Systematic Review

Comparative Efficacy and Safety of Anti-PD-1/PD-L1 Immune Checkpoint Inhibitors for Refractory or Relapsed Advanced Non-Small-Cell Lung Cancer—A Systematic Review and Network Meta-Analysis

Koichi Ando 1,2,*, Ryo Manabe 1, Yasunari Kishino 1, Sojiro Kusumoto 1, Toshimitsu Yamaoka 3, Akihiko Tanaka 1, Tohru Ohmori 1, Tsukasa Ohnishi 1 and Hironori Sagara 1

1 Division of Respiratory Medicine and Allergology, Department of Medicine, Showa University School of Medicine, 1-5-8 Hatanodai, Shinagawa-ku, Tokyo 142-8666, Japan; r.manabe@med.showa-u.ac.jp (R.M.); ookiyookiy@med.showa-u.ac.jp (Y.K.); k-sojiro@med.showa-u.ac.jp (S.K.); tanakaa@med.showa-u.ac.jp (A.T.); tohmorit@med.showa-u.ac.jp (T.O.); tohnishi@med.showa-u.ac.jp (T.O.); sagarah@med.showa-u.ac.jp (H.S.)
2 Division of Internal Medicine, Showa University Dental Hospital Medical Clinic, Showa University Senzoku Campus, 2-1-1 Kita-senzoku, Ohta-ku, Tokyo 145-8515, Japan
3 Advanced Cancer Translational Research Institute (Formerly, Institute of Molecular Oncology), Showa University, 1-5-8 Hatanodai, Shinagawa-ku, Tokyo 142-8555, Japan; yamaoka.t@med.showa-u.ac.jp
* Correspondence: koichi-a@med.showa-u.ac.jp; Tel.: +81-3-3784-8532; Fax: +81-3-3784-8742

Simple Summary: Improving treatment strategies for refractory or relapsed advanced non-small-cell lung cancer (NSCLC) remains a challenge. Efficacy and safety of the immune checkpoint inhibitors (ICIs), nivolumab (Niv) plus atezolizumab (Atz), were compared with those of ramucirumab (Ram) plus docetaxel (Doc), and the efficacy and safety of these two ICIs were compared with each other, using patient groups without programmed cell death ligand-1 (PD-L1) constraint. Additionally, efficacy and safety of, Niv, Atz, and pembrolizumab (Pem) were compared using a PD-L1 positive (≥1%) subgroup with refractory or relapsed advanced NSCLC. Niv or Atz was found to be more effective and safer than Ram plus Doc in groups without PD-L1 constraint. In the PD-L1 positive subgroup, Pem (10 mg/kg) showed the highest efficacy for ensuring overall survival, followed by Niv, Pem (2 mg/kg), Atz, and Doc. These results may help clinicians select and evaluate treatment options for refractory or relapsed advanced NSCLC.

Abstract: The efficacy and safety of immune checkpoint inhibitors (ICIs) in refractory or relapsed advanced non-small-cell lung cancer (NSCLC) have not yet been compared with those of ramucirumab (Ram) plus docetaxel (Doc). Furthermore, comprehensive comparisons between ICIs have not been conducted to date. In the current study, a Bayesian network meta-analysis of related phase III clinical trials was performed to compare the efficacy and safety of Ram+Doc, Niv, Atz, and Doc treatments in patient groups lacking the PD-L1 constraint. Surface under the cumulative ranking area (SUCRA) revealed that the overall survival (OS) of patients treated with Niv was the highest, followed by Atz, Ram+Doc, and Doc. Regarding grades 3–5 treatment-related adverse events (G3–5AEs), the use of Niv was ranked the safest, followed by Atz, Doc, and Ram+Doc. Significant differences in OS were observed between Niv and Ram+Doc, while significant differences in G3–5AEs were observed between Ram+Doc and Niv or Atz. In the PD-L1 positive (≥1%) patient subgroup, Pem (10 mg/kg) ranked the highest in efficacy for OS, followed by Niv, Pem (2 mg/kg), Atz, and Doc. These findings may expectedly provide oncologists with useful insights into therapeutic selection for refractory or relapsed advanced NSCLC.

Keywords: non-small-cell lung cancer; immune checkpoint inhibitors; overall survival; progression-free survival; network meta-analysis; systematic review
1. Introduction

Advances in molecular and immunologic chemotherapies [1–3] have resulted in the development of novel strategies for lung cancer treatment [1–4]. Despite these advances, lung cancer remains a major source of cancer-related death, accounting for 13% of all cancer mortalities, which is attributed to the poor 5-year survival rate (18%) associated with this disease [5]. When detected at an advanced stage, as is common in the more prevalent (84%) non-small-cell lung cancer (NSCLC) cases, the 5-year survival rate is only 5% [2,3,5]. Furthermore, relapse is a common occurrence in NSCLC, even following a positive response to initial treatment. Although these findings are based on data obtained from patients who had not been treated with immunotherapy, the prognosis for advanced NSCLC remains mostly poor. Therefore, improved and modified treatment strategies are warranted for refractory or relapsed NSCLC [1–3,5–8].

Recently, the efficacy of a combination of Ram and Doc (Ram+Doc) was evaluated as a second-line treatment for advanced NSCLC [9]. Ram+Doc treatment significantly improved overall survival (OS) as well as progression-free survival (PFS) in comparison with Doc monotherapy. Consequently, the Ram+Doc combination remains the treatment of choice for driver mutation-negative refractory or relapsed advanced NSCLC, or cancer of unknown status [2,3,9,10]. Nintedanib plus Doc, a treatment that has been approved based on the results of previous phase III trials, may present a new treatment option for adenocarcinoma [11].

Immunotherapy, including the use of immune checkpoint inhibitors (ICIs), has recently been approved by the FDA for lung cancer treatment and is widely used for this purpose [1,4,12,13]. Treatment with Niv, Atz, and Pem, in particular, have significantly improved the outcome of NSCLC patients [1,3–5,12,13]. However, to date, only a few studies have compared ICIs with Ram+Doc treatments. Moreover, studies comparing the efficacy and safety profiles of ICIs with each other in refractory or recurrent advanced NSCLC are scant. Although ICIs have been approved for the treatment of refractory or relapsed NSCLC [1,6], there is limited evidence suggesting that ICIs are superior to conventional therapies, such as Ram+Doc. Furthermore, currently, there is insufficient evidence indicating which ICIs are the most effective as well as safe for use in refractory or recurrent advanced NSCLC. Thus, a comparative analysis of ICIs and Ram+Doc treatments leading to a ranking in the order of usefulness or efficacy against this disease may be warranted.

In general, randomized controlled trials (RCTs) are considered to be the most appropriate for comparing the efficacy and safety of treatment interventions. However, RCTs are beset by limitations such as time constraints as well as logistical and financial considerations. Therefore, we adopted another statistical method, network meta-analysis (NMA), as an alternative [14–16]. Our study design permits the efficacy and safety of any treatment pair to be compared and ranked more rapidly than it would be possible via RCTs, even in the absence of previous studies of directly comparative RCTs [14–22]. Thus, using this systematic review and meta-analysis (registration: UMIN-CTR no. UMIN000041086), we compared and ranked the efficacy and safety of ICIs, Doc and Ram+Doc in treating refractory or relapsed patients with NSCLC using statistical Bayesian NMA. Our results may expectedly provide insights into the most efficacious and safe treatments for NSCLC.

2. Results

2.1. Systematic Review

Of the 1526 articles identified via a systematic literature review that met the search criteria (121 from PubMed [23], 276 from Embase [24], 345 from the Cochrane Central Register of Controlled Trials (CENTRAL) [25], and 784 from SCOPUS [26]), 1084 remained following the removal of duplicates (Figure 1). The application of the collation of Patients, Intervention, Comparison, Outcome and Study (PICOS) criteria (Table 1) suggested that four studies were in order: one comparing Ram+Doc with Doc [9]; and the others comparing Doc with Niv [27,28] or Atz [29]; (Table 2).
The common comparative group in these four studies was the Doc group [9,27–29]. The data obtained from these studies were sufficient to perform an NMA for OS, including a subgroup analysis based on histology. However, these data were not sufficient to conduct a subgroup analysis of PFS. Therefore, PFS was analyzed only in the overall patient group.
In all analyses, the preferred model convergence was confirmed using the Brooks–Gelman–Rubin method [30,31].

Although Pem is indicated for the treatment of NSCLC, immunological regimens comprising Pem were not included in the present NMA owing to the heterogeneity of target patients. Phase III trials, in which refractory or relapsed NSCLC was treated with Pem [32], included a patient group with limited PD-L1 expression, which was considered unsuitable for inclusion in this NMA. By contrast, CheckMate057, CheckMate017, and OAK, like REVEL, did not place a limit on PD-L1 positivity in their patient inclusion criteria. Therefore, including these four RCTs in the analysis allowed each treatment group to be compared in a uniform PD-L1 status environment. The network map of the NMA conducted in the present study is shown (Figure 2).

Figure 2. Network map. Solid lines represent randomized controlled trials (RCTs) while relative thickness represents the number of included studies. The dashed line reveals the absence of RCTs, suggesting that an indirect treatment comparison could be attempted. Circle size reflects the proportion of patients included in each treatment group. Ram+Doc (ramucirumab plus docetaxel); Niv (nivolumab); Atz (atezolizumab); n, number of patients.
| Study Names | Treatment Arms | N  | Age, Year Median (Range) | Females No. (%) | ECOG PS No. (%) | Histological Type No. (%) |
|-------------|----------------|----|-------------------------|-----------------|----------------|--------------------------|
| REVEL [9]   | Ramucirumab (10 mg/kg) plus docetaxel (75 mg/m²) on day 1 of 21-day cycle | 628 | 62 (21–85) | 209 (33) | PS0: 207 (33) | Non-squamous: 465 (74) |
|             | Placebo plus docetaxel (75 mg/m²) on day 1 of 21-day cycle | 625 | 61 (25–86) | 210 (34) | PS0: 199 (32) | Squamous: 171 (27) |
|             | | Total: 1253 | | | | Unknown: 6 (1) |
| CheckMate057 [28] | Nivolumab (3 mg/kg e2w) | 292 | 61 (37–84) | 141 (48) | PS0: 84 (29) | Non-squamous: 292 (100) |
|             | Docetaxel (75 mg/m² e3w) | 290 | 64 (21–85) | 122 (42) | PS0: 95 (33) | Squamous: 0 (0) |
|             | | Total: 582 | | | | NR: 0 |
| CheckMate017 [27] | Nivolumab (3 mg/kg e2w) | 135 | 62 (39–85) | 24 (18) | PS0: 27 (20) | Non-squamous: 0 (0) |
|             | Docetaxel (75 mg/m² e3w) | 137 | 64 (42–84) | 40 (29) | PS0: 37 (27) | Squamous: 135 (100) |
|             | | Total: 272 | | | | NR: 0 |
| OAK [29]    | Atezolizumab (1200 mg e3w) | 425 | 63.0 (33.0–82.0) | 164 (39) | PS0: 155 (36) | Non-squamous: 313 (74) |
|             | Docetaxel (75 mg/m² e3w) | 425 | 64.0 (34.0–85.0) | 166 (39) | PS0: 160 (38) | Squamous: 110 (26) |
|             | | Total: 850 | | | | PS1: 270 (64) |

N, sample size; ECOG, Eastern Cooperative Oncology Group; PS, performance status; e3w, every 3 weeks; e2w, every 2 weeks; NR, not reported.
2.2. Primary Efficacy Endpoint: OS

The OS of patients treated with Atz, Niv, and Ram+Doc was significantly higher than that of patients treated with Doc (hazard ratio [HR] [95% credible intervals [CrIs]]: 0.732 [0.615–0.864], 0.683 [0.575–0.806], and 0.862 [0.752–0.983], respectively). The OS of patients in the Niv group was significantly higher than that of patients in the Ram+Doc group (HR: 0.796 [95% CrI: 0.639–0.982]); however, there were no significant differences in the OS of patients receiving Atz vs. Ram+Doc (HR: 0.854 [95% CrI: 0.683–1.054]) or those receiving Atz vs. Niv (HR: 1.080 [95% CrI: 0.843–1.360]) (Figure 3).

2.3. Subgroup Analysis of the OS of Patients with Non-Squamous NSCLC

The OS of patients in the Atz, Niv and Ram+Doc groups was significantly higher than that of patients in the Doc group (HRs [95% CrIs]: 0.733 [0.598–0.888], 0.734 [0.595–0.898], and 0.833 [0.708–0.972], respectively). However, no statistically significant differences were noted for the Atz vs. Niv groups, Atz vs. Ram+Doc groups, and Niv vs. Ram+Doc groups (HRs [95% CrIs]: 1.010 [0.750–1.327], 0.887 [0.682–1.133], and 0.888 [0.679–1.142], respectively) (Figure 4).

2.4. Subgroup Analysis of OS of Patients with Squamous NSCLC

The OS of patients in the Atz and Niv groups was significantly higher than that of patients in the Doc group (HRs [95% CrIs]: 0.738 [0.540–0.982] and 0.597 [0.441–0.793], respectively). The OS of patients in the Niv group was improved compared with that of patients in the Ram+Doc group (HR: 0.682 [95% CrI: 0.457–0.981]). By contrast, no statistically significant differences were noted for Atz vs. Niv, Atz vs. Ram+Doc or Ram+Doc vs. Doc (HRs [95% CrIs]: 1.264 [0.812–1.874], 0.843 [0.561–1.216], and 0.890 [0.691–1.126], respectively) (Figure 5).
Comparing the OS of patients with refractory or relapsed advanced non-squamous non-small-cell lung cancer in the Ram+Doc, Doc, Niv and Atz groups. The patient inclusion criteria for all RCTs included in this analysis did not place any limitations on PD-L1 status. Comparisons are represented as treatment A versus treatment B. Data are presented as hazard ratios (HRs) and 95% credible intervals (CrIs). Ram+Doc, (ramucirumab plus docetaxel); Doc (docetaxel); Niv (nivolumab); Atz (atezolizumab); HR (hazard ratio); CrI (credible interval).

Comparative efficacy of the four therapeutic regimens, Ram+Doc, Doc, Niv and Atz, pertaining to the OS of patients with refractory or relapsed advanced squamous non-small-cell lung cancer. The patient inclusion criteria for all RCTs in this analysis did not place any limitations on PD-L1 status. Comparisons are represented as treatment A versus treatment B. Data are presented as HRs (95% CrIs). Ram+Doc, (ramucirumab plus docetaxel); Doc (docetaxel); Niv (nivolumab); Atz (atezolizumab); HR (hazard ratio); CrI (credible interval).

2.5. Secondary Efficacy Endpoint: PFS

The PFS of patients in the Niv and Ram+Doc groups was significantly higher than that of patients in the Doc group (HRs [95% CrIs]: 0.817 [0.700–0.948] and 0.761 [0.676–0.855], respectively), whereas no significant differences were observed in PFS for Atz vs. Doc (HR: 0.952 [95% CrI: 0.819–1.099]). Atz was significantly inferior to Ram+Doc in terms of PFS.
Comparative safety of the four therapeutic regimens, Ram+Doc, Doc, Niv, and Atz, pertaining to grade 3-5 treatment-related adverse events in patients with refractory or relapsed advanced non-small-cell lung cancer. The patient inclusion criteria for all RCTs in this analysis did not place any limitations on PD-L1 status. Comparisons are expressed as treatment A versus treatment B. Data are presented as RRs and 95% CrIs. Ram+Doc (ramucirumab plus docetaxel); Doc (docetaxel); Niv (nivolumab); Atz (atezolizumab); HR (hazard ratio); CrI (credible interval).

2.6. Primary Safety Endpoint: Grade 3-5 Treatment-Related Adverse Events (G3–5AEs)

G3–5AEs were significantly more common in the Ram+Doc group than in the Doc group (risk ratio [RR]: 1.101; 95% CrI: 1.032–1.172). The incidence of G3–5AEs in the Atz and Niv groups was significantly lower than that of the Doc group (RRs [95% CrIs]: 0.342 [0.274–0.421] and 0.175 [0.127–0.235], respectively). The incidence of G3–5AEs in the Atz and Niv groups was significantly lower than that of the Ram+Doc group (RRs [95% CrIs]: 0.311 [0.247–0.386] and 0.159 [0.115–0.215], respectively). The incidence of G3–5AEs in the Atz group was significantly higher than that of the Niv group (RR: 2.005 [95% CrI: 1.350–2.862]) (Figure 7).

Figure 6. Comparing the progression-free survival (PFS) of patients with refractory or relapsed advanced non-small-cell lung cancer in the Ram+Doc, Doc, Niv, and Atz groups. The patient inclusion criteria for all RCTs in this analysis did not place any limitations on PD-L1 status. Comparisons are expressed as treatment A versus treatment B. Data are presented as HRs (95% CrIs). Ram+Doc (ramucirumab plus docetaxel); Doc (docetaxel); Niv (nivolumab); Atz (atezolizumab); RR (risk ratio); CrI (credible interval).

Figure 7. Comparative safety of the four therapeutic regimens, Ram+Doc, Doc, Niv, and Atz, pertaining to grade 3-5 treatment-related adverse events in patients with refractory or relapsed advanced non-small-cell lung cancer. The patient inclusion criteria for all RCTs in this analysis did not place any limitations on PD-L1 status. Comparisons are expressed as treatment A versus treatment B. Data are presented as RRs and 95% CrIs. Ram+Doc (ramucirumab plus docetaxel); Niv (nivolumab); Atz (atezolizumab); Doc (docetaxel); RR (risk ratio); CrI (credible interval).
2.7. Ranking Assessment

The efficacy and safety of Ram+Doc, Doc, Niv, and Atz treatments were ranked according to the surface under the cumulative ranking area (SUCRA) curve. Higher SUCRA values indicate better outcomes. SUCRA values were estimated for efficacy and safety outcome indicators, OS, PFS, and G3–5AEs, while a subgroup analysis for OS was conducted based on histological data. SUCRA analyses indicated that OS was highest for Niv (SUCRA, 90.0%) followed by Atz (SUCRA, 73.8%), Ram+Doc (SUCRA, 35.7%), and Doc (SUCRA, 0.5%); (Figure 8A).

SUCRA values for the OS of non-squamous NSCLC patients indicated that Atz treatment (78.1%) was the best, followed by Niv (77.7%), Ram+Doc (43.8%), and Doc (0.4%); (Figure 8B). The SUCRA values for OS of squamous NSCLC patients indicated that Niv treatment (94.0%) was the best, followed by Atz (65.9%), Ram+Doc (34.2%), and Doc (6.0%); (Figure 8C). The SUCRAs for PFS indicated that Ram+Doc treatment (91.6%) was the best, followed by Niv (72.0%), Atz (28.0%), and Doc (8.3%); (Figure S1). The SUCRAs for G3–5AEs indicated that Niv treatment (100.0%) was the best, followed by Atz (66.7%), Doc (33.3%), and Ram+Doc (0.1%) treatments (Figure 8A–C and Figure S1).

Figure 8. Scatter diagrams of SUCRAs for OS and safety in terms of G3–5AEs among Ram+Doc, Niv, Atz and Doc treatments in refractory or relapsed advanced NSCLC patients. (A) OS in the overall population was highest following Niv treatment, followed by Atz, Ram+Doc and Doc; (B) In a subgroup analysis of OS in patients with non-squamous NSCLC, Atz treatment was associated with the highest OS, followed by Niv, Ram+Doc, and Doc; (C) In a subgroup analysis of OS in squamous NSCLC patients, Niv treatment was associated with the highest OS, followed by Atz, Ram+Doc, and Doc.
Figure 8. Scatter diagrams of SUCRAs for OS and safety in terms of G3–5AEs among Ram+Doc, Niv, Atz and Doc treatments in refractory or relapsed advanced NSCLC patients (A) OS in the overall population was highest following Niv treatment, followed by Atz, Ram+Doc and Doc; (B) In a subgroup analysis of OS in patients with non-squamous NSCLC, Atz treatment was associated with the highest OS, followed by Niv, Ram+Doc, and Doc; (C) In a subgroup analysis of OS in squamous NSCLC patients, Niv treatment was associated with the highest OS, followed by Atz, Ram+Doc, and Doc. Meanwhile, Niv treatment ranked the highest in terms of safety, followed by Atz, Doc, and Ram+Doc (A–C). SUCRA (surface under the cumulative ranking area curve); G3–5AEs (grade 3–5 treatment-related adverse events); Niv (nivolumab); Atz (atezolizumab); Ram+Doc (ramucirumab plus docetaxel); Doc (docetaxel).

2.8. Sensitivity Analysis

The OAK study included a group of patients who had received one previous systemic treatment and another group of patients who had received two systemic treatments. By contrast, other studies, such as CheckMate017, CheckMate057, and REVEL, included only patients who had received one previous systemic treatment.

To evaluate the effect of including or excluding the patient group that received two prior systemic treatments in the OAK study, we performed a sensitivity analysis for the primary endpoint (OS). The sensitivity analysis of almost any paired-treatment comparison designed to determine statistical significance presented a similar result. Furthermore, there was negligible change in the effect size and SUCRA values, and the ranking of the primary efficacy outcome was similar for each treatment. The results of the sensitivity analyses are presented (Tables S1 and S2). Based on these results, we considered that the inclusion or exclusion of patients with two prior systemic treatments in OAK did not affect our conclusions.

2.9. Bias Assessment

The quality of the studies included in the analysis was assessed using the Cochrane-recommended risk of bias tool 2 (ROB2) [33]. Although three studies presented some concerns regarding the bias attributed to deviation from intended intervention, due to their open-label design, the remaining study was considered to have a low risk of bias. Other parameters evaluated, i.e., bias arising from the randomization process, bias due to missing outcome data, bias in measurement of the outcome and bias in the selection of the reported results, were judged to be low risk in all four studies; thus, the quality of the included studies was considered to be good (Figure S2).
2.10. Comparison between the Effect of ICIs on Refractory or Relapse PD-L1-Positive (≥1%) Advanced NSCLC

2.10.1. Network Meta-Analysis of Six RCTs from REVEL, CheckMate057, CheckMate017, OAK, LUME-lung 1 and KEYNOTE-010, for Predefined Efficacy and Safety Outcomes (OS, PFS, and G3-5AEs)

Prior to comparing ICIs in the PD-L1-positive (≥1%) group, we performed a network meta-analysis of predefined efficacy and safety outcomes by including the overall participants of six studies, REVEL, CheckMate057, CheckMate017, OAK, LUME-lung 1 and KEYNOTE-010, and based on its results, conducted a subgroup analysis of refractory or relapse PD-L1-positive (≥1%) advanced NSCLC. Although KEYNOTE-010 and LUME-Lung 1 did not satisfy the predetermined criteria for inclusion in the NMA for a patient group lacking PD-L1 constraint, we surmised that the additional inclusion of Pem and Nin+Doc into the treatment arms of NMA may lead to further interesting findings. Based on this contention, we attempted to assess predefined efficacy and safety outcomes using these two trials, in addition to the four already included, for a total of six trials (REVEL, CheckMate057, CheckMate017, OAK, LUME-Lung 1, KEYNOTE-010).

The key inclusion criteria, main characteristics and results of risk of bias assessment of LUME-Lung 1, a phase III study of Nin+Doc, and KEYNOTE-010, a Phase II/III study of Pem, are shown (Tables S3 and S4, and Figure S3, respectively).

A comparative analysis which compared the effect of the seven treatments, Doc, Ram+Doc, Niv, Atz, Pem2, Pem10, and Nin+Doc on the OS of the overall participants of six RCTs, REVEL, CheckMate057, CheckMate017, OAK, KEYNOTE-010, and LUME-Lung 1, was performed. The network map is shown in Figure 9, and the results are shown in Table S5.

Furthermore, five RCTs, REVEL, CheckMate057, OAK, KEYNOTE-010 and LUME-Lung 1, were also included in the analysis that compared the effects of the six treatments, Doc, Ram+Doc, Niv, Atz, pem 2 mg/kg or 10 mg/kg (Pem pooled), and Nin+Doc on the OS of the non-squamous subgroup (Table S6). In addition, four RCTs, REVEL, CheckMate017, OAK and KEYNOTE-010, were included in the analysis that compared the efficacy of the five treatments, Doc, Ram+Doc, Niv, Atz and Pem (pooled), on the OS of the squamous subgroup, (Table S7). A further analysis compared the effects of the seven treatments, Doc, Ram+Doc, Niv, Atz, Pem2, Pem10, and Nin+Doc, on the PFS for overall participants of six RCTs, REVEL, CheckMate057, CheckMate017, OAK, KEYNOTE-010, and LUME-Lung 1. (Table S8). In addition, another analysis compared the effects of seven treatments, Doc, Ram+Doc, Niv, Atz, Pem2, Pem10 and Nin+Doc, on safety, as indicated by G3-5AEs, of the overall participants of six RCTs, REVEL, CheckMate057, CheckMate017, OAK, KEYNOTE-010 and LUME-Lung 1 (Table S9).

The results of ranking assessment as indicated by SUCRA value are presented (Table S10).

Figure 9. Network map of the analysis comparing the efficacy of the seven treatments, Doc, Ram+Doc, Niv, Atz, Pem2, Pem10, and Nin+Doc on the overall survival of participants of six RCTs, REVEL, CheckMate057, CheckMate017, OAK, KEYNOTE-010, and LUME-Lung 1. Solid lines represent randomized controlled trials (RCTs), and relative thickness represents the number of included studies. The dashed line reveals the absence of RCTs, suggesting that an indirect treatment comparison could be attempted. Circle size reflects the proportion of patients included in each treatment group. Ram+Doc, ramucirumab plus docetaxel; Niv, nivolumab; Atz, atezolizumab; Pem2, pembrolizumab 2 mg/kg; Pem10, pembrolizumab 10 mg/kg; n, number of patients.
Furthermore, five RCTs, REVEL, CheckMate057, OAK, KEYNOTE-010 and LUME-Lung 1, were also included in the analysis that compared the effects of the six treatments, Doc, Ram+Doc, Niv, Atz, pem 2 mg/kg or 10 mg/kg (Pem pooled), and Nin+Doc on the OS of the non-squamous subgroup (Table S6). In addition, four RCTs, REVEL, CheckMate017, OAK and KEYNOTE-010, were included in the analysis that compared the efficacy of the five treatments, Doc, Ram+Doc, Niv, Atz and Pem (pooled), on the OS of the squamous subgroup, (Table S7). A further analysis compared the effects of the seven treatments, Doc, Ram+Doc, Niv, Atz, Pem2, Pem10 and Nin+Doc, on the PFS for overall participants of six RCTs, REVEL, CheckMate057, CheckMate017, OAK, KEYNOTE-010, and LUME-Lung 1. (Table S8). In addition, another analysis compared the effects of seven treatments, Doc, Ram+Doc, Niv, Atz, Pem2, Pem10 and Nin+Doc, on safety, as indicated by G3-5AEs, of the overall participants of six RCTs, REVEL, CheckMate057, CheckMate017, OAK, KEYNOTE-010 and LUME-Lung 1 (Table S9).

The results of ranking assessment as indicated by SUCRA value are presented (Table S10).

2.10.2. Subgroup Analysis for Refractory or Relapse PD-L1-Positive (≥1%) Advanced NSCLC

We performed a subgroup analysis for the PD-L1-positive (≥1%) patient population and compared the efficacy for OS between ICIs. Four RCTs, CheckMate057, CheckMate017, OAK, and KEYNOTE-010, were included in this subgroup analysis which compared the efficacy of five treatments, Doc, Niv, Atz, Pem2 and Pem10. The network map of the NMA conducted as the subgroup analysis for refractory or relapse PD-L1-positive (≥1%) advanced NSCLC is shown in Figure 10, and the results are shown in Figure 11.

**Figure 10.** Network map of subgroup analysis for refractory or relapse PD-L1-positive (≥1%) advanced NSCLC. Solid lines represent randomized controlled trials (RCTs), and relative thickness represents the number of included studies. The dashed line reveals the absence of RCTs, suggesting that an indirect treatment comparison could be attempted. Circle size reflects the proportion of patients included in each treatment group. Ram+Doc, ramucirumab plus docetaxel; Niv, nivolumab; Atz, atezolizumab; Pem2, pembrolizumab 2 mg/kg; Pem10, pembrolizumab 10 mg/kg; n, number of patients.
3. Discussion

The current NMA compared the efficacy and safety of ICIs with Ram+Doc for refractory or relapsed advanced NSCLC in patient groups lacking the PD-L1 status constraint. The primary efficacy endpoint, OS, was significantly improved following treatment with Niv compared with Ram+Doc treatment. However, no significant differences were observed between the Atz and Ram+Doc groups. The incidence of G3–5AEs was lower in both Niv and Atz groups compared with that of the Ram+Doc group. The Niv group displayed the highest SUCRA for OS, followed by Atz, Ram+Doc and Doc groups. The Niv group showed the highest SUCRA for G3–5AEs (lowest incidence of G3–5AEs), followed by Atz, Doc and Ram+Doc groups. In subgroup analysis of the PD-L1-positive (≥1%) patient population, ICI-to-ICI comparisons indicated that Pem10 displayed the highest efficacy for OS, followed by Niv, Pem2 and Atz, although there were no significant differences between these ICIs.

Previous studies have compared the efficacy and safety of Ram+Doc and Doc [9], Niv and Doc [27,28], Atz and Doc [29], and Pem and Doc in cases limited to PD-L1 (≥1%) positive patients [32]. In patients with refractory or relapsed advanced NSCLC, Ram+Doc, Niv, Atz, and Pem significantly improved OS compared with Doc [9,27–29,32]. However, no studies have compared the efficacy and safety of ICIs with those of Ram+Doc. Moreover, there are no reports comparing the efficacy and safety of ICIs. In this systematic review and meta-analysis, we adopted a unique approach which simultaneously compared and ranked the efficacy and safety of four treatments, Ram+Doc, Doc, Niv and Atz, which are indicated for relapsed or refractory NSCLC in patients lacking the PD-L1 constraint. We also compared the inter-ICI efficacy of Niv, Atz and Pem for affecting OS in the PD-L1-positive (≥1%) patient subgroup. The results of the present study suggest that Niv and Atz displayed superior efficacy and safety profiles than did Ram+Doc for...
relapsed or refractory advanced NSCLC in patients lacking the PD-L1 constraint. This is an important clinical finding that will enhance the ability of oncologists to treat patients more effectively. Our results showed that ICIs displayed favorable efficacy and safety profiles compared with those of Ram+Doc for relapsed or refractory advanced NSCLC. Although Pem could not be included in the treatment comparison analysis for patient groups lacking the PD-L1 constraint, due to non-compliance with our predetermined inclusion criteria, a comparison between Niv and Atz indicated that Niv showed relatively better efficacy and safety profiles than did Atz. Particularly, with respect to the safety profile, Atz had a significantly higher frequency of G3–5AEs than did Niv. In the subgroup analysis of the PD-L1-positive (≥1%) patient population, Pem10 had the best efficacy profile, followed by Niv, Pem2, Atz, and finally Doc, although the results did not show a significant difference between these. Furthermore, PD-L1-positive (≥1%) cases were analyzed by histological type, allowing comparison between Pem (pooled) and Niv. The results showed that no significant differences existed between the effects of Pem (pooled) and Niv on OS in non-squamous and squamous patients, respectively.

The molecular basis underlying these findings requires clarification: Cell death caused by systemic anticancer therapy, such as chemotherapy, elicits an inflammatory response that induces the accumulation of activated T cells. Activated T cells are suppressed by cancer cells via PD-L1, which is eliminated by treatment with Niv and Atz, during secondary and subsequent therapies, thereby restoring the anticancer activity of T cells as well as apoptosis of cancer cells [1,4,12,34]; (Figure 12). These findings explain the process by which treatment response in advanced NSCLC is improved by ICIs, Niv and Atz, in comparison to chemotherapy (Doc), with or without Ram, during second-line and subsequent treatments.

However, there are reported cases of primary or adaptive resistance to cancer immunotherapeutic agents such as Niv [1,4,6]. Moreover, some patients who are resistant to immunotherapy have been reported to respond to a treatment regimen that includes Ram [10]. These reports not only suggest the importance of appropriate therapeutic selection for each patient, but also the importance of conducting a detailed investigation of patient characteristics that are most likely to benefit from each therapeutic agent.

Few reports have evaluated the efficacy of Pem in a refractory or relapsed advanced PD-L1-negative (<1%) NSCLC patient group. Therefore, we were unable to include Pem in an analysis of a patient group lacking the PD-L1 constraint. Although an efficacy analysis was performed on the whole patient population, including Pem, the results of which are shown (Tables S5–S8), these results must be interpreted with caution on account of the heterogeneity of PD-L1 status. Further validation of the efficacy and safety profile of Pem in a refractory or relapsed PD-L1-negative (<1%) advanced NSCLC group of patients may be necessary.

In an inclusive efficacy analysis of the six RCTs (REVEL, CheckMate057, Check-Mate017, OAK, LUME-Lung 1, KEYNOTE-010), conceptual heterogeneity existed in patient inclusion criteria for PD-L1 status in each RCT. That is, while only PD-L1-positive (≥1%) patients were included in KEYNOTE-010, PD-L1 status was not taken into consideration in patient inclusion for other RCTs. This may be considered as a limitation when comparing ICIs. By contrast, comparisons between Ram+Doc and Nin+Doc were considered to be feasible. Our analysis showed that there was no statistically significant difference in OS between Ram+Doc and Nin+Doc in the non-squamous subgroup.

Although only Niv and Atz were included and Pem was excluded, in the comparison of ICIs in the patient group lacking the PD-L1 constraint, the results led to some interesting observations, especially regarding efficacy outcomes. Subgroup analysis by histological type showed that the highest OS for a drug differed according to histological type as follows: in squamous cell carcinoma, Niv showed the highest OS, whereas in non-squamous cell carcinoma, Atz showed the highest OS. Hence, although safety outcomes cannot be analyzed via histology, these results suggested that histology-specific treatment strategies should be considered for recurrent NSCLC. However, direct comparative RCTs may be required to confirm our results.
Figure 12. Mechanisms contributing to the efficacy of immunotherapy in refractory and relapsed cases. Initial treatments, such as chemotherapy, eliminate cancer cells but treatment-resistant cancer cells may survive. Treatment induces inflammation in major vicinities, leading to the formation of active T cells. The antitumor effects of these activated T cells are suppressed by cancer cells. However, immunotherapy reactivates T cells, thereby causing an anticancer effect in tissues with recurrent cancer.

Analyses of safety profiles that were performed using the six RCTs, REVEL, CheckMate057, CheckMate017, OAK, LUME-Lung 1, and KEYNOTE-010, allowed the incidence of G3-5AEs among Doc, Ram+Doc, Niv, Atz, Pem and Nin+Doc to be compared. The results indicated that Niv was the safest, followed by Atz, Pem2, Pem10, Doc, Ram+Doc, and Nin+Doc (Table S10). The differences between Ram+Doc and Doc, Niv and Doc, Atz and Doc, Pem2 and Doc Pem10 and Doc, Nin+Doc and Doc, Niv and Ram+Doc, Atz and Ram+Doc, Pem2 and Ram+Doc, Pem10 and Ram+Doc, Atz and Niv, Pem2 and Niv, Pem10 and Niv, Nin+Doc and Niv, Nin+Doc and Atz, Nin+Doc and Pem2, and Nin+Doc and Pem10 were found to be statistically significant (Table S9). The results of this safety profile analysis also provided clinically relevant information for the selection of therapeutic agents for refractory or relapsed advanced NSCLC.

Several limitations were associated with this study. As this was an NMA of RCTs that were performed separately, inconsistencies between these RCTs were not addressed in this study. Secondly, the inclusion criteria for each study differed. For example, there was a slight difference in the classification of patient stages; only stage IV patients were included in the REVEL trial, whereas stage IIIB and IV patients were included in the remaining three trials (CheckMate057, CheckMate017, and OAK). Furthermore, the number of prior treatments also differed; the OAK trial included one or two prior systemic cancer treatments, whereas the remaining three trials (REVEL, CheckMate017, and CheckMate057) included...
only one systemic cancer treatment. To address this limitation, we performed a sensitivity analysis which confirmed that the inclusion/exclusion of patients who had received two prior treatments had little effect on the final conclusions of this NMA. Nevertheless, this heterogeneity cannot be completely ignored since the sensitivity analysis cannot eliminate all effects of heterogeneity on the results as well as conclusions. Thirdly, the efficacy profile of Pem in a population that included a PD-L1-negative (<1%) group was not evaluated in sufficient detail in the present NMA. This was due to the paucity of existing RCTs of Pem in the patient groups including refractory or recurrent PD-L1-negative (<1%) NSCLC patients. Therefore, the efficacy profile of Pem in a patient population including a group of patients with PD-L1-negative refractory or relapsed advanced NSCLC may need to be investigated in more detail in the future. Finally, the group of patients included in this analysis received platinum-based chemotherapy as the first-line treatment; these patients did not receive the current standard of care, that is, a combination of ICIs and platinum-based chemotherapy. Therefore, the results of this analysis are applicable only to a group of patients who received platinum-based chemotherapy without ICIs as the first-line treatment.

4. Materials and Methods

4.1. Systematic Review

Four databases, PubMed, CENTRAL, Embase, and SCOPUS, were searched in July 2020 for reported RCTs of ICIs or Ram+Doc for advanced NSCLC from 1946 onward. Our search strategy used keywords such as NSCLC, ICIs, ramucirumab, docetaxel, and their Medical Subject Headings in only English articles (Appendix A). All reference lists were reviewed to prevent any relevant studies from being missed.

The present study was conducted in accordance with the Preferred Reporting Items for Systematic Review and Meta-Analysis (PRISMA) [35] and the PRISMA extension statement for network meta-analysis [21]. The literature search was performed independently by two investigators (K.A. and Y.K.); in cases of conflict, a third researcher (T.Y.) participated for the purpose of conflict resolution. For the studies retrieved via a systematic review, inclusion and exclusion criteria were adapted using the PICOS approach to address clinical or methodological inter-study heterogeneity and ensure validity of the NMA.

4.2. Quality Evaluation

We evaluated the qualities of RCTs included in the present NMA using ROB2 recommended by the Cochrane Collaboration [33]. We assessed the following parameters as having a low risk, some concerns, or a high risk: (1) bias arising from the randomization process, (2) bias due to deviations from intended interventions, (3) bias due to missing outcome data, (4) bias in measuring the outcome and (5) bias in selection of the reported result. Quality evaluation was performed independently by two investigators (K.A. and S.K.), where in cases of conflict, a third investigator (T.Y.) was consulted for the purpose of conflict resolution.

4.3. Inclusion and Exclusion Criteria (Predefined PICOS)

4.3.1. Patients

The following inclusion criteria were used for this analysis: age over 18, histological or cytological confirmation of refractory or relapsed NSCLC and performance status of 0 to 2 (on a 5-point scale, with a higher number indicating poorer general condition).

4.3.2. Interventions/Comparisons

For the meta-analysis, we included patients that received at least one of the following treatments: (1) 3 mg/kg of Niv every 2 weeks, (2) 1200 mg/body of Atz every 3 weeks and (3) 10 mg/kg Ram every 3 weeks plus 75 mg Doc per square meter of body surface area every 3 weeks. The above dosages were selected based on licensed dosages and those reported in phase III trials. Doc was considered as the common comparative group since
it was the standard treatment regimen prior to the development of immunotherapy and molecular cancer therapies for refractory or recurrent advanced NSCLC [3].

4.3.3. Outcomes

The primary and secondary efficacy outcomes were OS and PFS, respectively, which were expressed using the HRs and associated 95% CrIs. The primary safety endpoint was G3–5AEs, which were expressed using RRs and associated 95% CrIs. To assess efficacy rankings, we calculated SUCRAs for OS and PFS in the overall population and in the subgroups based on histological types (non-squamous and squamous). In order to assess safety rankings, we calculated SUCRAs for the frequency of G3–5AEs. These analyses were performed only if they were obtained from studies that included data required for the analysis.

4.3.4. Study Design

Inclusion criteria for the present NMA comprised randomized, parallel group, phase III studies.

4.4. Statistical Method of NMA

We performed Bayesian NMA according to a rigorously constructed framework established by the National Institute for Health and Care Excellence [14–18,20,31,36]. These formulations have also been applied in various other clinical studies [37–43]. In this study, we used the standard Bayesian model described by Dias et al. [44–46], which assumes inconsistency and heterogeneity across all included studies [19]. The posterior distribution of the treatment effect was estimated by applying a non-informative prior distribution and Gibbs sampling based on the Markov chain Monte Carlo method. The number of iterations was set at 50,000, and the first 10,000 iterations were considered as a burn-in sample to exclude the effects of the initial values. Effect sizes were expressed as HR and RR along with associated 95% CrIs; if the 95% CrI of a HR or RR did not include 1, the difference between treatment groups was considered statistically significant. To estimate the ranking of each treatment effect, SUCRA was calculated. SUCRA values ranged from 0% to 100%, with higher SUCRA values indicating a relatively more favorable treatment [22]. We used the Brooks–Gelman–Rubin (BGR) diagnostic approach [30,31] to conduct convergence diagnoses of all comparisons. Both visual assessments and diagnoses used to confirm model convergence were conducted using the BGR method. OpenBUGS 1.4.0 (MRC Biostatistics Unit, Cambridge Public Health Research Institute, Cambridge, UK) was used for analysis and STATA (ver. 14; StataCorp, Cambridge, UK) was used to graphically display the results (College Station, TX, USA).

4.5. Ethical Aspects

Owing to the characteristics of the review and meta-analysis carried out in the present study, the need for institutional review board approval and informed consent was waived.

5. Conclusions

The Bayesian NMA comparing ICIs and Ram+Doc for refractory or relapsed advanced NSCLC in patient groups lacking the PD-L1 status constraint revealed that the OS corresponding to Niv treatment was superior to that of Ram+Doc. G3–5AEs were more common after treatment with Ram+Doc than after Atz or Niv. SUCRA values indicated that Niv displayed the highest OS and G3–5AEs. Our results indicated that treatment of refractory or relapsed advanced NSCLC with either Atz or Niv monotherapy was more effective than treatment with Doc or Ram+Doc for patients lacking the PD-L1 status constraint. Regarding PD-L1-positive (≥1%) refractory or relapsed advanced NSCLC subgroup, Pem10 was the most effective treatment, followed by Niv, Pem2, Atz, and finally Doc. Our findings may lead to more precise clinical decision-making in the future. However, verification using direct head-to-head RCTs seems to be necessary. Furthermore, characterizing the
patient profiles that benefit the most from each treatment agent prescribed for refractory or relapsed advanced NSCLC is an important topic that needs to be addressed via future clinical research.

**Supplementary Materials:** The following are available online at https://www.mdpi.com/2072-6694/13/1/52/s1. Figure S1: A scatter diagram of the SUCRA, of the efficacy in terms of PFS and safety in terms of G3–5AEs, of the four therapeutic regimens. Figure S2: Risk of bias summary. Figure S3: Risk of bias summary of LUME-Lung 1 and KEYNOTE-010. Table S1: Sensitivity analysis. Table S2: Sensitivity analysis for ranking assessment. Table S3: Key inclusion criteria of LUME-Lung 1 and KEYNOTE-010. Table S4: Main characteristic of LUME-Lung 1 and KEYNOTE-010. Table S5: The results of the OS comparison (six RCTs of REVEL, CheckMate057, CheckMate017, OAK, LUME-Lung 1, KEYNOTE-010 were included). Table S6: Results of subgroup analysis for non-squamous OS comparisons (five RCTs of REVEL, CheckMate057, OAK, LUME-Lung 1, KEYNOTE-010 were included). Table S7: Results of subgroup analysis for squamous OS comparisons (four RCTs of REVEL, CheckMate017, OAK, KEYNOTE-010 were included). Table S8: The results of the PFS comparison (six RCTs of REVEL, CheckMate057, CheckMate017, OAK, LUME-Lung 1, KEYNOTE-010 were included). Table S9: Results of the comparison of G3-5AEs incidences (six RCTs of REVEL, CheckMate057, CheckMate017, OAK, LUME-Lung 1, KEYNOTE-010 were included). Table S10: The results of SUCRA values (rank) for efficacy and safety outcomes (six RCTs of REVEL, CheckMate057, CheckMate017, OAK, LUME-Lung 1, KEYNOTE-010 were included). Table S11: Results of OS comparison in PD-L1-positive non-squamous subgroup (two RCTs of CheckMate057 and KEYNOTE-010 were included). Table S12: Results of OS comparison in PD-L1-positive squamous subgroup (two RCTs of CheckMate017 and KEYNOTE-010 were included). Table S13: SUCRA values (ranks) for OS efficacy in the PD-L1-positive group and its subgroups by histology (non-squamous or squamous) (four RCTs of CheckMate057, CheckMate017, OAK, KEYNOTE-010 were included).

**Author Contributions:** Conceptualization, K.A., R.M., Y.K., S.K., T.Y., A.T., T.O. (Tohru Ohmori), T.O. (Tsukasa Ohnishi) and H.S.; methodology, K.A., T.Y. and A.T.; validation, K.A., R.M., Y.K., S.K., T.Y., A.T., T.O. (Tohru Ohmori), T.O. (Tsukasa Ohnishi) and H.S.; formal analysis, K.A., T.Y. and A.T.; investigation, K.A., R.M., Y.K., S.K., T.Y., A.T., T.O. (Tohru Ohmori), T.O. (Tsukasa Ohnishi) and H.S.; resources, K.A., R.M., Y.K., S.K. and T.Y.; data curation, K.A., R.M., Y.K., S.K. and T.Y.; writing—original draft preparation, K.A.; writing—review and editing, K.A., Y.K., S.K., T.Y., A.T., T.O. (Tohru Ohmori), T.O. (Tsukasa Ohnishi) and H.S.; supervision, K.A., T.Y., A.T., T.O. (Tohru Ohmori), T.O. (Tsukasa Ohnishi) and H.S.; project administration, K.A., Y.K., S.K., T.Y., A.T., T.O. (Tohru Ohmori), T.O. (Tsukasa Ohnishi) and H.S.; funding acquisition, K.A. and H.S. All authors have read and agreed to the published version of the manuscript.

**Funding:** This research received no external funding.

**Institutional Review Board Statement:** This clinical study is a systematic review and meta-analysis, which used only previously published, de-identified patient data. Therefore, ethical review and approval were waived.

**Informed Consent Statement:** Written patient consent was also waived due to the retrospective nature of this study, as a systematic data review and meta-analysis.

**Data Availability Statement:** The authors confirm that the datasets analyzed during the current study are available from the corresponding author upon reasonable request.

**Conflicts of Interest:** The authors declare no conflict of interest.

**Appendix A. Search Strategies in PubMed**

Search query (Searched on 1 July 2020):
(((("nivolumab"[ALL] OR "nivolumab"[Supplementary Concept] OR “opdivo”[ALL] OR “BMS-936558”[ALL] OR “BMS936558”[ALL] OR “BMS 936558”[ALL] OR “ONO-...})})})

---

Cancers 2021, 13, 52

---

18 of 21
4538"[ALL] OR “ONO4538"[ALL] OR “ONO 4538"[ALL] OR “MDX-1106"[ALL] OR “MDX1106"[ALL] OR “MDX 1106"[ALL] OR “pembrolizumab"[ALL] OR “pembrolizumab” [Supplementary Concept] OR “keytruda”[ALL] OR “lambrolizumab”[ALL] OR “MK-3475"[ALL] OR “MK3475"[ALL] OR “MK 3475"[ALL] OR “SCH-900475"[ALL] OR “SCH90 0475"[ALL] OR “SCH 900475"[ALL] OR “Atezolizumab”[ALL] OR “Atezolizumab” [Supplementary Concept] OR “MDP3280A"[ALL] OR “MPDL3280A"[ALL] OR “MDP 3280A"[ALL] OR “RG7446"[ALL] OR “RG-7446"[ALL] OR “RG 7446"[ALL] OR “Tece- triq"[ALL] OR “immune checkpoint inhibitor”[ALL] OR “anti-PD-1”[ALL] OR “anti- PD1"[ALL] OR “anti-PD 1"[ALL] OR “pd1 inhibitor”[ALL] OR “pd1 inhibitor*”[ALL] OR “programmed cell death 1 receptor/antagonists and inhibitors”[MeSH Terms] OR “programmed cell death 1 receptor antagonists and inhibitors”[ALL] OR “programmed cell death 1 receptor antagonist*[ALL] OR “programmed cell death 1 receptor inhibitor*[ALL]) AND (“Docetaxel "[ALL] OR “Docetaxel” [Supplementary Concept] OR “Taxotere”[ALL] OR “RP-56976"[ALL] OR “RP56976"[ALL] OR “ RP 56976"[ALL] OR “ NSC-628503"[ALL] OR “ NSC628503"[ALL] OR “ NSC 628503"[ALL]) OR (“Ramucirumab”[ALL] OR “Ramucirumab” [Supplementary Concept] OR “IMC1121B “[ALL] OR “ IMC-1121B ”[ALL] OR “ IMC 1121B ”[ALL] OR “Cyramza “[ALL] OR “anti VEGFR2”[ALL] OR “anti VEGFR-2”[ALL] OR “vascular endothelial growth factor receptor-2"[ALL] OR “anti- vascular endothelial growth factor receptor-2”[ALL] OR “MOAB VEGFR-2”[ALL] OR “monoclonal antibody VEGFR-2”[ALL] OR “monoclonal antibody VEGFR2”[ALL]) AND (“Docetaxel “[ALL] OR “Docetaxel” [Supplementary Concept] OR “Taxotere”[ALL] OR “RP-56976"[ALL] OR “RP56976"[ALL] OR “ RP 56976"[ALL] OR “ NSC-628503"[ALL] OR “ NSC628503"[ALL] OR “ NSC 628503"[ALL]) AND (“Non-Small Cell Lung Cancer" OR “Non-Small Cell Lung Carcinoma" OR “Non-Small-Cell Lung Carcinoma" OR “Nonsmall Cell Lung Cancer" OR “Non-Small-Cell Lung Carcinomas" OR “NSCLC") AND (((“randomized controlled trial”[Publication Type] OR “controlled clinical trial”[Publication Type]) OR “randomized”[Title/Abstract]) OR “placebo”[Title/Abstract]) OR “clinical trials as topic”[MeSH Terms]) OR “randomly”[Title/Abstract] OR “trial”[Title]) NOT (“animals”[MeSH Terms] NOT “humans”[MeSH Terms]).

Search returned 121 results.

References

1. Wu, Z.; Man, S.; Sun, R.; Li, Z.; Wu, Y.; Zuo, D. Recent advances and challenges of immune checkpoint inhibitors in immunothera-ピー of non-small cell lung cancer. Int. Immunopharmacol. 2020, 85, 106613. [CrossRef]
2. Nasim, F.; Sabath, B.F.; Eapen, G.A. Lung cancer. Med. Clin. N. Am. 2019, 103, 463–473. [CrossRef] [PubMed]
3. Ettinger, D.S.; Wood, D.E.; Aggarwal, C.; Aisner, D.L.; Akerley, W.; Bauman, J.R.; Bharat, A.; Bruno, D.S.; Chang, J.Y.; Chirieac, L.R.; et al. NCCN guidelines insights: Non-Small Cell Lung Cancer, Version 1.2020. J. Natl. Compr. Cancer Netw. 2019, 17, 1464–1472. [CrossRef] [PubMed]
4. Onoi, K.; Chihara, Y.; Uchino, J.; Shimamoto, T.; Morimoto, Y.; Iwasaku, M.; Kaneko, Y.; Yamada, T.; Takayama, K. Immune checkpoint inhibitors for lung cancer treatment: A review. J. Immunother. Cancer 2020, 8, 1–15. [CrossRef] [PubMed]
5. Siegel, R.L.; Miller, K.D.; Jemal, A. Cancer statistics, 2020. CA Cancer J. Clin. 2020, 70, 7–30. [CrossRef] [PubMed]
6. Pasello, G.; Pavan, A.; Attili, I.; Bortolami, A.; Bonanno, L.; Menis, J.; Conte, P.; Guarneri, V. Real world data in the era of immune checkpoint inhibitors (ICIs): Increasing evidence and future applications in lung cancer. Cancer Treat. Rev. 2020, 87, 102031. [CrossRef] [PubMed]
7. Duma, N.; Santana-Davila, R.; Molina, J.R. Non-small cell lung cancer: Epidemiology, screening, diagnosis, and treatment. Mayo Clin. Proc. 2019, 94, 1623–1640. [CrossRef] [PubMed]
8. Arbour, K.C.; Riely, G.J. Systemic therapy for locally advanced and metastatic non-small cell lung cancer: A review. JAMA 2019, 322, 764–774. [CrossRef]
9. Garon, E.B.; Ciuleanu, T.E.; Arrieta, O.; Prabhak, K.; Syrigos, K.N.; Goksel, T.; Park, K.; Gorbunova, V.; Kowalsyszn, R.D.; Piikel, J.; et al. Ramucirumab plus docetaxel versus placebo plus docetaxel for second-line treatment of stage IV non-small-cell lung cancer after disease progression on platinum-based therapy (REVEL): A multicentre, double-blind, randomised phase 3 trial. Lancet 2014, 384, 665–773. [CrossRef]
10. Popat, S.; Grohé, C.; Corral, J.; Reck, M.; Novello, S.; Gottfried, M.; Radonjic, D.; Kaiser, R. Anti-angiogenic agents in the age of resistance to immune checkpoint inhibitors: Do they have a role in non-oncogene-addicted non-small cell lung cancer? Lung Cancer 2020, 144, 76–84. [CrossRef]
11. Reck, M.; Kaiser, R.; Mellemgaard, A.; Douillard, J.Y.; Orlov, S.; Krzakowski, M.; Von Pawel, J.; Gottfried, M.; Bondarenko, I.; Liao, M.; et al. Docetaxel plus nintedanib versus docetaxel plus placebo in patients with previously treated non-small-cell lung cancer (LUME-Lung 1): A phase 3, double-blind, randomised controlled trial. *Lancet Oncol.* 2014, 15, 143–155. [CrossRef]

12. Han, Y.; Liu, D.; Li, L. PD-1/PD-L1 pathway: Current researches in cancer. *Ann. J. Cancer Res. 2020, 10, 727–742.* [PubMed]

13. Chen, Y.; Zhou, Y.; Tang, L.; Peng, X.; Jiang, H.; Wang, G.; Zhuang, W. Immune-checkpoint inhibitors as the first line treatment of advanced non-small cell lung cancer: A meta-analysis of randomized controlled trials. *J. Cancer 2019, 10, 6261–6268.* [CrossRef] [PubMed]

14. Lu, G.; Ades, A.E. Combination of direct and indirect evidence in mixed treatment comparisons. *Stat. Med. 2004, 23, 3105–3124.* [CrossRef] [PubMed]

15. Lumley, T. Network meta-analysis for indirect treatment comparisons. *Stat. Med. 2002, 21, 2313–2324.* [CrossRef] [PubMed]

16. Bucher, H.C.; Guyatt, G.H.; Griffith, L.E.; Walter, S.D. The results of direct and indirect treatment comparisons in meta-analysis of randomized controlled trials. *J. Clin. Epidemiol. 1997, 50, 683–691.* [CrossRef]

17. White, I.R. Network meta-analysis. *Stat. J. 2015, 15, 951–985.* [CrossRef]

18. Hoaglin, D.C.; Hawkins, N.; Jansen, J.P.; Scott, D.A.; Itzler, R.; Cappelleri, J.C.; Boersma, C.; Thompson, D.; Larholt, K.M.; Diaz, M.; et al. Conducting indirect-treatment-comparison and network-meta-analysis studies: Report of the ISPOR Task Force on Indirect Treatment Comparisons Good Research Practices: Part 2. *Value Health 2011, 14, 429–437.* [CrossRef]

19. Jansen, J.P.; Crawford, B.; Bergman, G.; Stam, W. Bayesian meta-analysis of multiple treatment comparisons: An introduction to mixed treatment comparisons. *Value Health 2008, 11, 956–964.* [CrossRef]

20. Tonin, F.S.; Rotta, I.; Mendes, A.M.; Pontarolo, R. Network meta-analysis: A technique to gather evidence from direct and indirect comparisons. *Pharm. Pract. (Granada) 2017, 15, 943.* [CrossRef]

21. Hutton, B.; Salanti, G.; Caldwell, D.M.; Chaimani, A.; Schmid, C.H.; Cameron, C.; Ioannidis, J.P.; Straus, S.; Thorlund, K.; Jansen, J.P.; et al. The PRISMA extension statement for reporting of systematic reviews incorporating network meta-analyses of health care interventions: Checklist and explanations. *Ann. Intern. Med. 2015, 162, 777–784.* [CrossRef] [PubMed]

22. Salanti, G.; Ades, A.E.; Ioannidis, J.P. Graphical methods and numerical summaries for presenting results from multiple-treatment meta-analysis: An overview and tutorial. *J. Clin. Epidemiol. 2011, 64, 163–171.* [CrossRef] [PubMed]

23. PubMed. Available online: https://pubmed.ncbi.nlm.nih.gov/ (accessed on 1 July 2020).

24. Embase. Available online: https://www.embase.com/login (accessed on 1 July 2020).

25. Cochrane Central Register of Controlled Trials [CENTRAL]. Available online: https://www.cochranelibrary.com/ (accessed on 1 July 2020).

26. SCOPUS. Available online: https://www.scopus.com/home.uri (accessed on 1 July 2020).

27. Brahmer, J.; Reckamp, K.L.; Baas, P.; Crinò, L.; Eberhard, W.E.; Poddubskaya, E.; Antonia, S.; Pluzanski, A.; Vokes, E.E.; Holgado, E.; et al. Nivolumab versus docetaxel in advanced squamous-cell non-small-cell lung cancer. *N. Engl. J. Med. 2015, 373, 123–135.* [CrossRef] [PubMed]

28. Borghaei, H.; Paz-Ares, L.; Horn, L.; Spigel, D.R.; Steins, M.; Ready, N.E.; Chow, L.Q.; Vokes, E.E.; Felip, E.; Holgado, E.; et al. Nivolumab versus docetaxel in advanced nonsquamous non-small-cell lung cancer. *N. Engl. J. Med. 2015, 373, 1627–1639.* [CrossRef] [PubMed]

29. Hutton, B.; Salanti, G.; Caldwell, D.M.; Chaimani, A.; Schmid, C.H.; Cameron, C.; Ioannidis, J.P.; Straus, S.; Thorlund, K.; Jansen, J.P.; et al. The PRISMA extension statement for reporting of systematic reviews incorporating network meta-analyses of health care interventions: Checklist and explanations. *Ann. Intern. Med. 2015, 162, 777–784.* [CrossRef] [PubMed]

30. Borghaei, H.; Paz-Ares, L.; Horn, L.; Spigel, D.R.; Steins, M.; Ready, N.E.; Chow, L.Q.; Vokes, E.E.; Felip, E.; Holgado, E.; et al. Nivolumab versus docetaxel in advanced nonsquamous non-small-cell lung cancer. *N. Engl. J. Med. 2015, 373, 1627–1639.* [CrossRef] [PubMed]

31. Brooks, S.P.; Roberts, G.O. Convergence assessment techniques for Markov chain Monte Carlo. *Stat. Comput. 1998, 8, 319–335.* [CrossRef] [PubMed]

32. Brooks, S.P.; Gelman, A. General methods for monitoring convergence of iterative simulations. *J. Comput. Graph. Stat. 1996, 7, 434–455.* [CrossRef]

33. Herbst, R.S.; Baas, P.; Kim, D.W.; Felip, E.; Pérez-Gracia, J.L.; Han, J.Y.; Molina, J.; Kim, J.H.; Dubos Arvis, C.; Ahn, M.J. Pembrolizumab versus docetaxel for previously treated, PD-L1-positive, advanced non-small-cell lung cancer (KEYNOTE-010): A randomised controlled trial. *Lancet 2016, 387, 1540–1550.* [CrossRef]

34. Higgins, J.P.; Thomas, J.; Chandler, J.; Cumpston, M.; Li, T.; Page, M.J.; Welch, V. *Cochrane Handbook for Systematic Reviews of Interventions,* 9th ed.; John Wiley & Sons, Inc.: Hoboken, NJ, USA, 2019.

35. Chamoto, K.; Al-Habsi, M.; Honjo, T. Role of PD-1 in immunity and diseases. *Curr. Top. Microbiol. Immunol. 2017, 410, 75–97.* [CrossRef]

36. Moher, D.; Liberati, A.; Tetzlaff, J.; Altman, D.G.; PRISMA Group. Preferred reporting items for systematic reviews and meta-analyses: The PRISMA statement. *PLoS Med. 2009, 6, e1000097.* [CrossRef]

37. Siemieniuk, R.A.; Bartoszko, J.J.; Ge, L.; Zeraatkar, D.; Izcovich, A.; Pardo-Hernandez, H.; Rochwerg, B.; Lamontagne, F.; Han, M.A.; Kum, E.; et al. Drug treatments for covid-19: Living systematic review and network meta-analysis. *BMJ 2020, 370, m2980.* [CrossRef] [PubMed]
38. Hodkinson, A.; Bower, P.; Grigoroglou, C.; Zghebi, S.S.; Pinnock, H.; Kontopantelis, E.; Panagioti, M. Self-management interventions to reduce healthcare use and improve quality of life among patients with asthma: Systematic review and network meta-analysis. *BMJ* 2020, 370, m2521. [CrossRef] [PubMed]

39. Ge, L.; Sadeghirad, B.; Ball, G.D.C.; Da Costa, B.R.; Hitchcock, C.L.; Svendrovski, A.; Kiflen, R.; Quadri, K.; Kwon, H.Y.; Karamouzian, M.; et al. Comparison of dietary macronutrient patterns of 14 popular named dietary programmes for weight and cardiovascular risk factor reduction in adults: Systematic review and network meta-analysis of randomised trials. *BMJ* 2020, 369, m696. [CrossRef] [PubMed]

40. Elaidi, R.; Phan, L.; Borchiellini, D.; Barthelemy, P.; Ravaud, A.; Oudard, S.; Vano, Y. Comparative efficacy of first-line immune-based combination therapies in metastatic renal cell carcinoma: A systematic review and network meta-analysis. *Cancers* 2020, 12, 1673. [CrossRef]

41. Ando, K.; Kishino, Y.; Homma, T.; Kusumoto, S.; Yamaoka, T.; Tanaka, A.; Ohnori, T.; Ohnishi, T.; Sagara, H. Nivolumab plus ipilimumab versus existing immunotheapies in patients with PD-L1-positive advanced non-small cell lung cancer: A systematic review and network meta-analysis. *Cancers* 2020, 12, 1905. [CrossRef]

42. Ando, K.; Akimoto, K.; Sato, H.; Manabe, R.; Kishino, Y.; Homma, T.; Kusumoto, S.; Yamaoka, T.; Tanaka, A.; Ohnori, T.; et al. Brigatinib and alectinib for ALK rearrangement-positive advanced non-small cell lung cancer with or without central nervous system metastasis: A systematic review and network meta-analysis. *Cancers* 2020, 12, 942. [CrossRef]

43. Zhao, Y.; Liu, J.; Cai, X.; Pan, Z.; Liu, J.; Yin, W.; Chen, H.; Xie, Z.; Liang, H.; Wang, W.; et al. Efficacy and safety of first line treatments for patients with advanced epidermal growth factor receptor mutated, non-small cell lung cancer: Systematic review and network meta-analysis. *BMJ* 2019, 367, l5460. [CrossRef]

44. Dias, S.; Welton, N.J.; Sutton, A.J.; Caldwell, D.M.; Lu, G.; Ades, A.E. Evidence synthesis for decision making 4: Inconsistency in networks of evidence based on randomized controlled trials. *Med. Decis. Mak.* 2013, 33, 641–656. [CrossRef]

45. Dias, S.; Sutton, A.J.; Welton, N.J.; Ades, A.E. Evidence synthesis for decision making 3: Heterogeneity—subgroups, meta-regression, bias, and bias-adjustment. *Med. Decis. Mak.* 2013, 33, 618–640. [CrossRef]

46. Dias, S.; Sutton, A.J.; Ades, A.E.; Welton, N.J. Evidence synthesis for decision making 2: A generalized linear modeling framework for pairwise and network meta-analysis of randomized controlled trials. *Med. Decis. Mak.* 2013, 33, 607–617. [CrossRef] [PubMed]