The best platinum regimens for chemo-naive incurable non-small cell lung cancer: network meta-analysis

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Platinum regimens still play a key role in chemotherapy for incurable non-small cell lung cancer (NSCLC). Although guidelines list many platinum regimens, the best regimens have not yet clarified. Electronic searches were carried out during November 26th-28th, 2016. We included individually randomized trials comparing two or more platinum regimes for incurable chemo-naive NSCLC published in English full papers. The platinum doublets should be either Cisplatin (CDDP), Carboplatin (CBDCA), or Nedaplatin (CDGP) plus one of the third-generation agents. The platinum triplet should be the doublet plus bevacizumab (BEV). The data were independently extracted and cross-checked by two investigators. We did not observe heterogeneity (whole network level $Q = 28.9, df = 34, P = 0.717$) among 59 pairwise comparisons from 45 studies with 16141 cases for the primary outcome, hazard ratio for overall survival (HRos). Using CBDCA + Paclitaxel (PTX) + BEV as a common comparator, CDGP + Docetaxel (DTX) (HRos = 0.98, 95%CI: 0.75–1.29, $P = 0.884$), CDDP + Tegafur gimeracil oteracil (S1) (HRos = 1.23, 95%CI: 0.96–1.57, $P = 0.099$), CBDCA + S1 (HRos = 1.23, 95%CI: 0.99–1.53, $P = 0.062$), and CDGP + Gemcitabine (GEM) (HRos = 1.24, 95%CI: 0.71–2.17, $P = 0.45$) did not have significantly poorer HRos. We suggest that these regimens as acceptable first-choice regimens.

Non-small cell lung cancer (NSCLC), which is currently the most common malignant neoplasm in the world, is one of the leading causes of cancer death worldwide¹. In more than a half of cases, the NSCLC is detected after the disease has already progressed to an incurable stage. For such patients, chemotherapy is usually the first-choice treatment option because accumulated evidence has revealed that current standard chemotherapy treatments have substantial benefits for advanced, locally advanced, and recurrent NSCLC. Traditionally, some of the platinum regimens have been regarded as the standard first-line regimens for NSCLC for non-elderly patients with good performance status who do not have major co-morbidities. The currently preferred platinum doublets are combinations of one of the platinum agents and one of the third-generation chemotherapy agents. Currently accepted platinum triplets are combinations of these platinum doublets and bevacizumab (BEV)². Even though epidermal growth factor receptor-tyrosine kinase inhibitors, anaplastic lymphoma kinase inhibitors, and immune check-point inhibitors have recently been preferred for certain subgroups of NSCLC patients, these platinum regimens still play a key role in chemotherapy for incurable NSCLC².³

Current guidelines list many platinum doublets and triplets as recommended therapeutic options for advanced NSCLC. However, the single best regimen among platinum regimens has not yet been clarified for various reasons: inconsistent results from trials, difficulty in interpreting results from non-inferiority trials, lack of statistical power to detect subtle survival difference, differences of inclusion criteria such as performance status and age, and inconsistency of primary outcomes of original trials. To solve this, meta-analysis is a useful method to identify the best regimen⁴. Nonetheless, classical head-to-head pairwise meta-analysis cannot satisfactorily answer this clinical question because of the deficiency of direct comparisons among the numerous potentially best regimens.

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Therefore, some previous meta-analyses compared groups of regimens, for example, cisplatin (CDDP) regimens versus carboplatin (CBDCA) regimens or BEV regimens versus non-BEV regimens. On the other hand, network meta-analysis is a recently developed technique to integrate available data. This analysis has the advantages of allowing collective comparison among multiple treatment arms and the potential precision gains from combining direct and indirect evidence.

The goal of the current network meta-analysis is to identify and rank the best standard regimens by comparing the effectiveness and safety of a variety of the platinum regimens as first-line chemotherapies for advanced, locally advanced, and recurrent NSCLC.

Material and Methods

Protocol registration. This protocol of the systematic review and network meta-analysis has been uploaded on the website of International Prospective Register of Systematic Reviews (42016052455). We have composed this protocol following the Preferred Reporting Items for Systematic Reviews and Meta-Analyses statement and that for network meta-analysis. Institutional Review Board approval and patient informed consent were waived due to the review nature of this study.

Study search. Search formulas for electronic databases were created with the support of Cochrane Japanese. Search formulas for MEDLINE, EMBASE, the Web of Science Core Collection, and the Cochrane Central Register of Controlled Trials are presented in Supplementary Text 1. The search for each electronic database was carried out during November 26th-28th, 2016. An additional manual search was conducted by two investigators (NH and AN) independently. Candidate articles were first screened and then scrutinized independently by the two investigators. Discrepancies found during the study selection process were resolved by discussion between the two investigators.

Inclusion criteria. Publication type and trial design. We included individually randomized trials comparing two or more platinum regimens for incurable NSCLC, which have been reported and published in English full papers. We allowed a trial with three or more arms. We allowed all of superiority, non-inferiority, phase II, phase III, non-blinded, single-blinded, and double-blinded trials. A trial evaluating only the co-secondary outcomes of our analysis was allowed. Included patients should have been randomized before the first-line chemotherapy initiation. Thus, randomization just before the maintenance therapy was not accepted. We carefully checked for and avoided duplicate uses of the same study.

Treatments. Our concern was with the first-line platinum doublet and triplet chemotherapy regimens. Platinum agents should be either CDDP, CBDCA or Neadapl atin (CDGP). We disregarded regimens with Oxaliplatin. The counterpart of the platinum doublet had to be one of the following third-generation chemotherapy agents: Docetaxel (DTX), Paclitaxel (PTX), Vinorelbine (VNR), Gemcitabine (GEM), Irinotecan (CPT-11), Pemetrexed (PEM), and Tegafur gimeracil oteracil (S1). We regarded albumin-bound PTX and PTX-poliglumex as PTX. A platinum triplet had to be a combination of a platinum doublet and BEV. We did not include the following regimens: single agent chemotherapies, non-platinum doublets/triplets, regimens without the third-generation chemotherapy agent, regimens with Oxaliplatin, regimens with immune check-point inhibitors, and regimens that contained any targeted therapies for NSCLC with oncogenic driver mutation such as tyrosine kinase inhibitors. Any perioperative chemotherapy, adjuvant chemotherapy, neoadjuvant chemotherapy, and radio-chemotherapy were also excluded. We did not include studies that planned to stop the first-line regimen before administration of the third course.

Regimens that used the same medication were evaluated collectively regardless of administration root, speed, dosage, and schedule. We classified the treatment regimen based on the first-line chemotherapy regardless of maintenance, second-line, and later-line treatment. Similarly, we focused only on the first-line regimen of the cross-over trial. We equated placebo with “no treatment.” For example, “CDDP + PEM + (placebo of BEV)” arm was identical to the “CDDP + PEM” arm for our analysis.

Patients. Chemo-naive patients with advanced, or locally-advanced, or recurrent NSCLC were included. Although the tumor, node, metastasis classification has been updated every 4–8 years, we accepted the TNM classification regardless of version difference. Recurrent cancer patients with a history of operation or radiotherapy were accepted unless these patients had never undergone chemotherapy. Patients with a history of any adjuvant chemotherapy, neoadjuvant chemotherapy, or radio-chemotherapy were excluded. The age, sex, performance status, co-morbidities, and organ functions of patients were not questioned. Any study focusing on patients with large cell neuroendocrine carcinoma was planned to be excluded though this carcinoma is usually classified as NSCLC. If a regimen included PEM or BEV, the pathological type had to be limited to adenocarcinoma or non-squamous carcinoma.

Quality assessment. We assessed the quality of original studies using six domains of the Cochrane Risk of Bias evaluation sheet: selection, performance, detection, attrition, reporting, and other biases.

Outcomes. The primary outcome was hazard ratio (HR) for overall survival (OS, HROS). The co-secondary outcomes were HR for progression-free survival (FPS, HRFPS), odds ratio (OR) for response rate (RR, ORRR), and OR for severe adverse event (SAE) including neutropenia, anemia, thrombopenia, febrile neutropenia, and nausea. Adverse events with a severity, defined with Common Terminology Criteria for Adverse Events, of grade-three or higher were counted.

Evaluation of disease progression to assess the PFS and evaluation of objective response to assess RR should not have greatly deviated from the Response Evaluation Criteria In the Solid Tumors 2000 guidelines and the
2009 revised guidelines. Time to progression and time to treatment failure were not regarded as PFS. When disease progression and objective response were evaluated both by physicians caring for the patients and by the blinded independent central review board, we chose the data based on a pre-specified endpoint in each original report. If this was not clear, we used data from the blinded independent central review. The number of SAEs were counted on a patient basis, not on a per-cycle basis.

**Data extraction.** Data for the included studies, such as author name, publication year, country of origin, numbers of patients randomized, chemotherapy regimen, and data related to the study outcomes such as OS, PFS, RR, and SAE were extracted by the two investigators (NH and AN) independently. The data extracted by the two investigators were cross-checked and any discrepancies were discussed between them. We extracted data from non-inferiority studies using the same method as for superiority trials. For studies with three or more arms, data on every pairwise comparison were extracted. For example, if a three-arm study evaluated CDDP + GEM, CDDP + DTX, and DTX monotherapy, we used only the data concerning CDDP + GEM and CDDP + DTX. When two arms used the same anti-cancer medication in a three-arm trial, the outcomes in the two arms sharing the same medication were merged prior to the main analysis. For example, for a three-arm trial with (a) CDDP + PEM + high-dose BEV, (b) CDDP + PEM + low-dose BEV, and (c) CDDP + GEM, (a + b) and (c) were compared. When updated data for survival was available, the most recently updated data were preferred. When necessary, we adopted Parmar’s method to obtain survival data. Intention-to-treat analysis was preferred over full-analysis-set analysis and per-protocol analysis when two or more of these were available.

**Statistical analyses.** We pooled the logarithm of OR, HR, and their SE using the frequentist weighted least squares approach random-model network meta-analysis. All the binary outcomes were transformed to OR preceding the network meta-analysis. When one or more cells in a two-by-two contingency were zero, 0.5 was added to all the cells. When a network diagram showed two or more independent loops, we evaluated only the loop that contained major platinum regimes such as CDDP added to all the cells. When a network diagram showed two or more independent loops, we evaluated only the preceding the network meta-analysis. When one or more cells in a two-by-two contingency were zero, 0.5 was used only the data concerning CDDP + GEM and CDDP + DTX. When two arms used the same anti-cancer medication in a three-arm trial, the outcomes in the two arms sharing the same medication were merged prior to the main analysis. For example, for a three-arm trial with (a) CDDP + PEM + high-dose BEV, (b) CDDP + PEM + low-dose BEV, and (c) CDDP + GEM, (a + b) and (c) were compared. When updated data for survival was available, the most recently updated data were preferred. When necessary, we adopted Parmar’s method to obtain survival data. Intention-to-treat analysis was preferred over full-analysis-set analysis and per-protocol analysis when two or more of these were available.

**Results**

**Study search.** We first found 3405 and six articles by electronic and hand searches, respectively. Of 3411 articles that met the preliminary criteria, 162, 3112, and 89 were excluded through removal of duplication, title/abstract screening, and full-article scrutinizing, respectively (Fig. 1). We finally found 48 eligible articles (Fig. 1, Table 1, Supplementary Text 3).

**Characteristics of included studies.** The included studies were reported from a variety of countries all over the world, most of which were European or East Asian nations, and the USA (Table 1). The articles were published during 2000–2015. Among 48 articles, 18 were phase II studies, 26 were phase III studies, and evaluated OS as primary endpoints, 21 included ECOG 0–1 cases, and other 21 included ECOG 0–2 cases. We regarded two studies as three-arm studies, two studies as four-arm studies, and the other 44 as two-arm studies. Therefore, we eventually evaluated 102 arms, of which 92 were platinum doublet and 10 were platinum triplet. For example, if a three-arm study evaluated CDDP + PEM + high-dose BEV, CDDP + PEM + low-dose BEV, and CDDP + GEM, (a + b) and (c) were compared. When updated data for survival was available, the most recently updated data were preferred. When necessary, we adopted Parmar’s method to obtain survival data. Intention-to-treat analysis was preferred over full-analysis-set analysis and per-protocol analysis when two or more of these were available.

**Sensitivity analyses are planned:** (i) Fixed-model network meta-analysis instead of random-model. (ii) Random-model network meta-analysis using data from the phase III trials.

**Efficacy analysis.** Data for HRs was obtainable in 45 studies with 16141 cases (Table 1, Fig. 2). HRs presented in 59 pairwise comparisons ranged from 0.68 to 1.22 with a median of 0.95. Q statistics and a test for heterogeneity did not reveal inconsistency at any level: whole network level (Q = 28.9, df = 34, P = 0.717), within designs (Q = 11.8, df = 16, P = 0.760), and between design (Q = 17.1, df = 18, P = 0.516) (Fig. 2).

CDDP + DTX, which was evaluated in a phase III trial recruiting only squamous cancers, showed the best OS followed by CBDC + PTX + BEV, CBDC + PEM + BEV, CDDP + PEM, CBDC + PEM, and CDDP + CPT-11 in this order (Fig. 3, Supplementary Table 2). HRs between any pair of these six regimens were not significant (P = 0.516) (Supplementary Table 2). We selected CBDC + PTX + BEV as the common comparator throughout this study because this regimen showed the best OS among the CDDP/CBDC regimens. Using CBDC + PTX + BEV as a common comparator, CDDP + S1 (HRs = 1.23, 95%CI: 0.96–1.57, P = 0.099),
CBDCA + S1 (HRos = 1.23, 95%CI: 0.99–1.53, P = 0.062), and CDGP + GEM (HRos = 1.24, 95%CI: 0.71–2.17, P = 0.45) did not have significantly poorer HRos. Compared to CBDCA + PTX + BEV, the other regimens showed poorer survival assessed by HRos (P < 0.05 for all, Fig. 3, Supplementary Table 2).

Sensitivity analyses for HRos using the fixed model and using data only from phase III trials generally replicated the results (Supplementary Figure 1). A forest plot of HRos using CDGP + DTX and CDDP + CPT-11 as a common comparator is also shown to compare squamous NSCLC regimens (Supplementary Figure 1).

Three BEV regimens were high ranked for both PFS and RR. The lowest HRpfs was observed in CBDCA + PEM + BEV followed by CDDP + GEM + BEV, CDDP + PEM, and CBDCA + PTX + BEV in that order. The highest RR was shown by CBDCA + PEM + BEV followed by CBDCA + PTX + BEV, CDDP + GEM + BEV, and CBDCA + PEM. Notably, the CDGP + DTX arm had the best OS despite PFS and OR with lower ranks (Fig. 3).

Safety analysis. Neutropenia was most frequently observed for CDGP + DTX and CDDP + VNR, while S1 regimens were associated with significantly less neutropenia (P < 0.01 for both). Patients treated by PEM and GEM regimens such as CBDCA + PEM + BEV, CBDCA + PEM, CDDP + GEM + BEV, CDDP + GEM, and CBDCA + GEM had a significantly higher risk of both grade III anemia and thrombopenia when compared to CBDCA + PTX + BEV (P < 0.05 for all). Lack of statistical power due to low occurrence made it difficult to detect the difference in risk for febrile neutropenia and nausea (Fig. 3).

Discussion
We carried out the first network meta-analysis to compare platinum doublet and triplet regimens for chemo-naïve incurable NSCLC. Among the 18 platinum regimens, CDGP + DTX, CBDCA + PTX + BEV, CBDCA + PEM + BEV, CDDP + PEM, and CDDP + CPT-11 in this rank order had the best performance in the primary endpoint, OS. In addition, we evaluated PFS, RR, and adverse events of grade III or higher as the
| Study                  | Country | Phase | Primary outcome | Pathology | Arm   | Stage, Performance status | Regimens                                                                 | Patients | median age |
|-----------------------|---------|-------|-----------------|-----------|-------|--------------------------|--------------------------------------------------------------------------|----------|------------|
| Bennouna (2014)        | France  | II    | DCR             | NSq       | 2     | IIIb, IV, Rec KPS ≥ 80%   | CDDP (75 mg/m²), PEM (500 mg/m²) CDDP (80 mg/m²), VNR (80 mg/m² (d 1.8 po)) | 153      | 62         |
| Biesma (2011) NVALT-3 | Netherlands | III | QOL             | NSCLC     | 2     | III, IV ECOG 0-2         | CBDA (AUC 5), GEM (1250 mg/m² (d 1.8)) CBDA (AUC 5), PTX (175 mg/m²) | 182      | 74         |
| Chang (2008)           | China   | NS    | RR              | NSCLC     | 2     | IIIb, IV ECOG 0-2        | CDDP (80 mg/m²), GEM (1000 mg/m² (d 1.8,1,15)) CDDP (80 mg/m²), VNR (20 mg/m² (d 1.8,1,15)) | 83       | 62         |
| Chen (2004)            | Taiwan  | II    | NS              | NSCLC     | 2     | IIIb, IV ECOG 0-2        | CBDA (AUC 6), PTX (160 mg/m²) CBDA (60 mg/m²), PTX (160 mg/m²) | 81       | 75         |
| Chen (2006)            | Taiwan  | II    | Neuropathy      | NSCLC     | 2     | IIIb, IV ECOG 0-2        | CDDP (60 mg/m²), VNR (25 mg/m² (d 1.8)) CDDP (60 mg/m²), DTX (60 mg/m²) | 94       | 63         |
| Comella (2000)         | Italy   | III   | OS              | NSCLC     | 3     | IIIb, IV ECOG 0-1        | CBDA (120 mg/m²), VNR (30 mg/m² (weekly)) CDDP (100 mg/m²), GEM (1000 mg/m² (d 1.8,1,15)) | 120      | 62         |
| Douillard (2005)       | France  | II    | RR              | NSCLC     | 2     | IV                        | CDDP (100 mg/m²), DTX (75 mg/m²) CDDP (100 mg/m²), VNR (30 mg/m² (d 1.8,1,15)) | 239      | 57         |
| Edelman (2004)         | USA     | II    | OS              | NSCLC     | 2     | IIIb, IV ECOG 0-1        | CBDA (AUC 5.5), GEM (1000 mg/m² (d 1.8)) CBDA (100 mg/m²), VNR (25 mg/m² (d 1.8)) | 204      | 60         |
| Fossella (2003) TAX 326 | USA     | III   | OS (Non-inf)    | NSCLC     | 3     | IIIb, IV, Rec KPS ≥ 70%   | CBDA (AUC 5), DTX (75 mg/m²) CBDA (AUC 6), DTX (75 mg/m²) | 1218     | 60         |
| Galetta (2015) ERACLE  | Italy   | III   | QOL             | NSq       | 2     | IIIb, IV ECOG 0-1        | CDDP (75 mg/m²), PEM (500 mg/m²) CBDA (AUC 6), PTX (200 mg/m²), Bev (15 mg/kg) | 118      | 62         |
| Gebbia (2003)          | Italy   | III   | TTP, OS         | NSCLC     | 4     | IIIb, IV ECOG 0-2        | CDDP (100 mg/m²), VNR (25 mg/m² (d 1.8)) CDDP (100 mg/m²), GEM (1400 mg/m² (d 1.8)) | 278      | 62         |
| Gebbia (2010)          | Italy   | II    | QOL, AE, symptom| NSCLC     | 2     | IIIb, IV ECOG 0-1        | CDDP (75 mg/m²), DTX (75 mg/m²) CDDP (80 mg/m²), VNR (30 mg/m² (d 1.8,1,15)) | 86       | 62         |
| Gronberg (2009)        | Norway  | III   | QOL             | NSq #     | 2     | IIIb, IV ECOG 0-2        | CBDA (AUC 5), PEM (500 mg/m²) CBDA (AUC 5), GEM (1000 mg/m² (d 1.8,1,15)) | 329      | 65         |

Continued
| Study                  | Country    | Phase | Primary outcome | Pathology | Arm | Stage | Performance status | Regimens                                                                 | Patients | median age |
|-----------------------|------------|-------|-----------------|-----------|-----|-------|-------------------|---------------------------------------------------------------------------|----------|------------|
| Helbekkmo (2007)      | Norwegian  | III    | OS              | NSCLC     | 2   | IIIb, IV | ECOG 0-2          | CBDCA (AUC 5), VNR (25 mg/m² (d 1,8)) CBDCA (AUC 5), GEM (1000 mg/m² (d 1,8)) | 444      | 67         |
| Johnson (2004)        | USA        | II     | TTP, RR         | NSq #     | 3≥2 | IIIb, IV, Rec | ECOG 0-2          | CBDCA (AUC 6), PTX (200 mg/m²), BEV (7.5 or 15 mg/kg) CBSCA (AUC 6), PTX (200 mg/m²) | 79       | 63         |
| Kader (2013)          | Egypt      | II     | Toxicity, PFS   | NSq       | 2   | IIIb, IV | ECOG 0-2          | CBDCA (AUC 5), PTX (60 mg/m²), BEV (7.5 mg/kg) CBDDP (75 mg/m²), PEM (500 mg/m²) | 41       | 52         |
| Kawahara (2013)       | Japan      | II     | PFS             | NSCLC     | 2   | IIIb, IV, Rec | ECOG 0-1          | CBDCA (AUC 6), DTX (60 mg/m²) CBSCA (AUC 6), PTX (200 mg/m²) | 90       | 67         |
| Khodadad (2014)       | Iran       | NS     | PFS             | NSCLC     | 2   | IIIb, IV | ECOG 0-2          | CBDDP (75 mg/m²), DTX (75 mg/m²) CBSCA (AUC 5), PTX (200 mg/m²) | 100      | 51         |
| Kabota (2015)         | Japan      | III    | OS (Non-inf)    | NSCLC     | 2   | IIIb, IV, Rec | ECOG 0-1          | CBDDP (60 mg/m² (d8)), S1 (80 mg/m² (d 1-14 po bid)) CBDDP (80 mg/m²), DTX (60 mg/m²) | 608      | 62         |
| Langer (2007)         | USA        | II     | OS              | NSCLC     | 2   | IIIb, IV, Rec | ECOG 2             | CBSCA (AUC 6), PTX (200 mg/m²) CBDDP (60 mg/m²), GEM (1000 mg/m² (d 1,8)) | 103      | 66         |
| Martoni (2005)        | Italy      | III    | OR, TTP *       | NSCLC     | 2   | IIIb, IV, Rec | KPS ≥ 70%          | CBDDP (75 mg/m²), VNR (25 mg/m² (d 1,8)) CBDDP (75 mg/m²), GEM (1200 mg/m² (d 1,8)) | 286      | 63         |
| Mazzanti (2003)       | Italy      | II     | RR              | NSCLC     | 2   | IIIb, IV | ECOG 0-2          | CBDDP (60 mg/m²), GEM (1200 mg/m² (d 1,8)) CBSCA (AUC 5), GEM (1200 mg/m² (d 1,8)) | 125      | 63         |
| Minami (2013)         | Japan      | II     | PFS             | NSCLC     | 2   | IIIb, IV | ECOG 0-1          | CBSCA (AUC 6), PTX (200 mg/m²) CBSCA (AUC 5), GEM (1000 mg/m² (d 1,8)) | 50       | 64         |
| Niho (2012)           | Japan      | II     | PFS             | NSq       | 2   | IIIb, IV, Rec | ECOG 0-1          | CBSCA (AUC 6), PTX (200 mg/m²), BEV (15 mg/kg) CBSCA (AUC 6), PTX (200 mg/m²) | 180      | 61         |
| Ohe (2007)            | Japan      | III    | OS (Non-inf)    | NSCLC     | 4   | IIIb, IV | ECOG 0-1          | CBDDP (80 mg/m²), CPT-11 (60 mg/m² (d1,8,15)) CBSCA (AUC 6), PTX (200 mg/m²) | 602      | 62         |
| Okamoto (2010)        | Japan      | III    | OS (Non-inf)    | NSCLC     | 2   | IIIb, IV | ECOG 0-1          | CBSCA (AUC 6), S1 (80 mg/m² (d 1-14 po bid)) CBSCA (AUC 6), PTX (200 mg/m²) | 564      | 64         |

Continued
| Study                  | Country      | Phase | Primary outcome | Pathology | Arm | Stage, | Patients | median age |
|-----------------------|--------------|-------|-----------------|-----------|-----|--------|----------|------------|
| Patel (2013) PointBreak | USA          | III   | OS              | NSq       | 2   | IIIb, IV | 939      | 65         |
|                       |              |       |                 |           |     | ECOG 0-1 |          |            |
|                       |              |       |                 |           |     | CBDDCA (AUC 6), PEM (500 mg/m2), BEV (15 mg/kg) |          |            |
|                       |              |       |                 |           |     | CBDDCA (AUC 6), PTX (200 mg/m2), BEV (15 mg/kg) |          |            |
| Reck (2009) AVal      | Germany      | III   | PFS             | NSq       | 3≈ 2 | IIIb, IV, Rec | 1043     | 58         |
|                       |              |       |                 |           |     | ECOG 0-1 |          |            |
|                       |              |       |                 |           |     | CBDDPA (80 mg/m2), GEM (1250 mg/m2 (d 1,8)), BEV (7.5/15 mg/kg) |          |            |
|                       |              |       |                 |           |     | CBDDPA (80 mg/m2), GEM (1250 mg/m2 (d 1,8)) |          |            |
| Rodrigues (2011)      | Argentina    | III   | G3/4PFS         | NSq       | 2   | IIIb, IV | 260      | 60         |
|                       |              |       |                 |           |     | ECOG 0-2 |          |            |
|                       |              |       |                 |           |     | CBDDCA (AUC 5), PEM (500 mg/m2) |          |            |
|                       |              |       |                 |           |     | CBDDCA (AUC 5), PTX (200 mg/m2) |          |            |
| Rosell (2002)         | Spain        | III   | RR (Non-inf)    | NSCLC     | 2   | IIIb, IV, Rec | 618      | 58         |
|                       |              |       |                 |           |     | ECOG 0-2 |          |            |
|                       |              |       |                 |           |     | CBDDPA (80 mg/m2), PTX (200 mg/m2) |          |            |
|                       |              |       |                 |           |     | CBDDCA (AUC 6), PTX (200 mg/m2) |          |            |
| Sandler (2010) E4599, Updated | USA     | III   | OS              | Ad #      | 2   | IIIb, IV | 602      | 63         |
|                       |              |       |                 |           |     | ECOG 0-1 |          |            |
|                       |              |       |                 |           |     | CBDDCA (AUC 6), PTX (200 mg/m2), BEV (15 mg/kg) |          |            |
|                       |              |       |                 |           |     | CBDDCA (AUC 6), PTX (200 mg/m2) |          |            |
| Scagliotti (2002)     | Italy        | III   | NS              | NSCLC     | 3   | IIIb, IV, Rec | 612      | 63         |
|                       |              |       |                 |           |     | ECOG 0-2 |          |            |
|                       |              |       |                 |           |     | CBDDCA (AUC 6), GEM (1250 mg/m2 (d 1,8)) |          |            |
|                       |              |       |                 |           |     | CBDDCA (AUC 6), PTX (225 mg/m2) |          |            |
| Scagliotti (2008)     | Italy        | III   | OS (Non-inf)    | NSq #     | 2   | IIIb, IV | 1252     | 61         |
|                       |              |       |                 |           |     | ECOG 0-1 |          |            |
|                       |              |       |                 |           |     | CBDDPA (75 mg/m2), PEM (500 mg/m2) |          |            |
|                       |              |       |                 |           |     | CBDDPA (75 mg/m2), GEM (1250 mg/m2 (d 1,8)) |          |            |
| Schiller (2002) ECOG 1594 | USA       | NS    | OS              | NSCLC     | 4   | IIIb, IV, Rec | 1207     | 63         |
|                       |              |       |                 |           |     | ECOG 0-2 |          |            |
|                       |              |       |                 |           |     | CBDDPA (75 mg/m2), GEM (1000 mg/m2 (d 1,8,15)) |          |            |
|                       |              |       |                 |           |     | CBDDPA (75 mg/m2), DTX (75 mg/m2) |          |            |
|                       |              |       |                 |           |     | CBDDCA (AUC 6), PTX (225 mg/m2) |          |            |
| Schuette (2013)       | Germany      | II    | PFS             | NSq #     | 2   | IIIb, IV | 133      | 64         |
|                       |              |       |                 |           |     | ECOG 0-1 |          |            |
|                       |              |       |                 |           |     | CBDDPA (75 mg/m2), PEM (500 mg/m2) |          |            |
|                       |              |       |                 |           |     | CBDDCA (AUC 6), PEM (500 mg/m2) |          |            |
| Shukuya (2015) WJCOSG5208L | Japan   | III   | OS              | Sq        | 2   | IIIb, IV, Rec | 355      | 64         |
|                       |              |       |                 |           |     | ECOG 0-1 |          |            |
|                       |              |       |                 |           |     | CBDDPA (100 mg/m2), DTX (60 mg/m2) |          |            |
|                       |              |       |                 |           |     | CBDDPA (80 mg/m2), DTX (60 mg/m2) |          |            |
| Smit (2003) EORTC08975 | Netherlands | III   | OS              | NSCLC     | 3≈ 2 excluded | IIIb, IV | 319      | 57         |
|                       |              |       |                 |           |     | ECOG 0-2 |          |            |
|                       |              |       |                 |           |     | CBDDPA (80 mg/m2), PTX (175 mg/m2) |          |            |
|                       |              |       |                 |           |     | CBDDPA (80 mg/m2), GEM (1250 mg/m2 (d 1,8)) |          |            |
| Sun (2015)            | Korea        | II    | RR              | NSq       | 2   | IIIb, IV, Rec | 321      | 60         |
|                       |              |       |                 |           |     | ECOG 0-1 |          |            |
|                       |              |       |                 |           |     | CBDDPA (70 mg/m2), PEM (500 mg/m2) |          |            |
|                       |              |       |                 |           |     | CBDDPA (70 mg/m2), GEM (1000 mg/m2 (d 1,8)) |          |            |
| Tan (2009) GLOB3      | Singapore    | III   | TTF             | NSCLC     | 2   | IIIb, IV, Rec | 390      | 61         |
|                       |              |       |                 |           |     | KPS ≥ 80% |          |            |
|                       |              |       |                 |           |     | CBDDPA (80 mg/m2), VNR (30 (d1), 80 (d 8 po) mg/m2) |          |            |
|                       |              |       |                 |           |     | CBDDPA (75 mg/m2), DTX (75 mg/m2) |          |            |

Continued
secondary endpoints. The main advantage of this study over published systematic reviews on chemotherapy for NSCLC that used the conventional head-to-head meta-analysis is that we could compare a variety of chemotherapy regimens simultaneously by applying the network method (Fig. 2). In addition, the low heterogeneity (Fig. 2), the consistent results from sensitivity analyses (Supplementary Figure 1), the sound methodology following updated meta-analysis guidelines4,11, and the sufficient statistical power supported by the sufficient number of included studies and patients (Table 1) ensured the validity of the results. Although the results from this research could not recommend the single best regimen for NSCLC, we believe the current study provides useful data for the daily practice and for future chemotherapy trials.

### Table 1. Characteristics of included studies.

| Study            | Country | Phase | Primary outcome | Pathology | Arm | Stage, Performance status | Regimens | Patients | Median age |
|------------------|---------|-------|-----------------|-----------|-----|--------------------------|----------|----------|------------|
| Thomas (2006) GGFPC99-01 | France | II    | RR              | NSCLC     | 2   | IIIb, IV ECOG 0-2        | CDDP (80 mg/m2), VNR (10 mg/m2 (weekly)) CDDCA (AUC 6), GEM (1250 mg/m2 (d 1,8)) | 100      | 58        |
| Treat (2010)     | USA     | III   | OS              | NSCLC     | 3>2 | IIIb, IV Rec             | CDDCA (AUC 5.5), GEM (1000 mg/m2 (d 1,8)) | 758      | 64        |
| Wu (2014) JMIL   | China   | III   | OS              | NSq       | 2   | IIIb, IV ECOG 0-1        | CDDP (75 mg/m2), PEM (500 mg/m2) CDDP (75 mg/m2), GEM (1250 mg/m2 (d 1,8)) | 256      | 57        |
| Yang (2012)      | China   | NS    | RR              | NSCLC     | 2   | IIIb, IV ECOG 0-2        | CDDP (80 mg/m2), GEM (1250 mg/m2 (d 1,8)) CDDCA (AUC 5), GEM (1200 mg/m2 (d 1,8)) | 62       | 57        |
| Zatloukal (2003) | Czech   | III   | G3/4 toxicity   | NSCLC     | 2   | IIIb, IV KPS ≥ 70%       | CDDP (80 mg/m2), GEM (1200 mg/m2 (d 1,8)) CDDCA (AUC 5), GEM (1200 mg/m2 (d 1,8)) | 176      | 62        |
| Zhang (2013)     | China   | II    | PFS             | NSq #     | 2   | IIIb, IV Rec             | CDDP (75 mg/m2), PEM (500 mg/m2) CDDP (75 mg/m2), GEM (1000 mg/m2 (d 1,8)) | 205      | 54        |
| Zhou (2015)BEYOND| China   | III   | PFS             | NSq       | 2   | IV, Rec ECOG 0-1         | CDDCA (AUC 6), PTX (175 mg/m2), BEV (15 mg/kg) CDDCA (AUC 6), PTX (175 mg/m2) | 276      | 57        |
| Zinner (2015) PRONOUNCE | USA | III   | G4PFS           | NSq       | 2   | IV                      | CDDCA (AUC 6), PTX (200 mg/m2), BEV (15 mg/kg) CDDCA (AUC 6), PEM (500 mg/m2) | 361      | 66        |

Notes: First author, publication year, specific study name if available are presented. Updated: Updated data that were published later were available. NS: not specified. OS: overall survival. PFS: progression-free survival. QOL: quality of life. RR: response rate. DCR: disease control rate. TTP: time to progression. AE: adverse event. G3/4PFS: PFS without grade 3/4 AE. Non-inf: Primary outcome was evaluated by non-inferiority analysis. NS: not specified. A data for OS was not obtainable. Pathology: NSCLC: non-small cell lung cancer. NSq: non-squamous carcinoma. Ad: adenocarcinoma. #: The study was originally designed for NSCLC. However, we extracted data only for NSq or Ad because regimen included Pemetrexed or Bevacizumab. Arm: 3>2 (excluded), 4>2 (excluded): The original study evaluated three/four arms. However, only two arms were included for our analysis because one/two arm(s) evaluated regimen(s) out of our concern. 3>2 (combined): The original study evaluated three arms. However, we combined two arms with different dose of Bevacizumab. Stage: Rec: recurrent. Performance status: ECOG: Eastern Cooperative Oncology Group performance status. Karnofsky Performance Status. Regimens: CDDP: Cisplatin. CBDCA: Carboplatin. CDGP: Nedaplatin. DTX: Docetaxel. PTX: Paclitaxel. VNR: Vinorelbine. GEM: Gemcitabine. CPT-11: Irinotecan. PEM: Pemetrexed. S1: Tegafur gimeracil oteracil. d: day. po: oral administration. bid: twice daily. Patients: Numbers of patients randomized for evaluated arms. Median Age: When median age (years) is not available, average age (years) is presented instead.
CDGP administration root, because there was not a large difference of OS among them. We anticipate further trials of treatment choice among CDDP/CBDCA regimens is predominantly based on the adverse event profile and often selected as second-line regimens after the failure of the first-line treatment by epidermal growth factor rank order among some first-choice regimens is informative. Third, platinum doublet and triplet treatments are the very large sample size in the analysis, we could not clearly reveal the single best regimen. Nonetheless, the indirect comparisons. However, the consistency between study designs dispels such doubt. Second, despite

For the treatment of non-squamous NSCLC, CBDCA + PTX + BEV and CBDCA + PEM + BEV resulted in the best OS (Fig. 3). The main SAEs concerned were anemia and thrombopenia by CBDCA + PEM + BEV and alopecia by CBDCA + PTX + BEV. Although PEM and CDDP are known to cause severe nausea and appetite loss, thanks to the recent development of anti-emesis drugs, PEM and CDDP regimens are no longer associated with severe nausea (Fig. 3). Although PFS and RR were inferior to regimens above, CDDP + PEM is another excellent regimen showing almost equivalent OS to these BEV regimens. The adverse event profile of CDDP + PEM was also similar to that of CBDCA + PTX + BEV. In the last few decades, it has been believed that daily hydration is mandatory for CDDP administration to avoid nephrotoxicity. However, the recent development of Mg-containing short hydration enables the administration of the CDDP regimen in an outpatient setting without a large amount of hydration21. Another advantage of the CDDP + PEM regimen is being able to avoid the economically expensive BEV. Actually, these three regimens have been often selected as the first-choice regimens. CBDCA + PEM is another promising regimen despite frequent anemia and thrombopenia. This regimen showed good indications for the elderly especially those with deteriorated renal function22. CDDP + CPT11 has been one of the classical standard regimens since it was shown to be superior to CDDP + Vindesine for treatment of NSCLC23. HRos by CDDP + CPT-11 compared to CBDCA + PTX + BEV was 1.16 (95%CI 0.90–1.50). This does not show significance; however, this may suggest that CDDP + CPT-11 is related to poorer survival. In addition, frequent severe diarrhea and anemia made it questionable to regard this regimen as the first choice. Imprecise estimation for OS by CDGP + GEM made the survival benefits of this regimen inconclusive. Given the non-promising results from the phase II study, we do not anticipate a phase III trial for CDGP + GEM. In the last few decades, it has been believed that daily hydration is mandatory for CDDP administration to avoid nephrotoxicity. However, the recent development of Mg-containing short hydration enables the administration of the CDDP regimen in an outpatient setting without a large amount of hydration21. Another advantage of the CDDP + PEM regimen is being able to avoid the economically expensive BEV. Actually, these three regimens have been often selected as the first-choice regimens. CBDCA + PEM is another promising regimen despite frequent anemia and thrombopenia. This regimen showed good indications for the elderly especially those with deteriorated renal function22. CDDP + CPT11 has been one of the classical standard regimens since it was shown to be superior to CDDP + Vindesine for treatment of NSCLC23. 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This does not show significance; however, this may suggest that CDDP + CPT-11 is related to poorer survival. In addition, frequent severe diarrhea and anemia made it questionable to regard this regimen as the first choice. Imprecise estimation for OS by CDGP + GEM made the survival benefits of this regimen inconclusive. Given the non-promising results from the phase II study, we do not anticipate a phase III trial for CDGP + GEM. In the last few decades, it has been believed that daily hydration is mandatory for CDDP administration to avoid nephrotoxicity. However, the recent development of Mg-containing short hydration enables the administration of the CDDP regimen in an outpatient setting without a large amount of hydration21. Another advantage of the CDDP + PEM regimen is being able to avoid the economically expensive BEV. Actually, these three regimens have been often selected as the first-choice regimens. 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receptor-tyrosine kinase inhibitors, anaplastic lymphoma kinase inhibitors, and immune check-point inhibitors. Our study does not directly provide data for second-line platinum regimens. Fourth, most of the evaluated original trials had a high risk of bias judged from the Cochrane tool. Unfortunately, in practical terms, it is very difficult to conduct a double-blinded trial without sponsorship from pharmaceutical companies and we believe that these factors do not largely flaw the credibility of our analysis.

In conclusion, we conducted a systematic review and network meta-analysis. Based on 16842 NSCLC patients constituting 48 RCTs, CBDCA + PTX + BEV, CBDCA + PEM + BEV, CDDP + PEM, CBDCA + PEM, and

Figure 3. Forest plots for primary and secondary outcomes.
CDDP + CPT-11 seemed reasonable first-choice regimens for non-squamous NSCLC. Even though other platinum regimens are also recommended in the guidelines, the results from our analysis do not support regular use of these regimens. CDP + DTX and some CDDP/CBDCA regimens seemed acceptable first-choice regimens for squamous NSCLC.

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Author Contributions
H.N. contributed for conception, data extraction, analysis, and drafting, N.A. worked for data extraction and drafting, N.K. and S.Y. interpreted the results. I.K. critically revised the protocol and main manuscript. G.A. and Y.T. provided statistical advices. K.T. conceived the study and critically revised the manuscript.

Additional Information
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