Pharmacy Cancer Clinical Trial Feasibility Checklist Template

**Verison 1**

**Approved: 06/12/2024 by NCCP Cancer Clinical Trials Pharmacist Subgroup**

* **This checklist in not intended to be used with Advanced Therapy Medicinal Products (ATMPs).**
* **To note, this template is intended to be a guide only; the fields included in this form are not mandatory or exhaustive. Some fields will not be relevant to all cancer clinical trials.**
* **This checklist may be adapted for use at each site as deemed applicable to local practice.**

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| **General Information** | |
| Study Protocol Name: | ABC |
| Protocol Number: | 123 |
| Description: |  |
| Principal Investigator: |  |
| Investigational Medicinal Product (IMP): |  |
| Standard Of Care (SOC): |  |
| Site Number: |  |
| Likely Site Initiation Visit (SIV) Date: |  |
| Study Type: | Open -label  Blinded  (All staff are blinded)  **(**Pharmacy staff are unblinded) |
| Research nurse: |  |

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| **Clinical Trial Impact Assessment** | |
| Recruitment period |  |
| Target patient number for this site |  |
| Oral agents only in trial? | Yes No |
| Aseptic Compounding required?  To note, the complexity band definitions outlined in Appendix 2 of the NCCP Parenteral SACT Capacity Planning Toolkit User Manual may be useful – linked [here](https://www.hse.ie/eng/services/list/5/cancer/profinfo/medonc/sactguidance/aseptic%20unit%20capacity%20planning%20user%20manual.pdf) | Yes No  Compounding complexity / time to compound: |
| Special storage requirements, e.g. freezer |  |
| Frequency of dispensing episodes |  |
| Frequency of monitoring visits | In person Remote |
| Specific non-routine testing (e.g. molecular) required? |  |
| Is Pharmacy specific training required (post SIV)? | Yes No |
| Are any specific biosimilar (s) / brand of drug required for use in the trial? | Yes No  Details \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ |
| If CSTD in use at site | CRA informed of local practice |

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| **Impact on Pharmacy Services**  Questions to consider in addition to the information above: | |
| How does this study differ from the normal clinical pathway? |  |
| How many patients on this trial can pharmacy reasonably accommodate per week considering current ACU/ dispensary capacity? |  |

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| **Other relevant notes (Practical aspects) / Items which require follow up** |
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