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POSTER

Patient Characteristics in a Cohort of Advanced Non-Small Cell Lung Cancer (NSCLC) Patients Treated in the Community SettingE. Nadler¹, C. Chen², M. Forsyth², J.R. Penrod³, M. Corral³, M. Salvati⁴.¹US Oncology Inc., Texas Oncology PA – Sammons Cancer Ctr/Baylor University Medical Center, Dallas TX, ²US Oncology Inc., Informatics, The Woodlands TX, ³Bristol-Myers Squibb, Health Economics and Outcomes Research, Princeton NJ, ⁴Bristol-Myers Squibb, US Medical Affairs, Princeton NJ, USA

Background: An array of palliative 1st line treatment options for patients (pts) with advanced (metastatic) NSCLC improving survival and reducing disease related symptoms are available. Limited data exists on patient characteristics that drive treatment options. Review of electronic medical records offers an efficient new approach to understanding clinic aspects leading to treatment decisions. The objective of this study was to describe clinical and demographic characteristics that determine whether pts receive 1st line therapy for advanced NSCLC. Pt characteristics and determinants of appropriate treatment outlined in evidence-based treatment guidelines were examined.

Methods: Advanced NSCLC pts receiving active care at US Oncology practices utilizing the iKnowMed electronic medical record system with documentation of 1st line chemotherapy regimen initiated between 1/1/2007–2/28/2010.

Results: 6,862 pts met eligibility criteria, 5,313 pts (77.4%) initiated chemotherapy in 1st line setting and 1,549 pts (22.6%) did not. Mean BMI was higher among treated v untreated pts (26.0±5.6 v 25.4±5.6, p=0.0087). Variation in treatment rates by patient characteristics are given in the table.

| Characteristic | No. of patients (Treated %) N = 6,862 | P values |
|----------------|--|----------|
| Age | | 0.0011 |
| <65 yrs | 2498 (79%) | |
| 65–79 yrs | 3051 (78%) | |
| 80+ yrs | 895 (73%) | |
| Gender | | 0.0233 |
| Male | 3,660 (78%) | |
| Female | 3,202 (76%) | |
| Stage | | <0.0001 |
| Recurrent | 1,380 (82%) | |
| 1st Diagnosis | 4,948 (75%) | |
| Unknown | 534 (86%) | |
| Region | | <0.0001 |
| Midwest | 1,063 (60%) | |
| Northeast | 610 (76%) | |
| South | 3,667 (82%) | |
| West | 1,522 (79%) | |
| Histology | | <0.0001 |
| Squamous | 1,478 (79%) | |
| Non-squamous | 3,463 (80%) | |
| NOS | 1,921 (72%) | |
| Baseline ECOG | | <0.0001 |
| 0 | 546 (71%) | |
| 1 | 3,338 (83%) | |
| 2 | 1,342 (75%) | |
| 3+ | 232 (60%) | |
| Unknown | 1,404 (71%) | |
| Payer Status | | 0.1246 |
| Medicare | 3,643 (77%) | |
| Medicaid | 123 (84%) | |
| Private | 1,729 (79%) | |
| Other | 1,350 (77%) | |
| Unknown | 17 (71%) | |

Conclusions: These real world data from a national oncology network describe patient characteristics used by physicians to initiate treatment for advanced NSCLC. 77.4% of pts were given 1st line systemic treatment with factors such as age, gender, BMI, stage, region, histology and performance score as strong influencers in treatment decision. Payer status did not significantly impact decision to treat.

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POSTER

Pattern of Chemotherapy for Advanced Non-small Cell Lung Cancer in Relation to EGFR Mutation Status – Clinical Experience in a Comprehensive Cancer Center in KoreaK. Park¹, J. Lee¹, J. Sun¹, J. Cho², E. Guallar³, B. Parasuraman⁴, G. Lee⁵, J. Han⁶, Y. Choi⁶, Y.M. Shim⁵. ¹Samsung Medical Center, Division of Hematology-Oncology Department of Medicine, Seoul, ²Samsung Medical Center, Cancer Education Center, Seoul, Korea; ³Johns Hopkins Bloomberg School of Public Health, Department of Epidemiology, Baltimore, ⁴Astra Zeneca, Health Economics and Outcome Research, Willmington, USA; ⁵Samsung Medical Center, Department of Thoracic and Cardiovascular Surgery, Seoul, ⁶Samsung Medical Center, Department of Pathology, Seoul, Korea

Background: Epidermal growth factor receptor (EGFR) mutation is an established predictive marker for EGFR tyrosine kinase inhibitor (TKI) treatment in advanced non-small cell lung cancer (NSCLC). A growing body of evidence supports EGFR mutation testing to guide initial treatment decisions for patients with advanced NSCLC. This study describes the pattern of practice for chemotherapy for NSCLC in a clinical setting in a comprehensive cancer center in Korea.

Method: Retrospective cohort study of 1,527 patients diagnosed with stage IIIB/IV NSCLC at Samsung Medical Center (SMC) in Seoul, Korea, from January 2007 through July 2010.

Results: Data were analyzed from 1,243 patients not involved in clinical trials. The proportions of patients with adenocarcinoma and squamous cell carcinoma were 68.2%, and 18.8%, respectively. The number (percent) of patients who received at least one, two, or three lines of systemic treatment were 1,140 (91.7%), 775 (62.3%) and 487 (39.2%), respectively. EGFR mutation testing was successfully performed in 446 (35.9%) patients, of whom 163 (36.6%) were EGFR positive. The treatments received by EGFR mutation status are shown in the Table. In the pretreated setting, EGFR-TKIs (e.g. erlotinib or gefitinib) were more frequently administered to EGFR M(+) patients compared to EGFR M(–) patients (73.6% vs 44.9%; p<0.001).

| | EGFR mutation (+) | EGFR mutation (–) | EGFR mutation unknown |
|--------------------------------------|-------------------|-------------------|-----------------------|
| 1st line treatment | N = 161 | N = 270 | N = 709 |
| Gemcitabine/platinum | 88 (54.7%) | 159 (58.9%) | 446 (62.9%) |
| Pemetrexed/platinum | 33 (20.5%) | 49 (18.2%) | 34 (4.8%) |
| Gefitinib | 16 (9.9%) | 19 (7.0%) | 55 (7.8%) |
| Other | 24 (14.9%) | 43 (15.9%) | 174 (24.5%) |
| 2nd line treatment | N = 137 | N = 184 | N = 454 |
| Gefitinib/erlotinib | 93 (67.9%) | 79 (42.9%) | 176 (38.8%) |
| Pemetrexed | 26 (19.0%) | 56 (30.4%) | 133 (29.3%) |
| Gemcitabine/platinum | 5 (3.6%) | 14 (7.6%) | 37 (8.2%) |
| Other | 13 (9.5%) | 35 (19.0%) | 108 (23.8%) |
| 3rd line treatment | N = 75 | N = 119 | N = 293 |
| Pemetrexed | 32 (42.7%) | 34 (28.6%) | 78 (26.6%) |
| Gefitinib/erlotinib | 29 (38.7%) | 49 (41.2%) | 148 (50.5%) |
| Other | 14 (18.7%) | 36 (30.2%) | 67 (22.9%) |

*Percentages may not add to 100% due to rounding errors.

Conclusion: During the study period, EGFR testing and use of EGFR-TKIs were not reimbursed in Korea. Less than 40% of subjects were successfully tested, but the results of EGFR mutation testing influenced the selection of chemotherapy in pretreated patients. Accumulating evidence on the efficacy of EGFR-TKIs and changes in reimbursement policies may change clinical management of advanced NSCLC patients in the future.