to 3+, ranging
from 0 to 300), with optimal threshold for ROS1 positivity
defined as >100.20 ROS1-rearranged tumors usually show
diffuse immunohistochemical expression (>75% of tumor
cells) with a 2+/3+ staining intensity, basically correspon-
ding to an h-score >100.

ROS1 expression at IHC level typically reveals finely
granular cytoplasmic and/or membranous staining (Figure 7),
although the staining pattern may depend on the different
gene fusion partners.19 The same IHC protocols used in
formalin-fixed, paraffin-embedded (FFPE) tissue specimens
may be employed on cell blocks obtained from fine-needle
aspirates and pleural/pericardial effusions Although ROS1
IHC works on conventional alcohol-fixed cytology, FISH
methodology is superior to IHC on nonbloody cytologic smears
and cytospin slides.65,66 Conversely, IHC should be preferred
in FFPE small biopsy and cell block containing few tumor
cells (Figure 8). Some authors suggested to consider IHC
ROS1 expression only when at least 20 tumor cells are
clearly identifiable in the samples to be analyzed,22 but, in the
absence of a clear-cut IHC cut-off, even a positive result on
a few neoplastic cells should be considered appropriate and
sufficient to quote a tumor as positive/rearranged.

When using clone D4D6 together with highly sensitive
amplification kits, the correlation between IHC and FISH is
good, although presently, IHC seems to be less specific than
FISH when compared with the more robust similar diagnostic

Figure 5 The combination of FISH and protein expression by IHC is the main methodology used in routine practice to detect ROS1 rearrangement. In this, even in the presence of IHC positivity, it is necessary to confirm ROS1 positivity by FISH.

Abbreviations: FISH, fluorescence in situ hybridization; IHC, immunohistochemistry; NGS, next-generation sequencing; NSCLC, non-small cell lung cancer; PCR, polymerase chain reaction.
Table 1 Summary of studies comparing identification of ROS1 rearrangement with FISH and IHC

<table>
<thead>
<tr>
<th>Reference</th>
<th>Number of analyzed cases</th>
<th>Histology</th>
<th>Positive cases</th>
<th>ROS1-positive histology</th>
<th>Clone</th>
<th>Scoring method</th>
<th>Platform</th>
<th>Amplification kit</th>
<th>FISH</th>
<th>Sensitivity, %</th>
<th>Specificity, %</th>
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<td>≥2 +</td>
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<td>BACPAC Resource</td>
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<td>Abbrev</td>
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<td>99</td>
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Abbreviations: NSCLC, non-small cell lung cancer; adeno, adenocarcinoma; LCC, large cell carcinoma; retro, retrospective; pros, prospective; NA, not available.
scenario already established for \textit{ALK} rearrangement. Several other promising anti-ROS1 primary monoclonal and polyclonal antibodies are commercially available, although no overt data have been published so far. Another promising primary rabbit monoclonal antibody (EPMGHR2; Abcam, Cambridge, UK) is now commercially available.

**Extractive technologies**

Extractive assays are based on RT-PCR and massive parallel NGS with kits including several fusion genes (\textit{ROSI}, \textit{ALK}, \textit{RET}, \textit{NTRK}). While the sensitivity and specificity of RT-PCR are quite good, the presence of numerous \textit{ROSI} fusion partners identified (e.g., \textit{CD74}, \textit{FIG}, \textit{SLC34A2}, \textit{TPM3}, \textit{SDC4}, \textit{EZR}, \textit{LRIG3}, \textit{GOPC}, \textit{MSN}, \textit{KDELr2} and \textit{CCDC6}) or still unknown in the absence of a clear predictive value and the availability of good-quality RNA obtained from FFPE small specimens could limit the use of RT-PCR in routine practice.

nCounter platform (NanoString assay) is a multiplex mRNA-based promising test showing high specificity and sensitivity with referenced tests in detecting ROS1. This method may detect several gene fusions by direct profiling using a limited amount of RNA.

Reguart et al recently demonstrated the identification of ALK, RET and ROS1 fusions using nCounter technology in FFPE samples of advanced stage NSCLC. Agreement in detecting ROS1 was 87.2% and 86% with IHC and FISH, respectively.

A combined assay involving a pan-receptor tyrosine kinase immunohistochemical cocktail of antibodies (NTRK1, NTRK2, NTRK3, ROS1 and ALK) followed by an RNA-based anchored multiplex PCR NGS assay has been recently proposed.

The promising results of extractive methodologies applied in clinical practice to determine \textit{ROSI} rearrangement seem to candidate non-in situ methods as the near-standalone assays, thus limiting the role of IHC and FISH.

\textit{ROSI} rearrangement is commonly mutually exclusive with other alterations involving driver oncogenes in lung cancer. According to this view, a recent study on 62 patients with \textit{ROSI}-rearranged NSCLC failed to evidence concurrent \textit{ALK} and \textit{EGFR} gene alterations, while concomitant \textit{KRAS} mutations were detected in two cases (3.2%). No concurrent mutations in \textit{BRAF}, \textit{ERBB2}, \textit{PIK3CA}, \textit{AKT1} or \textit{MAP2K1} were detected. While no concomitant \textit{ALK} and \textit{ROSI} rearrangements were identified, exceedingly rare mutations in \textit{EGFR} and \textit{KRAS} were observed.
Of note, the occurrence of KRAS mutation in ROS1-rearranged lung adenocarcinoma seems to lead to crizotinib resistance in preclinical studies.73

By contrast, a recent work evidenced frequent (36%) concomitant oncogenic driver mutations involving EGFR (6 cases), KRAS (2 cases), PIK3CA and BRAF among 25 lung adenocarcinomas showing ROS1 positivity by IHC.74

Similarly, among 15 cases of ROS1-positive lung adenocarcinomas, 2 patients had TP53 mutations, 1 patient showed an R248L mutation co-occurring with a MAP2K1 missense mutation (K57N) and 1 had an EGFR mutation in exon 21 (P848L). Two out of 17 patients analyzed with NGS or by Sanger sequencing had BRAF mutations and 1 patient showed c-MET mutation (R988C). Overall, 66.7% of patients with ROS1-positive adenocarcinoma analyzed by NGS had further genetic aberrations.73

**ROS1 testing into a practical algorithm**

The availability of specific inhibitors of ROS1 should lead to ROS1 testing into a practical algorithm.

ROS1 rearrangement is generally mutually exclusive with other genetic alterations in NSCLC, but a subset may concurrently harbor EGFR or KRAS mutations.

The role of KRAS mutations in an algorithm of predictive biomarkers in routine practice is still debatable. However, when considered as the most common oncogenic driver occurring in about one-third of adenocarcinomas in the Caucasian population, its value as a negative selecting biomarker precluding useless search of further rare targetable gene alterations seems at least reasonable.

According to the recent proposal by an European Board of Pathologists, a routine clinical practice algorithm based on IHC screening with further confirmation by ROS1 break-apart FISH assay in IHC-positive and doubtful cases seems entirely appropriate (Figure 9).19 In the near future, the advent of highly sensitive extractive methods using a small amount of tumor RNA in a single-tube assay to simultaneously detect several oncogenic fusions (ALK, RET, ROS1 and NTRK1 gene rearrangements; Figure 10) and mutations, coupled to a decrease in the cost of instruments will be implemented even in routine practice, possibly limiting the use of IHC and FISH methods.33,77,78

**Disclosure**

The authors report no conflicts of interest in this work.

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