Access to the NCCN Chemotherapy Order Templates (NCCN Templates®) for non-commercial users is available via subscription.

Prior to accessing the NCCN Templates®, users must accept an End-User License Agreement (EULA) and create a free account or login with an existing account on NCCN.org.

About the NCCN Templates®

NCCN continues to add to the library of chemotherapy order templates to improve the safe use of drugs and biologics in cancer care. NCCN Templates are intended for personal and practical use only. The information contained in the NCCN Templates is based on the NCCN Clinical Practice Guidelines in Oncology (NCCN Guidelines®) and the NCCN Drugs & Biologics Compendium (NCCN Compendium®). The NCCN Templates include chemotherapy, immunotherapy, supportive care agents, monitoring parameters, and safety instructions. Special instructions for self-administered chemotherapeutic agents are also provided.

NCCN Templates enhance patient safety by allowing you to:

- Standardize patient care
- Reduce medication errors
- Anticipate and manage adverse events

An NCCN Template does not constitute an order. Any clinician seeking to treat a patient using the NCCN Templates is expected to use independent medical judgement in the context of the individual clinical circumstances specific to the patient’s care or treatment.

The NCCN Templates Committee and the NCCN Templates reviewers play a critical role in the development and maintenance of the NCCN Templates. The NCCN Templates Committee and NCCN Templates reviewers consist of physicians, pharmacists, and nurses from NCCN Member Institutions. They are selected based on their clinical expertise with regard to systemic therapies as well as disease-specific subspecialty areas. NCCN Template content is reviewed annually based on the NCCN Guidelines®, the NCCN Compendium®, published drug information and research, and clinical experience.

NCCN recognizes and thanks committee members and volunteer reviewers for contributing their time and expertise by listing their names on NCCN.org/templates.
The NCCN Templates website contains a drop-down menu for displaying the template library by cancer type and/or agent name.

NCCN Chemotherapy Order Templates (NCCN Templates®)

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NCCN Templates® include regimens for the following diseases:

Please choose a cancer type:

And/or

Please choose an agent:

Reset Filters

Appendices A through G provide supplementary information about common topics across the library of chemotherapy order templates.

Endorsed resources are listed that may be helpful in applying the information contained in the chemotherapy order templates.
To display the template library for the cancer type or agent of your choice, select any item from the drop-down menus.

<table>
<thead>
<tr>
<th>Regimen Name</th>
<th>Disease Name</th>
<th>Indication(s)</th>
<th>Template ID</th>
<th>Last Modified Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>VinCREL-Bine</td>
<td>Cervical Cancer</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Topotecan/CISplatin</td>
<td>Cervical Cancer</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Topotecan Weekly</td>
<td>Cervical Cancer</td>
<td>Recurrent or Metastatic</td>
<td>CRV22</td>
<td>06/09/2018</td>
</tr>
<tr>
<td>Topotecan Every 28 Days</td>
<td>Cervical Cancer</td>
<td>Recurrent or Metastatic</td>
<td>CRV5</td>
<td>06/09/2018</td>
</tr>
<tr>
<td>PEM/Etreted</td>
<td>Cervical Cancer</td>
<td>Recurrent or Metastatic</td>
<td>CRV19</td>
<td>06/09/2018</td>
</tr>
<tr>
<td>Pembrolizumab</td>
<td>Cervical Cancer</td>
<td>Recurrent or Metastatic; Subsequent therapy - MSI-H/dMMR tumors or PD-L1 positive</td>
<td>CRV20</td>
<td>06/09/2018</td>
</tr>
<tr>
<td>PACLItaxel+Topotecan + Bevacizumab</td>
<td>Cervical Cancer</td>
<td>Recurrent or Metastatic</td>
<td>CRV25</td>
<td>06/09/2018</td>
</tr>
<tr>
<td>PACLItaxel+Topotecan</td>
<td>Cervical Cancer</td>
<td>Recurrent or Metastatic</td>
<td>CRV10</td>
<td>06/09/2018</td>
</tr>
<tr>
<td>PACLItaxel+CISplatin + Bevacizumab</td>
<td>Cervical Cancer</td>
<td>Recurrent or Metastatic</td>
<td>CRV1</td>
<td>06/09/2018</td>
</tr>
<tr>
<td>PACLItaxel+CISplatin</td>
<td>Cervical Cancer</td>
<td>Recurrent or Metastatic</td>
<td>CRV27</td>
<td>06/09/2018</td>
</tr>
</tbody>
</table>

Text can be entered into the “Search” box to narrow down the results using information in any of the columns.

Click the “Sort” icon on any of the columns to sort alphabetically.

Click on the Regimen Name hyperlink to open a template.
Below is an example of an NCCN Chemotherapy Order Template, CRV1: PACLitaxel/CISplatin for Cervical Cancer. Each section is described in more detail using the associated letters.

A. Template Header/Regimen Name

The template header lists the cancer type for which the regimen is recommended, and is associated with a specific NCCN Guideline. The regimen name is listed below the cancer type and includes the regimen acronym (if applicable), the agents included in the regimen, and may also include the length of the regimen if the same regimen has more than one option for cycle length.

Tall Man lettering is included where applicable, as described in more detail in Appendix G: Tall Man Lettering.

B. Indication

The indication(s) is/are derived directly from the associated NCCN Guidelines. These are usually summarized, thus it is recommended to refer to the associated NCCN Guidelines for more detailed information. NCCN Templates are also linked to the corresponding entry (or entries) in the NCCN Compendium.

C. References

The active links in this section include the associated NCCN Guidelines as well as published literature that supports the listed regimen. Each reference is assigned a superscript according to the classification outlined in Appendix E: Regimen References.

Continued on next page.
Below is an example of an NCCN Chemotherapy Order Template, CRV1: PACLitaxel/CISplatin for Cervical Cancer. Each section is described in more detail using the associated letters (continued).

D. NCCN Supportive Care

This section addresses emetic risk and febrile neutropenia risk levels.

Emetic Risk

The emetic risk level listed on the NCCN Templates is based on recommendations in the NCCN Guidelines for Antiemesis. The highest emetic risk level for each day of therapy is listed in this section and includes all days of treatment.

For more information on emetic risk levels, please refer to Appendix D: Nausea/Vomiting.

Febrile Neutropenia Risk

The febrile neutropenia risk level listed on the NCCN Templates is based on recommendations in the NCCN Guidelines for Myeloid Growth Factors. If the specific regimen is not included in the NCCN Guidelines for Myeloid Growth Factors, NCCN may add a febrile neutropenia risk level to the template if appropriate based on a review of the literature.

Risk levels of either “High Risk” or “Intermediate Risk” are called out specifically in this section of the templates. Regimens with unique considerations, unknown risk, or low risk based on the available literature refer back to the NCCN Guidelines for consideration of additional variables including patient- and disease-specific factors.

For more information on febrile neutropenia risk, please refer to Appendix C: Growth Factors.
E. Chemotherapy Regimen

This section focuses on drug administration, including cycle definition (which contains the cycle length, number of cycles, and other schedule-related information), dosing, frequency, and routes of administration. For standardization, regimens with continuous daily dosing are represented using a 28-day cycle length.

The NCCN Templates designate a specific order of administration if conclusive evidence is available to support a suggested chemotherapy sequence based on improved efficacy, decreased toxicity, or established clinical practice. Regimens with a recommended order of administration are designated with connecting phrases such as “concurrent with” or “followed by” as listed in CRV1 above. For more information, please refer to Appendix F: Chemotherapy Administration Sequence.

For more information regarding chemotherapy calculations, please refer to Appendix A: Chemotherapy Calculations.

For more information regarding carboplatin dosing, please refer to Appendix B: Carboplatin Dosing.

Below is an example of an NCCN Chemotherapy Order Template, CRV1: PACLitaxel/CISplatin for Cervical Cancer. Each section is described in more detail using the associated letters (continued).
F. Supportive Care

This section addresses specific recommendations for Premedications, Antiemetic Therapy, Myeloid Growth Factor Therapy, and Other Supportive Therapy. Only the sections that are relevant to a particular regimen will display on the template.

Premedications
This section includes specific recommendations for premedication(s) for reasons including, but not limited to, infusion reactions/hypersensitivity, fluid retention, and arachnoiditis. Doses may appear as ranges if clinically appropriate, to allow for provider or institutional customization based on product availability and other considerations.

Antiemetic Therapy
This section includes general guidance for selection of antiemetic therapy based on the emetic risk designated for the regimen. Links to the NCCN Guidelines and Appendix D: Nausea/Vomiting are included for more information.

Myeloid Growth Factor Therapy
This section includes general guidance for selection of prophylactic colony stimulating factor (CSF) support based on the febrile neutropenia risk level. Links to the NCCN Guidelines and Appendix C: Growth Factors are included for more information.

Continued on next page.
F. Supportive Care (continued)

Other Supportive Therapy
This section includes general recommendations with examples for supportive care medications, such as hydration, anti-infectives, or anti-diarrheals. These notes are not meant to be prescriptive, but rather to alert clinicians that patients may require additional treatment support.

G. Monitoring and Hold Parameters
The information in this section includes recommendations for monitoring found in the NCCN Guidelines, drug package insert, other drug information resources, and clinical experience. Adverse effects, including those listed as warnings and precautions are assessed for frequency of occurrence, as well as for actionable measures that could be taken either via routine monitoring or via treatment once the adverse event has occurred.

When appropriate, recommendations for laboratory tests or other assessments to monitor for toxicities and adverse reactions are provided in a general format to allow for discretion of the ordering prescriber or institutional preference as clinically appropriate. The level of specificity may vary depending on the available information, and clinicians are encouraged to refer to the package insert for more information. Examples of adverse effects that are generally excluded from the templates include fatigue, weakness, and malaise.

Notes in this section may state that potential dose modification or discontinuation may be required based on toxicity or tolerability. Dose modification refers to actions including, but not limited to, dose reduction, change in frequency, and/or holding the drug for a period of time. Clinicians are encouraged to review the package insert for more detailed information.

H. Safety Parameters and Special Instructions
This section reviews specific safety considerations as well as unique administration instructions. Examples of the information in this section include the use of filters or specific tubing requirements, vesicant/irritant properties, drug interactions, administration of oral medications with or without food, and REMS program requirements.