Lung cancer trial of RET inhibitor selpercatinib achieves durable responses in patients with RET gene fusions

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For patients with non-small cell lung cancers (NSCLC) marked by RET gene fusions, the targeted therapy selpercatinib was well tolerated and achieved durable objective responses, or tumor shrinkage, in the majority of participants in the Phase I/II LIBRETTO-001 trial, according to researchers from The University of Texas MD Anderson Cancer Center.

Among previously untreated patients, the objective response rate (ORR) was 85%, and those receiving at least prior platinum-based chemotherapy had an ORR of 64%. For patients with brain metastases, there was a 91% ORR in the brain.

Evaluating selective RET inhibitor to answer significant unmet need

RET fusions occur when a portion of the chromosome containing the RET gene breaks and rejoins with another piece of chromosome, creating a fusion protein capable of fueling cancer growth. RET alterations occur in roughly 2% of NSCLC, 10-20% of papillary thyroid cancers (PTC) and the vast majority of medullary thyroid cancers (MTC). Up to half of all RET fusion-positive cancers metastasize to the brain.

Several targeted therapies, such as cabozantinib and vandetanib, have ancillary activity against RET alterations, but clinical trials found that NSCLC patients saw only limited benefit from these drugs, explained Subbiah, who is co-principal investigator of the trial.

The study reports findings from 144 patients with advanced NSCLC enrolled on the open-label,
international trial. The primary endpoint was ORR assessed by an independent review committee, and secondary endpoints included safety, intracranial response, duration of response, and progression-free survival (PFS). Results from investigator assessments were not significantly different from that of independent reviewers.

The trial included previously untreated patients (39) and patients having received at least platinum-based chemotherapy (105). Previously treated patients had a median of three prior lines of therapy, including immune checkpoint blockade in more than half. Across both cohorts, trial participants were 57.6% Caucasian, 32.6% Asian, 5.6% Black, 2.8% other and 1.4% unknown. The median age was 61, with women accounting for 58.3% and men 41.7% of participants.

Among previously treated patients, 2% had a complete response, 62% had a partial response and 29% had stable disease. The median duration of response was 17.5 months and 63% of responses were ongoing at a median follow-up of 12 months. Median PFS was 16.5 months.

In previously untreated patients, 85% had a partial response and 10% had stable disease. At six months, 90% of responses were ongoing, and neither median duration of response nor median PFS has been reached at the time of analysis.

On the study, 11 patients had measurable brain metastases. Ten of these patients (91%) saw an objective response in the brain, including three complete responses. The median duration of CNS response was 10.1 months.

The most common adverse events of grade 3 or higher were hypertension (14%), increased aminotransferase (13%), increased aspartate aminotransferase (10%), hyponatremia (6%) and lymphopenia (6%). Four patients discontinued selpercatinib treatment due to treatment-related adverse events.

Strong response also reported in patients with thyroid cancers

In additional cohorts of LIBRETTO-001, selpercatinib also showed activity in RET-altered thyroid cancers, including MTC with RET mutations and papillary/anaplastic thyroid cancers with RET fusions, with a similar safety profile.

Among patients with MTC, there was a 73% ORR in previously untreated patients and 69% ORR in those receiving prior targeted therapies. In patients with previously treated papillary/anaplastic thyroid cancer, the ORR was 79%. These data also were published today in New England Journal of Medicine, with Subbiah and Maria Cabanillas, M.D., professor of Endocrine Neoplasia and Hormonal Disorders, as co-senior authors.

"The data show these patients benefit from this treatment and it is safe compared with multi-kinase inhibitors and chemotherapy," said Subbiah. "The continued implementation of a robust molecular screening strategy in frontline lung and thyroid cancers with the ability to detect RET and other gene fusions will be critical for identifying patients who may benefit from specifically targeted therapies like selpercatinib."

Provided by University of Texas M. D. Anderson Cancer Center