

(2) Phone entry (emergency only)

In the event that the web-based registration system is not available due to system failure or it is not possible to use the system during the surgery for any reason, the iPocc Trial Coordinating Center will complete the patient registration process on behalf of the physician. Please follow the procedure below.

<Before the surgery date>

After tentative registration has been completed, inform the iPocc Trial Coordinating Center of the scheduled date and time of surgery via email 2 business days before surgery starts.

<During the surgery>

- 1) Upon completion of the comprehensive staging surgery, but immediately before the abdomen or abdominal wall is closed, the investigator or his/her designee will call the iPocc Trial Coordinating Center and notify the center of the institution name, the name of the investigator, the patient's birth date, the tentative registration number, and other stratification factors (the size of the residual tumor and the FIGO stage).
- 2) Emergency registration by phone will be available on weekdays from 9:00 am to 5:00 pm Japan time. Prior notification to the iPocc Trial Coordinating Center is necessary if final registration/randomization is likely to be after 5:00 pm,.
- 3) The iPocc Trial Coordinating Center, on behalf of the investigator, will enter stratification factors and perform randomization via the Rave system. Randomization will take approximately 3-5 minutes.
- 4) The investigator or his/her designee will repeat the assigned regimen on the phone and confirm it with other attending medical staff at the site.
- 5) The IP port system will be placed only when the patient is assigned to regimen II (dd-TCip therapy).

[For emergency final registration]

iPocc Trial Coordinating Center

Kitasato University Research Center for Clinical Pharmacology,
Clinical Trial Coordinating Center

TEL: +81-3-5791-6419 or 6398

Emergency final registration time:

9:00 am to 5:00 pm Japan time on weekdays, in general

*No Emergency final registration will be accepted on Saturdays,
Sundays or public, and holidays.

[Japan only as a general]

6.3.2.4 Notes for final registration/randomization

- 1) An email with the final registration/randomization result will be automatically sent to the investigator who had completed the final registration/randomization.
- 2) The investigator should confirm the “Protocol patient ID (final registration number)” and assigned regimen in the Rave system.
- 3) The assignment regimen should be recorded in the patient’s medical record.
- 4) Once the final registration is completed, it is impossible to cancel this from the database. In case of a duplicate registration, the initial final registration data and the Protocol patient ID (final registration number) will be kept and used for the study analysis.
- 5) When a false registration or a duplicate registration is found, the investigator should promptly notify the iPocc Trial Coordinating Center.

6.3.3 Notes for final registration (For institutions other than those in Japan)

For institutions that prefer to obtain randomization results after histopathological confirmation for epithelial ovarian cancer and staging, please follow the subsequent procedure. Tentative registration is exactly the same as those given in the Japanese guidelines. 【See section 6.3.1】

Final registration can be done after histopathological diagnosis has been made. In this case, the registration procedure is the same as that described in Section 6.3.2.2.

Placement of the IP port can be performed after the patient has been assigned to regimen II. IP port can be placed during surgery for all patients, and can be removed, when the patient is assigned to regimen II.

Please consult your institutional review committee regarding port insertion procedure that will be applied.

7 TREATMENT PLAN AND TREATMENT MODIFICATION CRITERIA

Protocol treatment and treatment modification will be implemented according to the protocol, as long as patient safety is not jeopardized. In the event that compliance with the protocol is not medically appropriate for the patient, the investigator may modify the protocol treatment using their medical judgment. This would be classed as a “protocol deviation” but may be considered to be a “clinically reasonable deviation.”

7.1 Protocol treatment

Patients will be assigned to either of the following two regimens. The investigator should start the assigned protocol regimen within 8 weeks after the comprehensive staging surgery.

Regimen I (Standard treatment: dd-TCiv therapy)

Paclitaxel:	80 mg/m ²	1 hour IV infusion	Days 1, 8, and 15
Carboplatin:	AUC = 6.0	1 hour IV infusion	Day 1

The 3-week period (21 days) is 1 cycle. A total of 6 to 8 cycles will be repeated.*)

*) Protocol treatment comprises 6 cycles. However, if interval debulking surgery (IDS) is performed after 3, 4 or 5 cycles, the patients can receive up to 3 additional cycles of the protocol treatment.

Regimen II (Study treatment: dd-TCip therapy)

Paclitaxel:	80 mg/m ²	1 hour IV infusion	Days 1, 8, and 15
Carboplatin:	AUC = 6.0	IP injection	Day 1

The 3-week period (21 days) is 1 cycle. A total of 6 to 8 cycles will be repeated. *)

*) Protocol treatment comprises 6 cycles. However, if interval debulking surgery (IDS) is performed after 3, 4 or 5 cycles, the patients can receive up to 3 additional cycles of the protocol treatment.

7.2 Dose calculation

A dose calculation tool is posted on the STUDY WEB-PAGE.

7.2.1 Calculation of body surface area (BSA)

1) The DuBois formula will be used to determine BSA, which will be used for dose calculation.

DuBois Formula:

$$\text{BSA} = \text{Body Weight}^{0.425} \times \text{Height}^{0.725} \times 71.84 / 10000$$

where BSA is in m², Body Weight is in kg and Height is in cm.

2) In patients whose BSA is greater than 2.0 m², a BSA of 2.0 m² will be used.

7.2.2 Dose calculation for paclitaxel

The dose of paclitaxel will be calculated using the BSA determined by the DuBois formula.

7.2.3 Dose calculation for carboplatin

- 1) The dose of carboplatin will be calculated using the following Calvert formula ²⁶⁾.

Calvert Formula:

$$\text{Carboplatin dose (mg/body)} = \text{target AUC} \times (\text{GFR} + 25)$$

- 2) In this study, the GFR is considered to be equivalent to the creatinine clearance (Ccr).
- 3) The Ccr is calculated using the following modified-Jelliffe formula ²⁷⁾.

Modified-Jelliffe Formula:

$$\text{CCr} = [98 - \{0.8 \times (\text{age} - 20)\}] \times \text{BSA} \times 0.9 / (\text{serum creatinine} \times 1.73)$$

where Ccr is in mL/min, age is in years, BSA is in m², and serum creatinine is in mg/dL.

When expressed in SI units, serum creatinine should be converted to mg/dL using the following equation:

$\text{Serum creatinine (mg/dL)} = \text{SI unit } (\mu\text{mol/L}) / 88.4$
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- 4) In patients whose serum creatinine is less than 0.6 mg/dL, a serum creatinine value of 0.6 mg/dL will be used for the calculation.
- 5) Dose recalculation for carboplatin is not necessary unless the patient experiences a newly diagnosed urinary tract obstruction or CTCAE renal dysfunction of at least grade 2 in severity (serum creatinine greater than 1.5 times the institution's upper limit of normal).
- 6) The maximum dose of carboplatin will not exceed 1000 mg /person.

7.3 Regimen I (Standard treatment: dd–TCiv therapy)

Paclitaxel:	80 mg/m ²	1 hour IV infusion	Days 1, 8, and 15
Carboplatin:	AUC = 6.0	1 hour IV infusion	Day 1

The 3-week period (21 days) is 1 cycle. A total of 6 to 8 cycles will be repeated.*)

*) Protocol treatment comprises 6 cycles. However, if interval debulking surgery (IDS) is performed after 3, 4 or 5 cycles, the patients can receive up to 3 additional cycles of the protocol treatment.

7.3.1 Administration for paclitaxel

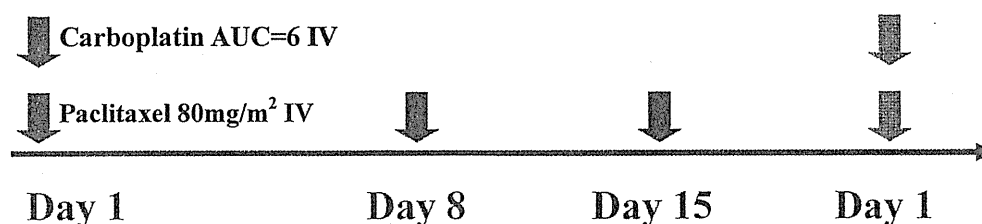
- 1) After the introduction of premedication, the prescribed dose of paclitaxel diluted in 250 mL of 5% dextrose or normal saline solution (NSS) and will be administered by intravenous infusion over 1 hour through an exclusive line.
- 2) Paclitaxel will be given on Days 1, 8 and 15 for each cycle.

7.3.2 Dosing regimen for carboplatin

- 1) Following the administration of paclitaxel, the prescribed dose will be diluted in 250 mL of 5% dextrose or NSS and administered by IV over 1 hour.
- 2) Carboplatin will be given on Day 1 for each cycle.

7.3.3 Example of Regimen I (Standard treatment: dd-TCiv therapy)

<u>Day 1</u>	
9:00 AM	Premedication 【See section 7.8.4】
9:30 AM	NSS (250 mL) + Paclitaxel (80 mg/m ²) IV infusion (approximately 60 minutes)
10:30 AM	NSS (250 mL) + Carboplatin (AUC=6) IV infusion (over 30 minutes)
<u>Day 8</u>	
9:00 AM	Premedication
9:30 AM	NSS (250 mL) + Paclitaxel (80 mg/m ²) IV infusion (approximately 60 minutes)
<u>Day 15</u>	
9:00 AM	Premedication
9:30 AM	NSS (250 mL) + Paclitaxel (80 mg/m ²) IV infusion (approximately 60 minutes)



7.4 Regimen II (Study treatment: dd-TCip therapy)

Paclitaxel:	80 mg/m ²	1 hour IV infusion	Days 1, 8, and 15
Carboplatin:	AUC = 6.0	single IP injection	Day 1

The 3-week period (21 days) is 1 cycle. A total of 6 to 8 cycles will be repeated.*)

*) Protocol treatment comprises 6 cycles. However, if interval debulking surgery (IDS) is performed after 3, 4 or 5 cycles, the patients can receive up to 3 additional cycles of the protocol treatment.

7.4.1 Administration of paclitaxel

- 1) After the introduction of premedication, the prescribed dose of paclitaxel diluted in 250 mL of 5% dextrose or normal saline solution (NSS) will be administered by intravenous infusion over 1 hour through an exclusive line.
- 2) Paclitaxel will be given on Days 1, 8 and 15 for each cycle.

7.4.2 Administration of carboplatin

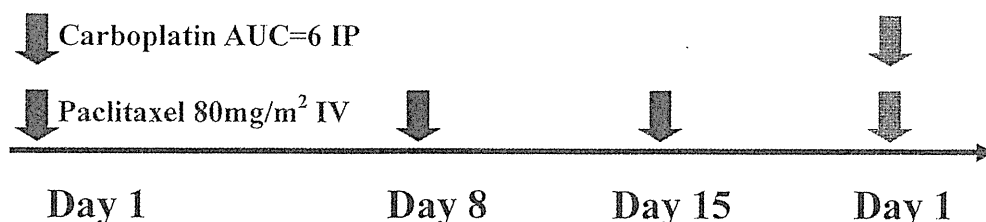
- 1) During IV administration of paclitaxel, the IP port will be punctured using the Huber needle, and 1000 mL to 1500 mL of normal saline solution will be intraperitoneally injected.
- 2) After completion of paclitaxel IV administration, the prescribed dose of carboplatin prepared in a syringe will be given as a single injection through the Huber needle.
- 3) Following injection of the total dose of carboplatin, 10 mL of heparinized normal saline solution will be injected, and the Huber needle will be removed.
- 4) Carboplatin will be given on Day 1 for each cycle.

7.4.3 Notes for IP administration (For details, [see Appendix1-A])

- 1) Pay careful attention to prevent infection. IP-related infection may possibly cause peritonitis.
- 2) Port puncture should be performed aseptically using surgical gloves.
- 3) Completely sterilize the skin at the puncture site with Isodine solution.
- 4) When carboplatin is given as a single injection, it is recommended to inject approximately 10 mL first, and inject the rest after observation to check for any allergic reactions.
- 5) Remove the IP port immediately if peritoneal infection is suspected.
- 6) Discontinue IP administration if catheter obstruction is suspected.
- 7) In patients with massive ascites, it is not always necessary to intraperitoneally administer 1000 mL to 1500 mL of normal saline solution. Based on the medical judgment of the attending physician, an adequate amount of normal saline solution should be administered. The attending physician should confirm that there is no difficulty associated with injection of carboplatin.
- 8) If there is a problem in IP administration of normal saline solution, catheter obstruction or bowel adhesion to the area surrounding the catheter may be suspected. In such a case, the investigator should discontinue IP administration and notify the iPocc Trial Coordinating Center.
- 9) If any allergic reaction to carboplatin is observed, stop carboplatin injection, and please inform the iPocc Trial Coordinating Center. 【See Appendix3-V】 Switching carboplatin (IP) to cisplatin (IP) is not generally allowed for this study.

7.4.4 Example of Regimen II (Study treatment: dd-TCip therapy)

<u>Day 1</u>	
9:00 AM	Premedication
9:30 AM	NSS (250 mL) + paclitaxel (80 mg/m ²) IV infusion (60 minutes) IP Port puncture, and injection NSS(1000mL to 1500mL) IP injection(full open)
10:30 AM	Completion of paclitaxel infusion Carboplatin (AUC = 6) Single IP injection Heparinized NSS (10 mL) IP injection, Removal of the Huber needle
<u>Day 8</u>	
9:00 AM	Premedication
9:30 AM	NSS (250mL) + paclitaxel (80mg/m ²) IV infusion (60 minutes)
<u>Day 15</u>	
9:00 AM	Premedication
9:30 AM	NSS (250mL) + paclitaxel (80mg/m ²) IV infusion (60 minutes)



7.5 Criteria for starting protocol treatment [For both Regimen I and II]

7.5.1 Criteria for starting protocol treatment on Cycle 1

7.5.1.1 Criteria for starting Day 1 of Cycle 1

After surgery, the investigator needs to confirm that the patient meets all clinical laboratory data criteria. [See the section 4.1.] The laboratory data referred to for each patient should be that obtained within 2 weeks before the first date of protocol treatment.

7.5.1.2 Criteria for starting Day 8 and Day 15 of Cycle 1

- 1) Within 2 days before starting administration of protocol treatment, the patient meets all criteria described in Table 1.
- 2) If the patient fails to meet all criteria in Table 1, treatment can be delayed by up to 3 weeks.
- 3) In case of failure to meet the criteria even after a 3-week delay, the protocol treatment should be discontinued. (However, if the same treatment as the protocol-directed treatment is to be given after more than 3 week delay, we request the treatment to be continued as close as possible to the protocol-directed treatment. Ssubmission of eCRFs is also necessary for such a case., see the section 7.7.2.)

Table 1. Criteria for starting paclitaxel administration on Day 8 and Day 15

Item	Criteria for starting
Absolute neutrophil count	$\geq 500 /\text{mm}^3$
Platelet count	$\geq 50,000 /\text{mm}^3$

7.5.2 Criteria for starting successive cycles

7.5.2.1 Criteria for starting administration on Day 1 of successive cycles

- 1) Within 2 days before starting administration of the protocol treatment, the patient meets all criteria described in Table 2.
- 2) If the patient fails to meet all the criteria in Table 2, treatment can be delayed by up to 3 weeks.
- 3) In case of failure to meet the criteria even after a 3-week delay, the protocol treatment should be discontinued. (However, if the same treatment as the protocol-directed treatment is to be given after more than 3 week delay, we request the treatment to be continued as close as possible to the protocol-directed treatment. Ssubmission of eCRFs is also necessary for such a case., see the section 7.7.2.)

Table 2. Criteria for starting administration on Day 1 after cycle 2

Item	Criteria for starting
Absolute neutrophil count	$\geq 1,000 /\text{mm}^3$
Platelet count	$\geq 75,000 /\text{mm}^3$
Peripheral neuropathy	Grade 1 or 0
Other non-hematologic toxicities (except hair loss, fatigue, nausea, constipation)	Grade 1 or 0

7.5.2.2 Criteria for starting administration on Day 8 and Day 15 of successive cycles

Same as section 7.5.1.2. criteria for starting administration Day 8 and Day 15 of cycle 1

7.6 Dose reduction criteria [For both Regimen I and II]

7.6.1 Dose reduction process

- 1) If patients meet any one of the reduction criteria shown in Table 3 during the previous cycle, the dose will be reduced by one level in subsequent cycles according to the dose reduction levels shown in Table 4.
- 2) Even if patients meet two or more of the reduction criteria, the dose will be reduced by only one level in the subsequent cycle. (Do not reduce the dose by two levels at once.)
- 3) If patients meet the criteria again after reducing the dose, the dose will be reduced again by a further one level in the subsequent cycle according to the dose reduction levels shown in Table 4.
- 4) Up to two dose reductions are allowed. Protocol treatment in subsequent cycles will be discontinued if patients meet the criteria after the second reduction. (In the event that the same treatment as in the protocol treatment is continued after discontinuation of the protocol treatment, see section 7.7.2.)
- 5) Once the dose has been reduced, it should not be increased for subsequent cycles.
- 6) If the attending physician determines that it is appropriate to modify the treatment for other reasons, notification should be provided to the iPocc Trial Coordinating Center.

Table 3. Dose reduction criteria

Reduction criteria	Drug to be reduced
Cycle delayed more than 2 weeks but for no more than 3 weeks because the patient did not meet the criteria for starting treatment described in Table 2	
Dose-limiting neutropenia (DLT-ANC)* or Dose-limiting thrombocytopenia (DLT-PLT)** during the previous cycle	The doses of both paclitaxel and carboplatin are reduced by -1 level .
Grade 3 non-hematological toxicities (Except for alopecia, fatigue, nausea, constipation, or peripheral neuropathy)	
Grade 2 or greater peripheral neuropathy observed in the previous cycle.	The dose of paclitaxel alone is reduced by -1 level.

* Neutropenia (DLT-ANC):

- ① Febrile neutropenia: neutrophil count decreased associated with fever as specified in CTCAEv4.0.
- ② Prolonged grade 4 neutropenia (absolute neutrophil count less than 500/mm³) for at least 7 days despite the use of G-CSFs. Neutropenia without complication lasting for less than 7 days is not DLT-ANC.

** Thrombocytopenia (DLT-PLT):

- ① Grade 4 thrombocytopenia (<25,000/mm³).
- ② Thrombocytopenia with a bleeding tendency (2,500/mm³ to <50,000/mm³) is observed, and/or platelet transfusion is required.

Table4. Dose modification for toxicity

Level	Paclitaxel (mg/m ²)	Carboplatin (AUC)
0	80	6.0
-1	70	5.0
-2	60	4.0
-3	Discontinued	Discontinued

7.6.2 Management of hypersensitivity reactions to paclitaxel [see Appendix 3-VI]

- 1) Hypersensitivity reactions to paclitaxel are NOT considered to be a dose-limiting toxicity.
- 2) It may be possible to administer the total dose of paclitaxel after treating hypersensitivity reactions.
- 3) When a patient is re-challenged after the occurrence of hypersensitivity reactions to paclitaxel, slower administration of paclitaxel should be performed for subsequent doses.

7.6.3 Recalculation of dose

7.6.3.1 Dose recalculation for renal dysfunction

Considering the dosage and regimen in this protocol, renal dysfunction is usually not directly associated with chemotherapy-induced adverse effects. However, if serum creatinine is greater than 1.5 times the institution's upper limit of normal, the dose of carboplatin should be recalculated for each cycle.

7.6.3.2 Dose recalculation for weight gain/loss

If patient weight change is less than 10% compared with the weight at the start of protocol treatment, recalculation of the dose for the subsequent cycle will not be required. In the event of weight gain/loss of more than or equal to 10%, the dose will be recalculated.

7.7 Criteria for protocol treatment completion/discontinuation

[For both Regimen I and II]

7.7.1 Criteria for protocol treatment completion

- 1) Protocol treatment should be completed at 6 cycles.
- 2) If interval debulking surgery (IDS) is performed, the patient can receive up to 3 additional cycles of the protocol treatment after IDS.
- 3) Treatment after completion of the protocol treatment is not specified. However, the details of such treatment (dose, dosing regimen, duration of administration, etc.) will be reported on the follow-up form.

7.7.2 Criteria for protocol treatment discontinuation

- 1) Discontinuation of protocol treatment due to adverse events.
 - ① Greater than 3-week delay^{*)}
 - ② Grade 4 non-hematologic toxicity^{**)}

- ③ The third dose reduction (level -3)*)
- 2) Discontinuation of protocol treatment due to a patient request.
- 3) Patient death during protocol treatment.
- 4) Development of disease progression after the starting the protocol treatment.
- 5) When the investigator determines that the protocol treatment is no longer appropriate for the patient
(including cases where treatment is switched from dd-TCip therapy to dd-TCiv therapy).
In such a case, notification should be provided to the iPocc Trial Coordinating Center.

*) In cases where the same treatment as the protocol treatment is continued, even after more than a 3-week delay and/or at a smaller dosage than the protocol-directed treatment, patients may, in some cases, not be excluded from the study. Therefore, it is necessary to continue the treatment according to the test/observation schedule specified in the protocol, and submit the case report form likewise.

**) If the attending physician considers that it is appropriate to continue the protocol treatment after grade 4 non-hematologic toxicities, the iPocc Trial Coordinating Center should be notified in advance to discuss this with the study chair.

7.8 Concurrent/supportive therapy [For both Regimen I and II]

7.8.1 G-CSF

- 1) No prophylactic administration of G-CSF, such as filgrastim or lenograstim, PEG-filgrastim (Neulasta), or sargramostim (GM-CSF) is allowed.
- 2) The ASCO Guidelines indicate the following [see Appendix 3-VII]:
 - (1) Conditions for G-CSF use
 - ① ANC is less than 1000/mm³ (grade3) and a fever (38°C/100.4°F or higher) is observed.
 - ② ANC is less than 500/mm³ (grade4).
 - ③ Either of the above conditions ① or ② were observed in the previous cycle with an ANC of less than 1000/mm³ in current cycle.
 - (2) Dosage and administration
A dose of 50 µg/m² will be administered daily by subcutaneous injection.
 - (3) Conditions for G-CSF discontinuation
 - ① When patients have an ANC of greater than or equal to 5000 /mm³ after daily injection of G-CSF.
 - ② When patients have an ANC of greater than or equal to 2000/mm³ without any symptoms suggestive of infection and with no anticipated additional risk to the patients due to good response to previous G-CSF administration, at the discretion of the investigator.

7.8.2 Treatment for anemia

Patients may receive erythropoietin (EPO), iron supplements, and/or blood transfusions as clinically indicated for management of anemia.

7.8.3 Antiemetic drugs

Prophylactic administration of a 5-HT₃ antagonist or other antiemetic drugs for reducing nausea and vomiting is recommended.

7.8.4 Antiallergic drugs

Prophylactic administration of a steroid or antihistamine agent for preventing allergic reaction to paclitaxel is recommended.

7.8.5 Premedications: an example

For all cycles when paclitaxel is administered, an appropriate premedication regimen should be employed.

- 1) 30 minutes prior to paclitaxel administration, patients will receive the following short premedication to reduce the risk of hypersensitivity reactions. This regimen should include dexamethasone 20 mg intravenously or orally, anti-histamine H1 (such as diphenhydramine 50 mg orally), anti-histamine H1 (such as ranitidine hydrochloride 50 mg intravenously²⁸).
- 2) If patients have no allergic reactions during cycle 1, the dose of dexamethasone may be gradually decreased to 20 mg, 8 mg, 4 mg, 2 mg and 1 mg for subsequent cycles.

7.8.6 Acceptable concurrent/supportive therapy and surgery

- 1) If concurrent use of drugs (antimicrobial agents, analgesic, blood transfusion, etc.) for treating complications or adverse events is necessary, such drugs can be used at the discretion of the investigator.
- 2) Various symptomatic therapies for treating peripheral neuropathy caused by paclitaxel can be used at the discretion of the investigator.
- 3) IDS is allowed to be performed after 3, 4 or 5 cycles of the protocol treatment. In such cases, the protocol treatment must be restarted within 8 weeks after IDS. If IDS is performed, patients can receive up to 3 additional cycles of the protocol treatment after IDS.

7.8.7 Unacceptable concurrent/supportive therapy and surgery

- 1) All anticancer treatments other than the protocol treatment with paclitaxel and carboplatin will not be accepted. Those include all cancer drugs except paclitaxel and carboplatin, immunotherapy, radiation therapy, etc.
- 2) Administration of any drugs undergoing clinical trials will not be accepted. (An exception may be granted after a diagnosis of progressive disease. For such a case, the iPocc Trial Coordinating Center should be notified in advance to discuss this with the study chair.)
- 3) Second-look surgery must not be carried out.

7.8.8 Post-protocol treatment after completion/discontinuation of the protocol treatment

No further anticancer therapy should be given after completion of 6-9 cycles of the protocol treatment until disease progression is documented.

Anticancer agents other than this protocol-directed therapy, radiation therapy, or immunotherapy must not be administered unless the protocol treatment is discontinued before completing the protocol-directed treatment.

Once the patient stops receiving protocol-directed therapy, post-protocol treatment is not specified. When post-treatment is given, treatment drugs, the regimen, duration of administration, etc. must be reported on the follow-up form.

8 OBSERVATIONS AND TEST SCHEDULE

8.1 The following surveys and examinations must be conducted in accordance with the specified schedule, and the evaluation forms must be filled out.

Parameters	Prior to tentative-registration	After surgery - prior to the start of therapy	During protocol therapy				At completion of therapy or withdrawal	After therapy - during follow-up period
			Every week	Prior to each cycle	Every 2 cycles	Every 3 cycles		
Explanation of the study	•							
Signed informed consent	•							
Background data	•							
Surgery and pathological findings		•						
Medical history (Past, Current)	•							
Physical examination and PS (ECOG)	•	•2)		•			•	•13)
CBC, Differential, Platelets	•1)	•2)	•4)5)	•5)			•	
Electrolytes (Na, K, Cl, Ca), BUN, serum creatinine, CA125 and other tumor markers (creatinine clearance, if necessary)	•1)	•2)		•5)			•	
Total bilirubin, ALT(GPT), AST(GOT), ALP, LDH	•1)	•2)		•5)			•10)	
Chest X-ray	•1)	•2)			•7)		•11)	
Electrocardiogram (ECG)	•1)	•2)					•11)	
Radiographic Tumor measurements		•3)			•8)		•12)	•13)
Adverse events		•2)		•			•9)	•9)
Serious adverse event reporting			•6)					
QOL assessment		•14)				•14)	•14)	•14)
Cost assessment		•15)					•15)	•15)

Notes:

1. To be conducted within 28 days prior to scheduled surgery.
2. To be conducted within 14 days prior to the start of protocol treatment after comprehensive staging surgery.
3. In the event that residual tumor(s) are found at the time of surgery, abdominal/pelvic CT or MRI must be performed within 28 days prior to the start of protocol therapy after surgery. If the use of contrast agents is contraindicated, plain CT or MRI must be performed. When IDS is conducted, abdominal/pelvic CT or MRI must be performed again within 28 days prior to the start of protocol treatment after the IDS.

4. To be conducted at least once a week.
5. To be conducted within 2 days prior to the next cycle of protocol treatment.
6. In the event that adverse events to be reported occur, prompt reporting is required. 【See Appendix 2-⑤】
7. If an abnormal condition is detected at the time of tentative registration, or if it is necessary to determine the effect of therapy, plain chest X-P must be performed repeatedly. It is not mandatory in other cases.
8. With or without measurable lesions, CT or MRI must be performed every 2 cycles (after 2 cycles, after 4 cycles, after 6 cycles, (after 8 cycles), and at the time of discontinuation of the protocol treatment). If the use of contrast agents is contraindicated, plain CT or MRI must be performed. In order to evaluate the effect of therapy, the same test method must be used for every evaluation, and RECIST (version 1.1) will be used for evaluation.
9. The final evaluation of adverse events must be conducted within 3 weeks after the final cycle of protocol treatment. Unless the protocol treatment-related adverse reactions are of at least grade 2 in severity at the time of this evaluation, follow-up observations of adverse events are not required. If adverse reactions with a grade of at least 2 are reported, evaluations must be conducted at each follow-up until such events resolve.
10. To be conducted within 7 days after the discontinuation of protocol treatment or within 7 days after the final administration in the last cycle.
11. To be conducted if necessary.
12. CT or MRI must be performed within 28 days after the discontinuation or completion of protocol treatment. However, this is not required when the discontinuation of protocol treatment is due to tumor progression.
13. Follow-ups must be conducted after the discontinuation or completion of protocol treatment as follows: 3-month intervals for the first 2 years, 6-month intervals for the following 2 years, and once a year thereafter. Follow-ups must include the evaluation of general clinical condition and CA-125 test as a minimum standard. Additionally, CT or MRI must be performed at times of suspected progression of the disease. Investigators may shorten the intervals of follow-ups and of CT or MRI if necessary.
14. The QOL survey【See Appendix 2-⑥】must be conducted as follows: after obtaining informed consent and prior to the start of protocol treatment, after 3 cycles of protocol treatment (or after 9 weeks from the start of treatment), after 6 cycles of protocol treatment (or after 18 weeks from the start of treatment), after 36 weeks from the start of protocol treatment, after 60 weeks from the start of protocol treatment, and after 84 weeks from the start of protocol treatment.
15. Survey forms in relation to the patient's expenses 【See Appendix 2-⑦】if possible with copies of receipts are to be mailed to the iPocc Trial Coordinating Center. The evaluation must be conducted by using the following three types of survey forms: ① to be used once “after the obtaining of informed consent and prior to the start of protocol treatment,” ② to be used 5 times simultaneously with the QOL evaluations, and ③ to be used “during the follow-up period after 84 weeks from the start of protocol treatment” after the completion of QOL evaluations.

9 EVALUATION CRITERIA

9.1 Adverse events

9.1.1 Definitions of adverse events (AE), adverse reactions (AR), and unexpected adverse reactions (UAR)

An adverse event (AE) is any unfavorable or unintended sign (including an abnormal laboratory finding), symptom, or disease that occurs in a patient receiving an investigational drug, whether or not it is considered related to this investigational drug. An adverse reaction (AR) is any unfavorable and unintended reaction that occurs at any dose level. An unexpected adverse reaction (UAR) is an adverse reaction (AR) that is of a nature or severity not consistent with the applicable product information.

9.1.2 Evaluation of AE and AR

AE and AR are to be evaluated using the NCI Common Terminology Criteria for Adverse Events (NCI-CTCAE) version 4.0.

AE and AR that occur or worsen during this study are to be recorded on case report forms with the details of the AE, date of occurrence, grade and relationship between the event and the investigational drug.

The investigator will assess the causal relationship of the event to the study drug in accordance with the following classification: “Reasonable possibility of a relationship to the study drug” is either Yes or No. If there is SOME evidence to suggest a causal relationship, it will be recorded as “Yes” on the CRF. When there is LITTLE evidence of any causal relationship and there may have been other factors that contributed to the AE, it will be recorded as “No” on the CRF. “A reasonable possibility of the relationship to the study drug” will include any AE that is deemed definitely, probably, possibly or unlikely. “No reasonable possibility of a relationship to the study drug” will include any AE that is deemed unrelated.

All AEs will be documented in the patient’s medical records and CRF. All AEs must be followed until resolution, or for at least 30 days after discontinuation of study treatment, whichever comes first or until toxicity has resolved to baseline, or until the toxicity is considered to be irreversible. Perceived lack of efficacy is not an AE. An exacerbation of a pre-existing condition is an AE.

Abnormal laboratory test results that are deemed clinically significant by the investigator and that lead to a change in the dosage of the study treatment or temporary or permanent discontinuation of the study treatment in the CRF.

9.1.3 Expected AEs, and actions to be taken

Please refer to the latest version of the Package Insert of each drug for expected adverse events related the study drugs used in this trial. Some of the expected AEs and the actions to be taken are listed below.

9.1.3.1 Hematotoxicity

Leukopenia, neutropenia, erythropenia, and thrombocytopenia may occur during the study treatment period. In severe cases, G-CSF and blood component transfusion (erythrocytes and thrombocytes) are to be administered. The necessity of blood component transfusion will be

determined by the attending physician/sub-attending physician. The frequency of administration and dose per administration shall be reported on case report forms.

【See the section 7.8】

9.1.3.2 Non-hematotoxicity

1 Anaphylactic reaction

Because the occurrence of anaphylactic reactions has been reported, sufficient medical history of the patient should be taken and the instructions for premedication described in section 7.8 should be followed. If an anaphylactic reaction occurs, the proper treatment should be promptly offered. If a paclitaxel-induced anaphylactic reaction occurs, it is recommended that extended premedication be provided for the following courses. If repeated anaphylactic reactions occur, discontinuation of the protocol therapy should be considered. If a carboplatin-induced anaphylactic reaction occurs, the discontinuation of protocol treatment should be considered.

2 Peripheral neuropathy

Peripheral neuropathy such as numbness in the extremities has been reported. In severe cases, dose reduction or discontinuation of study drug for following cycles is to be considered.

3 Myalgia and arthralgia

The symptoms of myalgia and arthralgia usually appear within 2 to 3 days after the administration of paclitaxel followed by a remission in a few days. In case of strong pain, the use of analgesic agents may be considered. In severe cases, dose reduction or discontinuation of the investigational drug for the following cycles is to be considered.

4 Abdominal pain

Abdominal pain while receiving an intraperitoneal injection of carboplatin has been reported. In severe cases, dose reduction or discontinuation of the investigational drug for the following cycles is to be considered.

5 Cardiovascular symptoms

Arrhythmia or asymptomatic bradycardia may occur. ECG monitoring should be performed before initial administration of the protocol treatment.

6 Gastrointestinal symptoms

Nausea, vomiting, and stomatitis may occur. A decision regarding the use of antiemetic agents will be made by the attending physician 【See section 7.8.2.】. Dose reduction or discontinuation of the study drug shall not be undertaken due to nausea and constipation.

7 Skin

Alopecia, skin rash, and skin flushing(紅潮) may occur. If any of these symptoms occur, the appropriate treatment is to be given by the attending physician.

8 Fever

If fever occurs, a prompt investigation of the cause and the proper treatment should be given. Particularly where febrile neutropenia is suspected, strict management according to guidelines is required.【See section 7.6 and 7.8】

9 Infections

If infection is suspected, the appropriate administration of drugs, such as antibiotics and antimycotics are to be given promptly. The patient's general status should be closely observed.

9.2 Tumor response evaluation

Patients participating in this trial include those who clinically have no residual tumors and those who have residual tumors to be measured by physical examinations and/or radiographic imaging.

Assessment of residual tumors must be performed within 28 days before the start of protocol treatment after surgery.

Responses in patients with measurable lesions identified by radiographic imaging performed right before starting tumor of the protocol treatment will be evaluated as below. The RECIST criteria (version 1.1) are to be used for response evaluation [See Appendix 3-IV]. Patients having only non-target lesions are also to be reported on case report forms.

9.2.1 Disease Parameters

Measurable disease:

Measurable lesions are defined as those that can be accurately measured in at least one dimension (longest diameter to be recorded) as >10 mm by CT scan, as >20 mm by chest x-ray, or >10 mm with calipers on clinical examination. All tumor measurements must be recorded in decimal fractions of centimeters.

Malignant lymph nodes:

To be considered pathologically enlarged and measurable, a lymph node must be >15 mm on the short axis when assessed by CT scan (CT scan slice thickness recommended to be no greater than 5 mm). At baseline and in follow-up, only the short axis will be measured and followed.

Non-measurable disease:

All other lesions (or sites of disease), including small lesions (longest diameter <10 mm or pathological lymph nodes with ≥ 10 to <15 mm short axis), are considered non-measurable disease. Leptomeningeal disease, ascites, pleural/pericardial effusions, lymphangitis cutis/pulmonitis, inflammatory breast disease, and abdominal/pelvic masses (identified by physical exam and not CT or MRI), are considered as non-measurable.

Bone lesions:

Lytic bone lesions or mixed lytic-blastic lesions, with identifiable soft tissue components, that can be evaluated by CT or MRI can be considered as measurable lesions if the soft tissue component meets the definition of measurability described above. Blastic bone lesions are non-measurable.

Cystic lesions

Lesions that meet the criteria for radiographically defined simple cysts should not be considered as malignant lesions (neither measurable nor non-measurable) since they are, by definition, simple cysts. "Cystic lesions" thought to represent cystic metastases can be considered as measurable lesions, if they meet the definition of measurability described above. However, if non-cystic lesions are present in the same patient, these are preferred for selection as target lesions.

Target lesions:

All measurable lesions up to a maximum of two lesions per organ and five lesions in total, representative of all involved organs, should be identified as **target lesions** and recorded and measured at baseline. Target lesions should be selected on the basis of their size (lesions with the longest diameter), be representative of all involved organs, and in addition should be those that lend themselves to reproducible, repeated measurements. It may be the case that, on occasion, the largest lesion does not lend itself to reproducible measurement, in which circumstance the next largest

lesion that can be reproducibly measured should be selected. A sum of the diameters (longest for non-nodal lesions, short axis for nodal lesions) for all target lesions will be calculated and reported as the baseline sum diameters. If lymph nodes are to be included in the sum, then only the short axis is added into the sum. The baseline sum diameters will be used as reference to further characterize any objective tumor regression in the measurable dimension of the disease.

Non-target lesions:

All other lesions (or sites of disease) including any measurable lesions over and above the five target lesions should be identified as non-target lesions and should also be recorded at baseline. Measurements of these lesions are not required, but the presence, absence, or in rare cases unequivocal progression of each should be noted throughout follow-up.

If multiple non-target lesions are identified in the same organ or location, it is possible to record them as a single item. (e.g., multiple enlarged pelvic lymph nodes or multiple liver metastases).

In addition, lymph nodes with the smallest short axis of less than 10 mm will be considered normal, and not included in non-target lesions.

9.2.2 Methods for Evaluation of Disease

All measurements should be taken and recorded in metric notation using a ruler or calipers. All baseline evaluations should be performed as close as possible to the beginning of treatment and never more than 4 weeks before the beginning of the treatment.

The same method of assessment and the same technique should be used to characterize each identified and reported lesion at baseline and during follow-up. Imaging-based evaluation is preferred to evaluation by clinical examination unless the lesion(s) being followed cannot be imaged but are assessable by clinical exam.

Clinical lesions:

Clinical lesions will only be considered measurable when they are superficial (e.g., skin nodules and palpable lymph nodes) and ≥ 10 mm diameter as assessed using calipers (e.g., skin nodules). In the case of skin lesions, documentation by color photography, including a ruler to estimate the size of the lesion, is recommended.

Chest x-ray:

Lesions on chest x-ray are acceptable as measurable lesions when they are clearly defined and surrounded by aerated lung. However, CT is preferable.

Conventional CT and MRI:

This guideline has defined measurability of lesions on CT scan based on the assumption that CT slice thickness is 5 mm or less. If CT scans have slice thickness greater than 5 mm, the minimum size for a measurable lesion should be twice the slice thickness. MRI is also acceptable in certain situations (e.g., for body scans), but NOT for the lungs.

As with CT, if an MRI is performed, the technical specifications of the scanning sequences used should be optimized for the evaluation of the type and site of disease. Furthermore, as with CT, the modality used at follow-up should be the same as was used at baseline and, the lesions should be measured/assessed on the same pulse sequence.

PET-CT:

At present, the low-dose or attenuation correction CT portion of a combined PET-CT is not always of optimal diagnostic CT quality for use with RECIST measurements. PET-CT scans are not always

done with oral and IV contrast. In addition, the PET portion of the CT introduces additional data which may bias an investigator if it is not routinely or serially performed. For these reasons, this study will not allow PET-CT use for RECIST 1.1 response criteria.

Ultrasound:

Ultrasound is not useful in assessment of lesion size and should not be used as a method of measurement. Ultrasound examinations cannot be reproduced in their entirety for independent review at a later date and, because they are operator dependent, it cannot be guaranteed that the same technique and measurements will be taken from one assessment to the next. If new lesions are identified by ultrasound in the course of the study, confirmation by CT or MRI is advised.

Endoscopy, Laparoscopy:

The utilization of these techniques for objective tumor evaluation is not advised. However, such techniques may be useful to confirm complete pathological response when biopsies are obtained or to determine relapse in trials where recurrence following complete response (CR) or surgical resection is an endpoint.

CA-125 (Ovarian, Fallopian tube and primary peritoneal cancer trials):

CA-125 alone cannot be used to assess response in this protocol.

9.2.3 Response Criteria

The first response evaluation will be conducted after the second cycle, followed by once every 2 cycles (after the fourth cycle, after the sixth cycle (after the eighth cycle), and when discontinuing the protocol treatment).

Determination of response should take into consideration for all target and non-target lesions and, if appropriate, biomarkers.

9.2.3.1 Evaluation of Target Lesions

Complete Response (CR):

Disappearance of all target lesions. Any pathological lymph nodes (whether target or non-target) must have reduction in short axis to <10 mm.

Partial Response (PR):

At least a 30% decrease in the sum of the diameters of target lesions, taking as reference the baseline sum diameters

Progressive Disease (PD):

At least a 20% increase in the sum of the diameters of target lesions, taking as reference the smallest sum (this includes the baseline sum if that is the smallest). In addition to the relative increase of 20%, the sum must also demonstrate an absolute increase of at least 5 mm. (Note: The appearance of one or more new lesions is also considered to be progression).

Stable Disease (SD):

Neither sufficient shrinkage to qualify for PR nor sufficient increase to qualify for PD, taking as reference the smallest sum diameters.

Not evaluable (NE):

When at least one target lesion is not evaluated at a particular time point.

$$\text{The percentage decrease in the sum of diameters} = \frac{\text{The sum of diameters measured prior to treatment} - \text{The sum of diameters measured at evaluation}}{\text{The sum of diameters measured prior to treatment}} \times 100 \%$$

$$\text{The percentage increase in the sum of diameters} = \frac{\text{The sum of diameters measured at evaluation} - \text{The smallest sum diameters}}{\text{The smallest sum diameters}} \times 100 \%$$

9.2.3.2 Evaluation of Non-Target Lesions

Complete Response (CR):

Disappearance of all non-target lesions. All lymph nodes must be non-pathological in size (<10 mm short axis).

Note: If CA-125 is initially above the upper normal limit, it must normalize for a patient to be considered to be in complete clinical response.

Non-CR/Non-PD:

Persistence of one or more non-target lesion(s).

Progressive Disease (PD):

Appearance of one or more new lesions and/or *unequivocal progression* of existing non-target lesions. *Unequivocal progression* should not normally take precedence over target lesion status. It must be representative of overall disease status change, not an increase in size of a single lesion.

Not evaluable (NE):

When at least one non-target lesion is not evaluated at a particular time point.

Although a clear progression of only “non-target” lesions is exceptional, the opinion of the treating physician should prevail in such circumstances, and the progression status should be confirmed at a later time by the principal Investigator.

9.2.3.3 Evaluation of biomarkers

Biomarker-based progression or recurrence involves assessing the patient's longitudinal CA-125 values. However, CA125 alone cannot be used to assess response in this protocol. If [an elevated?] CA125 value is observed, radiographic imaging should be obtained.

9.2.3.4 Evaluation of new lesions

New lesions are those that appear for the first time during the protocol treatment, and this category does not include any tumor lesions recorded prior to the start of treatment.

However, to be designated as “new lesions,” it is necessary to ensure that there is no change in images due to differences in scanning technique or imaging modality from the methods used for baseline evaluations, and that there is no change in images due to the presence of something other than a tumor. For example, a cystic lesion that appeared in the focal area due to a necrotic liver metastasis shall not be considered a new lesion. Newly identified lesions during follow-up in a location that was not examined at baseline (before registration) will be considered as new lesions.