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Guideline on the evaluation of anticancer medicinal products in man

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Appendices:

Appendix 1: Methodological consideration for using progression-free survival (PFS) or disease-free survival (DFS) in confirmatory trials - Rev. 1 – Appendix available here:
http://www.ema.europa.eu/ema/index.jsp?curl=pages/regulation/general/general_content_000406.jsp&mid=WC0b01ac0580034cf3

Appendix 2: Use of patient reported outcome (PRO) measures and health-related quality of life (HRQoL) in Oncology (draft -in preparation)

Appendix 3: Paediatric Oncology (Rev. 2 - ongoing)

Appendix 4: Condition Specific Guidance – Appendix available here:
http://www.ema.europa.eu/ema/index.jsp?curl=pages/regulation/general/general_content_000406.jsp&mid=WC0b01ac0580034cf3

Executive summary

The purpose of this guideline is to provide guidance on all stages of clinical drug development for the treatment of malignancies, including drug resistance modifiers or normal tissue protective compounds. Supportive measures such as anti-emetics and haematopoietic growth factors, however, are covered by separate guidelines.

Alongside conventional aims such as defining the proper dose(s) and schedule(s), the importance of identifying a target population with optimised benefit risk is emphasised in Section 6: Exploratory Studies. Guidance is also provided on combination studies. Combinations of drugs with minimal activity as monotherapy, but synergistic effects when combined, as well as combinations of conventional cytotoxics, are also discussed.

Convincingly demonstrated favourable effects on overall survival (OS) are from both a clinical and methodological perspective the most persuasive outcome of a clinical trial. Prolonged progression-free or disease-free survival (PFS/DFS), however, are in most cases as such considered relevant measures of patients benefit, but the magnitude of the treatment effect should be sufficiently large to outbalance toxicity and tolerability problems. In order to capture possible negative effects on the activity of next-line therapies and also treatment related fatalities, informative data on overall survival compatible with a trend towards favourable outcome are normally expected at time of submission. This has consequences with respect to interim analyses, other than for futility, and cross-over, which thus should be undertaken only when available survival data provide the information needed for a proper evaluation of benefit/risk.

An assessment of benefit/risk should encompass all relevant data on efficacy and safety, also taking into account uncertainties as well as external data of relevance in relation to the experimental compound and the disease to be treated. Therefore no precise definition of “trend towards favourable effects on survival” or “reasonably excluding negative effects on OS” is given in this document. If a major increase in toxicity is foreseeable (see section 7), it is recommended that confirmatory studies are undertaken with the aim to show an OS benefit. It is also acknowledged that improved safety without loss in efficacy may constitute tangible aims and the design of non-inferiority efficacy studies are discussed in 7.7.3.

In section 8, definitions and abbreviations used in this guideline are summarised. Appendix 1 provides methodological guidance on the use of PFS as endpoint in confirmatory studies. A planned appendix 2 will focus on the use of patient reported outcome (PRO) measures and health-related quality of life (HRQoL) from a regulatory perspective. A revised paediatric guideline is also foreseen as Appendix 3 and Appendix 4 is dedicated to condition specific guidance.

1. Introduction

The guideline on anticancer medicinal products adopted in 1996, and revised in 2001 and 2003, focused on conventional cytotoxic compounds. In 2005, a major revision was undertaken, aiming at covering non-cytotoxic compounds, to expand on the sections on exploratory trials and to provide more guidance with respect to methodological issues. Later, there followed an appendix on methodological issues related to use of PFS and in early 2010 an appendix on haematological malignancies followed. In this appendix disease specific guidance was introduced and the section on confirmatory studies based on aims of therapy and relative toxicity was restructured. These latter elements have now been incorporated in the revised main guideline. In this revision, the chapter on

exploratory trials for cytotoxic compound has been shortened as it was considered too detailed and too prescriptive.

The section on condition specific guidance (Appendix 4) has been expanded and now constitutes a separate Appendix.

2. Scope

Whilst the thrust of a regulatory guideline should be on confirmatory studies, the aim of this guideline is also to underline the importance of exploratory studies in order to identify the most appropriate target population in addition to the usual aims: to define dose, schedule, tumour type and line of therapy. The role of biomarkers to achieve these objectives is also further emphasised in this revised guideline.

There are numerous possible ways to classify anti-cancer drugs such as direct anti-tumoural vs. indirect anti-tumoural, or based on pharmacology or molecular target (e.g. hormones, immune modulators, nuclear-targeting, signal-transduction targeting, etc.). As this document is meant to provide guidance on clinical drug development, the aim has been to classify compounds according to reasonable designs of exploratory studies, i.e. cytotoxic compounds where toxicity and ORR are considered suitable markers of activity in dose finding studies vs. non-cytotoxic compounds where ORR and/or toxicity may not serve this purpose.

A very large number of anti-cancer compounds have been and currently are under development. Only a minority, however, have completed the clinical development and obtained a marketing authorisation, due to poor activity or evidence of a detrimental safety profile. Until non-clinical models with good predictive properties have been defined, this situation is likely to remain essentially unchanged and the absence of such models is considered to constitute the greatest hurdle for efficient drug development within the foreseeable future.

Since chemoprotective agents and drug resistance modifiers are used as part of anticancer regimens, some guidance on these agents will also be provided in appropriate sections of this guideline. Anti-emetics and haematopoietic growth factors, however, are covered in separate documents.

3. Legal basis and relevant guidelines

This document should be read in conjunction with Directive 2001/83/EC, as amended. Applicants should also refer to other relevant European and ICH guidelines on the conduct of clinical trials, including those on:

- Nonclinical evaluation for anticancer pharmaceuticals EMEA/CHMP/ICH/646107/2008 (ICH S9)
- Clinical Investigation of the Pharmacokinetics of Therapeutic Proteins CHMP/EWP/89249/2004
- Evaluation of the Pharmacokinetics of Medicinal Products in Patients with Impaired Hepatic Function - CPMP/EWP/2339/02
- Guideline on the investigation of drug interactions, CPMP/EWP/560/95/Rev. 1
- Points to Consider on Adjustment for Baseline Covariates - CPMP/EWP/2863/99
- Points to Consider on Multiplicity Issues in Clinical Trials - CPMP/EWP/908/99

- Guideline on the choice of non-inferiority margin - CPMP/EWP/2158/99
- Qualification of novel methodologies for drug development: guidance to applicants EMA/CHMP/SAWP/72894/2008 Rev.1
- Reflection paper on methodological issues associated with pharmacogenomic biomarkers in relation to clinical development and patient selection EMA/CHMP446337/2011
- Reflection paper on pharmacogenomics in oncology EMEA/CHMP/PGxWP/128435/2006.
- Guideline on clinical trials in small populations-CPMP/EWP/83561/2005
- Choice of Control Group in Clinical Trials CHMP/ICH/364/96 (ICH E10)
- Guideline on clinical evaluation of diagnostic agents - CPMP/EWP/1119/98
- Note for guidance on clinical safety data management: data elements for transmission of individual case safety reports - CPMP/ICH/287/95 (ICH E2B)
- Points to consider on application with 1. Meta-analyses 2. One pivotal study - CPMP/EWP/2330/99
- Reflection paper on methodological issues in confirmatory trials planned with an adaptive design – CHMP/EWP/2459/02

4. Pharmacokinetics

In general, the same recommendations are valid for anticancer products as for other medicinal products and reference is made to the clinical pharmacology guidelines available. For therapeutic proteins, reference is made to CHMP/EWP/89249/2004. This section is thus mainly meant to highlight some areas where missing information frequently has been encountered in submissions for marketing authorisation and to underline some areas considered to be of special interest.

In the past, human mass-balance studies (in vivo studies investigating the fate of a radiolabelled dose in plasma and excreta) have not been performed to the same extent for anticancer drugs as for other medicinal products. Due to the importance of the information gained in these studies for the understanding of the clinical pharmacology of the investigational drug, including the drug-drug interactions assessment, mass-balance studies are strongly recommended (CPMP/EWP/560/95/Rev. 1).

Food interaction studies should be performed prior to phase III and administration in fed or fasted state should be investigated and a rationale for administration in fed and/or fasted state should be provided. .

The potential for drug-drug interactions should be assessed. If in vitro data indicate that the anticancer product will give rise to, or be a victim of, important drug-interactions, this should as far as possible be investigated in vivo.

Studies to be undertaken in patients with impaired organ function should mainly be selected based on prior information on the mode of elimination of the drug and formation/elimination of potential pharmacologically active metabolites. If a study in hepatic impairment is needed and liver metastases are common in the target patient population, as a first step a study in patients with liver metastases is warranted. Whether studies in more advanced liver disease are needed should be decided on a case by case basis (CPMP/EWP/2339/02). Lack of data is reflected in the SmPC. Exploratory studies, including

PK, in patients with malignant ascites or other third space conditions such as massive pleura fluid are encouraged if seen in the condition being treated.

It is recommended to also evaluate the influence of intrinsic factors through population PK analyses. The plasma concentration data should optimally come from as many as possible of the clinical studies. Both sparse (few samples per patient) and rich data (full plasma concentration-time profiles) can be used. Factors to investigate as covariates could include age, weight, gender, renal function, S-bilirubin, liver enzymes, genotype, soluble receptors/ligands, tumour burden, inflammatory markers etc.

The use of PK and PD (biomarkers and clinical markers) sampling for PK/PD analysis related to efficacy and safety is encouraged. This information aids in understanding the exposure-response relationships for the drug, and may allow for a rational selection of treatment strategies in patients who are at risk for excessive toxicity or ineffective therapy. Exposure-efficacy and exposure-safety analysis/modelling is encouraged in the Phase II randomized trials (sections 6.2 and 6.3) to provide PK/PD information and to support Phase III dose selection. Ultimately, a pooled analysis of PK and PD data obtained in all phases of development is encouraged in order to fully characterize and summarize the PK/PD of the drug. In order to utilize all collected data efficiently, longitudinal PK/PD analysis of PD data e.g. tumour shrinkage as a continuous variable is recommended. Simulation based evaluations of the study design with respect to power of identifying PK/PD relationships and covariate effects are recommended. Due to high withdrawal rates leading to informative censoring, handling of missing data is of crucial importance in longitudinal analyses and sensitivity analyses, e.g. using early time points for tumour shrinkage should be considered.

5. Biomarkers

In order to optimise benefit – risk, it is essential to identify the proper target population for therapy. This might be possible to accomplish through the judicious use of biomarkers in all phases of clinical drug development. A biomarker should be capable of objectively measuring and evaluating a normal biological process, a pathological process or the pharmacological response to a therapeutic intervention, depending upon its purpose. A suitable biomarker may be identified and measured by a variety of different diagnostic approaches (e.g. expression profiling of transcripts, differential antigen expression, genetic diagnostics, including next generation sequencing, etc).

Irrespective of pharmacological class, it is assumed that entrance into clinical development of new molecule today is guided by translational research. This means that in most cases there are hypotheses to be tested and candidate biomarkers available. The utility of biomarkers is broad e.g. prospective stratification of clinical trial subjects according to biomarker status, determination of the biologically effective dose, early proof of mechanism or concept, assessment of toxicity and an indication of the natural course of a disease. However, although efforts to identify targets and explain variability in PK and PD are essential, the need to confirm the findings should not be overlooked in the planning of the drug development programme (technical and clinical validation). For patient stratification, if convincing evidence of biomarker selectivity is established early in the non-clinical and clinical development phase, confirmatory evidence in the negative population may not be required and such studies may be carried out in patients expressing the biomarker of interest.

It is acknowledged that biomarkers tested in early clinical trials are often exploratory in nature, but it is essential that technical/quantitative reliability is assured (EMA/CHMP/SAWP/72894/2008 Rev.1, EMEA/CHMP/PGxWP/128435/2006). While serum biomarkers or other sources of biological samples might be informative, tumour samples are expected to constitute an integral part of the biomarker exercise, if not otherwise justified. It is acknowledged, however, that single biopsies may not be

representative due to tumour heterogeneity. Normal tissues samples may also be used in early clinical studies, if non-clinical studies indicate that there is a correlation between the changes observed in normal tissues and the features of the tumour. The role of functional imaging in early drug development is not regarded as well established, but its use is encouraged.

The development of biomarker diagnostic methods should be considered early in clinical development, maximising the clinical application of the technology. A diagnostic assay complying with the requirements laid down in IVD Directive (98/79/EC), as appropriate, should be available at time of licensure

For the use in confirmatory studies and e.g. as measures of efficacy, biomarkers must be carefully and rigorously validated, ideally following systematic evaluation in well designed prospective clinical trials (EMA/CHMP/446337/2011). Of note, this guideline also opens for the possibility retrospective validation through replication of findings. In order to assist in interpretation of results across studies and limit sources of variability when developing biomarkers, the use of available reporting guidelines is encouraged.

6. Exploratory studies

Exploratory studies are essential in rational drug development. The distinction between Phase I/II exploratory and Phase III confirmatory trials has been adhered to in this Guideline. However, this does not mean that exploratory aims should not form an important part of Phase III trials. Similarly, hypothesis generation, testing and confirmation may form parts of Phase II trials.

So called phase 0 trials, i.e. trials exploring micro dosages may be informative in certain circumstances as regards tissue distribution and receptor binding, e.g. when it is considered important to early identify whether a compound is likely to penetrate sanctuaries, such as CNS, or, when feasible, to obtain early data on pharmacological activity at low drug concentrations.

6.1. Cytotoxic compounds

This section refers to conventional cytotoxic agents, i.e. compounds inducing irreversible lethal cellular damage following short-term exposure through interference with DNA replication, mitosis, etc. For these compounds, toxicity and tumour response are considered suitable indicators of activity.

Conceptually this section is also relevant to more targeted cytotoxic compounds such as monoclonal antibody coupled toxin products. In these circumstances however, tumour antigen expression and prodrug activating pathways should also be taken into consideration.

As for non-cytotoxic compounds, non-clinical and clinical studies encompassing aims to characterise prerequisites for activity/resistance and to identify markers of resistance are encouraged.

6.1.1. Phase I, single agent dose and schedule finding trials

The basic assumption governing the design of these trials is that, for dose finding purposes, toxicity is an acceptable endpoint. The main objective is thus to define dose-limiting toxicities and the dose to bring forward into further trials. While meeting this objective is generally straightforward, in spite of the fact that the inter-patient variability in PK might be large, it is often more complex to define reasonable dose schedules to study further.

Initial dosing may use flat doses or body surface area (BSA) scaled doses. The scientific support for the notion that BSA scaled dosing generally reduces inter-patient variability in exposure is weak and may

lead to over and under-exposure in patients with a high and low BSA, respectively. It is expected that the importance of BSA or weight for variability in exposure is explored through modelling & simulation using actual pharmacokinetic data.

The use of pharmacodynamic endpoints, where available, may also assist in dose selection

Main Objectives

- Maximum Tolerated Dose (MTD), Dose Limiting Toxicity (DLT) and recommended Phase II dose (RP2D) should be identified for defined schedules and modes of administration
- Frequent side effects and target organs for toxicity should be characterised as regards relationship to dose and schedule. Severity, duration and reversibility should be determined.
- Initial characterisation of pharmacokinetics including dose and time-dependencies. As appropriate, PK/PD related to target effects and adverse effects, exposures obtained with different routes of administration.

Eligibility of patients

These trials should normally be undertaken in cancer patients without established therapeutic alternatives.

Routes of administration and schedules

The choice of route and rate of administration of the first dose in man should be justified based on the non-clinical data. In most cases, intravenous administration, when feasible, is advisable for first use in man studies since it eliminates variability related to bioavailability.

For schedule finding, experience related to class of compounds is helpful. Non-clinical data with respect to cycle dependency and the ratio tumour / normal tissue cytotoxicity *ex vivo* may be of some interest.

Dose escalation

In case of minimal toxicity, or occasionally in case of non-significant toxicity, within-patient dose escalation may be appropriate in order to reduce the number of patients exposed to non-active doses. This may be acceptable after the end of the period of DLT assessment, if non-clinical data provide evidence of no cumulative toxicity.

If toxicity is acceptable, the patient may be re-exposed upon recovery and preferably should receive at least 2 cycles at the same dose level.

Evaluation of toxicity

The minimal requirements for evaluation of adverse effects include assessment of symptoms, physical examination, ECG, blood and urine laboratory analyses and radiological assessment as appropriate. Preclinical data should be used to guide the need for further examinations. If there are no signals with respect to QTc in preclinical studies or related to class of products, no dedicated QTc studies are expected, but inclusion of ECG as part of routine monitoring is recommended. Local toxicity at the site of administration should be specifically recorded. The toxicity should be graded according to a generally recognised system (e.g. Common Terminology Criteria for Adverse Events, CTCAE).

Factors influencing toxicity (organ dysfunction, concomitant therapy) should be explored as appropriate. These factors should be further elucidated in Phase II/III.

6.1.2. Phase II, single agent therapeutic exploratory studies

Phase II trials may investigate single-agent activity in a variety of tumour types, or in a selected tumour type, or investigate activity and feasibility of combination or multimodality regimens.

This section is focused on trials where the primary objective is to estimate single agent antitumour activity in patients with a defined tumour type in order to identify compounds to bring forward to confirmatory trial.

Objectives and design

Phase II trials may use a variety of study designs and early studies should provide initial evidence of treatment activity and tolerability. Inclusion of a randomised control arm is encouraged, particularly if only one confirmatory pivotal trial is foreseen (see section 7.1.2).

The studies are intended to:

- Assess the probability of response (and other relevant efficacy measures) in the target tumour type and conclude on the need for further studies (investigate earlier stages of the disease, combinations, compare with standard therapy).
- Investigate pharmacogenomics and biomarker characteristics, where appropriate
- Further characterise dose and schedule dependency, with respect to safety and activity
- Further characterise the side-effects of the medicinal product
- Further characterise PK and PK/PD (see section 4)
- When applicable, further characterise the optimum route of administration

Selection and number of patients

Exact definition of the target disease, previous therapy (if any) and stage should be given, in line with internationally agreed diagnostic criteria.

Provided safety and activity is reasonably established and there is a scientific rationale, it might be appropriate to conduct studies also in patients for whom alternative therapies are available. This includes the neo-adjuvant setting in treatment naïve patients scheduled for surgery, provided that delay in surgery cannot be unfavourable to the patient. The safety and interests of the patient must always be guaranteed and a detailed justification should be provided in the study protocol. In these cases, the use of sensitive measures of anti-tumour activity such as functional imaging is expected.

Dose and schedule

The dose and schedule should be clearly defined. Details on the administration of the medicinal product with special precautions (hydration of patients, protection against light and temperature, etc.) should be stated as well as other agents, which are contraindicated during the study period.

- Guidance should be supplied outlining dose reductions related to the severity of the observed toxicity.
- As appropriate, guidance outlining dose escalations in case of low toxicity may be incorporated.
- Consideration should be given to study high-risk patients (e.g. high risk with respect to target organ toxicity or compromised metabolic or excretory mechanisms for the experimental compound) separately.

Any evidence of cumulative toxicity should be recorded and estimated as a function of total dose. This should be specifically studied according to target organ or function.

Evaluation of activity

ORR should be documented according to international standards (e.g. RECIST, Volumetric RECIST or WHO criteria). Modifications of these criteria may be appropriate in certain situations, but should be justified.

In evaluating ORR, the ITT principle should be adhered to. In single arm studies, ORR in the per-protocol analysis set may be reported as primary outcome measure. External independent review of tumour response is encouraged, according to the objectives of the trial.

Data on duration of response, TTP/PFS, confirmed ORR and available data on OS should normally be reported. The use of tumour biomarkers and other dynamic measures of activity is encouraged.

In haematological malignancies, disease specific response criteria are unavoidable in many cases and full harmonization has not yet been accomplished for some disease entities. Therefore it is of importance to follow the progress made by international working groups on these issues. Especially if less conservative disease specific response criteria are introduced in new clinical guidelines, a justification with focus on aspects of drug development is expected from the sponsor.

In patients with symptomatic disease at base line, the assessment of symptom control is encouraged, if a randomised phase II trial is undertaken.

6.2. Non-cytotoxic compounds

This refers to a very heterogeneous group of compounds ranging from antihormonal agents to antisense compounds, signal transduction, angiogenesis or cell cycle inhibitors, immune modulators, etc. The common element affecting the design of clinical trials is that toxicity may not be an appropriate endpoint in dose and schedule finding trials and ORR may not be an appropriate measure of anti-tumour activity.

For these reasons, the early stages of clinical drug development are more complex and have to be tailored according to the assumed pharmacology of the individual compound as defined in non-clinical studies. The rather strict delineation between Phase I and II trials, as for conventional cytotoxic compounds, may be less relevant as measures of anti-tumour activity, e.g. based on assessment of biomarkers might be needed early in order to define dose and schedule.

Otherwise, most of the elements discussed in relation to cytotoxic drugs are of relevance also here such as restrictions with respect to patient eligibility, recommendations as regards routes of administration, evaluation of toxicity and anti-tumour activity, etc. These issues will not be further discussed here.

6.2.1. Phase I, single agent dose and schedule finding trials

Non-clinical data and, when available, data from healthy volunteers should be used to design the studies to be conducted in patients, e.g. as regards eligibility criteria and starting dose. In accordance with the guidance for cytotoxic compounds, availability of established therapies should normally be regarded as an exclusion criterion. Refractoriness to conventional cytotoxic compounds, however, may confer resistance also to some clearly non-related compounds. This obviously affects the possibility to define a dose/concentration – effect relationship. All sensible and ethically acceptable measures undertaken to increase the assay sensitivity of these clinical trials, including the conduct of window of

opportunity studies (Definitions and Abbreviations, 8) are encouraged. Whenever appropriate, this includes measuring the expression of the assumed target(s) for drug activity.

PD measures may include biochemical measures (receptor binding, enzyme inhibition, downstream events, etc. as defined in non-clinical studies), functional imaging, proteomics, immunological measures (antibody or T-cell response), etc. Population PK/PD studies are encouraged. For compounds shown to be cytostatic in non-clinical models, prolonged exposure may be needed to elicit tumour shrinkage in clinical studies. If in these cases unexpected, early tumour shrinkage is observed this constitutes a signal indicating that further studies exploring the underlying mechanisms behind early response are warranted.

While it is acknowledged that drug development for compounds with a single main target for activity, such as mutated BRAF, is more straight forward, it is still expected that the pharmacological rationale behind poly-targeting compounds is reflected in the exploratory studies programme, e.g. in terms of biomarkers selected in order to identify the proper target population for treatment.

Main objectives

- Tolerability, safety, PK and, if at all possible, PD measures of activity are appropriate objectives
- As for conventional cytotoxic drugs, the use of tumour markers and sensitive imaging techniques, in combination with conventional methods, are recommended in order to delineate possible antitumour activity. It is recommended that technical standardisation of, e.g. functional imaging techniques and biomarker assays, is implemented in order to reduce inter-centre variability.

Eligibility of patients

Based on preclinical tolerability and toxicology findings and the assumed pharmacology of the compound, early trials may sometimes be conducted in healthy volunteers.

Eligibility criteria and the number of patients should be defined according to the objectives of the study, also taking into account variability in PK and PD at doses and schedules selected for further studies.

If not pharmacologically justified, proper analyses of biopsies from accessible tumours (primaries and/or metastatic lesions), are expected to constitute a pivotal role in studies undertaken to identify the proper target population for confirmatory studies. This might be crucial and has to be considered in the recruitment of institutions, investigators and patients.

Dose escalation

Until now available experience indicates that tumour selectivity is not to be expected for most compounds. Tolerability and toxicity thus remain important measures in dose and schedule finding studies. However, there are cases where dose escalation to MTD is not adequate in order to define the recommended dose. In these cases, dose escalation can be based on pharmacodynamics and safety data in relevant animal models, and on human PK/PD data from initial and subsequent dose cohorts. Mechanism-based PK/PD modelling may also be useful to guide decision making.

6.2.2. Phase II, single agent therapeutic exploratory studies

For the purpose of simplification, it is assumed that a dose/exposure range has been defined that shows pharmacological activity/target occupancy with or without dose limiting toxicity. If not otherwise justified, it is postulated that activities related to identification of the proper target population, as discussed above, continues in these studies.

Study designs and measures of activity

ORR, despite all its shortcomings related to patient-selection, etc, is a rather convincing measure of anti-tumour activity as for most tumours, spontaneous regression fulfilling criteria for at least partial response is a rare phenomenon. For exploratory purposes, studies without a randomised reference are therefore considered interpretable and guidance provided in the section about cytotoxic compounds is relevant. Irrespective of this, inclusion of a randomised reference arm is encouraged and might be of special interest in order to explore whether, e.g. a selected biomarker is prognostic and/or predictive (see 7.1.2).

Time to progression (TTP) and progression-free survival (PFS), however, are in principle a function of underlying tumour growth rate and the activity of the anti-tumour compound. Also, if documented progressive disease is an inclusion criterion, underlying growth rate is hard to define in most patients and historical data will be even harder to interpret. Therefore, the interpretation of TTP/PFS data without a randomised reference is problematic. In particular in breast cancer, clinical benefit response rate (CBR), i.e. CR, PR and absence of progression at 6 months, is a well established measure of anti-tumour activity and might be used for between study comparisons, even though subject to the same principle problem as TTP/PFS.

Exploratory trials with time-related endpoints

There is probably no ideal yet feasible design of exploratory studies for compounds assumed to mainly elicit tumour growth control. In the following some design alternatives are discussed, all with pros and cons, but in principle acceptable from a regulatory perspective. Irrespective of design, it is recommended that only patients with documented tumour progression are enrolled.

- A randomised dose comparative trial, e.g. comparing the lowest dose likely to be pharmacologically active with higher dose(s), if showing a difference in TTP/PFS, will obviously provide evidence of activity, but not in absolute terms.
- Randomised withdrawal of therapy in a single arm study in patients with non-progressive disease after a defined period of time on experimental therapy. The acceptability of this design to patients and investigators, however, may constitute an obstacle and carry-over effects may be a reality for some compounds.
- In previously treated patients, a within patient comparison of TTP/PFS might provide evidence of activity. Here TTP on last prior therapy is compared with TTP/PFS on the experimental therapy. It should be noted, however, that the underlying assumption of at least similar growth rate over time cannot always be substantiated. For exploratory purposes this constitutes no major concern. It is advisable to recruit patients with secondary as well as primary resistance on prior therapy. This ensures at least to some extent, that the study population is relevant. It should also be noted that patients with early failure (primary resistance) on prior therapy may show some inversions in terms of TTP just due to fluctuations in tumour growth rate and variability related to imaging techniques.

For certain indications, a within patient comparison may be justified also in treatment naïve patients, i.e. patients are followed without therapy until progression followed by experimental therapy until progression.

- A randomised phase II study versus a compound known to be active in the selected population (or placebo/BSC if justified) provides another alternative. In a comparison in terms of TTP/PFS it should be noted, that a purely growth inhibitory compound is "favoured" compared with a compound inducing tumour shrinkage, as progression is defined in relation to best tumour response. At the time of tumour progression, the tumour burden in patients failing a purely

growth inhibitory compound will therefore be higher than in patients where tumour shrinkage was elicited.

- If no more refined techniques are applicable, TTP/PFS and CBR without an internal reference has to be accepted as a measure of Phase II anti-tumour activity. A systematic literature review, including methodology used, is advised in these cases.

In principle, a statistical approach similar to that for Phase II trials with ORR as outcome measure is applicable. It is harder to set up criteria for early termination, however. The number of patients should be sufficient to obtain a reasonably precise estimate of the percentage of progression-free patients at a predefined time point. The underlying assumptions as regards progression rate without therapy are more problematic and “promising activity” is harder to define.

For these studies, the use of conventional criteria for ORR and tumour progression is recommended and independent review is encouraged. It is recognised, however, that, e.g. an apparent increase in tumour size due to inflammatory oedema, “pseudoprogression”, might be a first sign of activity for certain compounds. If prior trials indicate that this is the case, it is accepted that this is accounted for in the study protocol. The use of ORR and TTP as key measures of activity should not be regarded as contradictory to the use of tumour/PD markers in parallel.

If a randomised design is considered appropriate, the use of generally accepted instrument to estimate HRQoL or symptom control may provide valuable information (see Appendix 2).

For window of opportunity studies and if sensitive measures of pharmacological activity are available, e.g. functional tumour imaging and/or biomarkers, and a target population has been identified with tumours likely to be sensitive, placebo-controlled trials with one or preferably more doses of the experimental compound might be feasible. Sensitive measures, even if not fully validated with respect to relationship to ORR, are from a regulatory perspective acceptable for exploratory purposes and allow not only for refined dose comparisons, but also early escape in case of absence of activity. It is advisable though to clearly define in the protocol criteria for progressive disease, whether a composite (e.g. biomarkers, or imaging, or symptoms) is used or not.

6.3. Immune modulating compounds and Monoclonal antibodies (MoAb)

This section is primarily meant to provide guidance as regards exploratory studies, but also on some aspects of relevance for confirmatory studies.

6.3.1. Monoclonal antibodies

Monoclonal antibodies may affect tumour cells directly, e.g. through ADCC and/or blocking of growth factor/anti-apoptotic receptor signalling, or indirectly through the targeting of growth factors for the tumour or tumour supportive structures, or by blocking T cell inhibitory signals (e.g. anti-CTLA4).

In vitro non-clinical studies should be performed to elucidate the prime activity of the MoAb. These studies may include relevant assays on:

1. Binding to target antigen(s): tumour cells or plasma should be screened for (over)-expression of the target and the relationship between target expression and activity should be investigated.
2. Unwanted targets. Tumour specificity may not be attainable, but it is possible to screen for “unwanted” targets in vitro, facilitating the safety assessment.
3. Fab-associated functions (e.g. neutralization of a soluble ligand, receptor activation or blockade)

4. Fc-associated functions (e.g. antibody-dependent cell-mediated cytotoxicity, ADCC; complement-dependent cytotoxicity, CDC; complement activation)

Target-mediated disposition may be seen with MoAbs. Adequate characterization of this form of non-dose proportional PK behaviour may not be possible until late phase studies, when patients with tumours having widely variable amounts of target are studied. Therefore, continued evaluation of MoAb PK during the clinical development program, which often involves different tumour types and stages of disease is encouraged.”

Clearance of MoAbs is typically influenced by FcRn IgG cycling, immunogenicity (Anti-Drug-Antibodies (ADA)) and may also be impacted by patient health status factors (e.g. albumin, soluble receptors/ligands, disease type and severity, tumour burden, etc.). Knowledge of these factors may contribute to understanding the nature of MoAb exposure and response. The experience as regards immunogenicity of MoAbs in other fields of clinical medicine should be taken into account with respect to choice of assays, markers for loss of activity and possible safety problems.

6.3.2. Immune modulating compounds including tumour vaccines

Immune therapies including therapeutic cancer vaccines are aimed to induce specific anti-tumour immunity toward existing malignant disease. Such immune therapies are normally aimed to induce adaptive T and B cell as well as innate immune responses in cancer patients. The nature of the drug substances used is highly variable, including synthetic peptides, recombinant proteins, virus-like particles, immune-modulating antibodies, gene therapy, and cell-based products. As it is difficult to break tolerance towards tumour antigens which are normally derived from self-antigens, cancer vaccines are often combined with pharmacologically active adjuvants such as cytokines or toll-like receptor agonists. One other approach to break immune tolerance is to block T cell inhibitory signals, e.g. with monoclonal antibodies. The resulting T-cell activation and proliferation leads to wanted and unwanted immune stimulatory effects: the desired anti-tumour effect as well as the appearance of immune related toxicities like colitis and endocrine insufficiency.

Non-clinical in vitro and in vivo proof-of-concept studies should be presented to justify the planned starting dose and schedule in phase I studies. Furthermore, and on a case-by-case basis, the rationale for the starting dose may be supported by using the ‘Minimal Anticipated Biological Effect Level’ (MABEL) approach, and by non-clinical and clinical data from related compounds (EMA/CHMP/SWP/28367/07).

It is acknowledged that for products relying on human-specific antigens which need to be presented on human MHC molecules, predictive animal models are often not available. Nevertheless, animal models using homologous antigens or animals being human MHC transgenic might be considered for non-clinical pharmacology and toxicology studies, if available. Information on the differential expression of the target antigen in human tumour and healthy tissues should be provided. In case that no relevant and predictive animal model is available, in vitro studies with human cells, like e.g. in vitro T-cell priming assays might be suitable to show proof-of-concept.

The aim of early clinical trials is to determine the safety and the dose and schedule that induced a desired immune response. Dose-finding studies are generally required to establish the recommended phase II dose. Monitoring the immune response, i.e. the induction of antigen-specific T cells or the presence of a humoral response are of interest to determine appropriate dose and schedule. To achieve this goal multiple monitoring assays may be necessary and these should be carefully explored. The analytical methods should be described in detail in the clinical trial protocol.

Tumour biopsies taken before and after treatment are expected to play a pivotal role in assessing the extent and type of immune activation in the target tissue and could serve as an early marker for possible anti-tumour activity.

The induction of tumour response in patients with high tumour burden might be a too high hurdle to overcome and may favour the inclusion of patients with minimal or low tumour burden. Examples are therapy of patients with NSCLC after complete tumour resection where cancer immunotherapy can be assessed in the adjuvant setting. Another example is patients suffering from non-resectable NSCLC who have responded to chemotherapy. The design of clinical studies using clearly experimental therapies in patients with limited and measurable disease, not heavily pretreated with cytotoxic regimens has to be carefully justified. As for other agents, evidence of anti-tumour activity is essential prior to the initiation of confirmatory studies.

Oncology patients are usually taken off treatment upon disease progression. Induction of an effective immune response and clinical response may need more time to develop (delayed effect) compared to classical cytotoxic compounds. Patients may thus experience disease progression prior to the onset of biological activities or clinical effects. Discontinuation of active cancer immunotherapy in case of slow progression may not be appropriate. In these situations a detailed definition of “slowly progressive disease” and/or withdrawal criteria is expected in the study protocol and close monitoring of patients is required. The definition of “slowly progressive disease” should be guided by the course of disease under investigation. Revised criteria defining progression is accepted if properly justified, in confirmatory studies, however, OS is the recommended outcome measure.

Possible toxicities like induction of autoimmune reactivity (cellular and humoral) and induction of tolerance should be carefully monitored during the clinical development.

6.4. Combination therapy studies

Conventional cytotoxic compounds have for long been used in combination in order to increase the anti-tumour activity at acceptable levels of toxicity. This may be accomplished by combining compounds with at least partly non-overlapping toxicity and, perhaps, partly non-overlapping prerequisites for activity/resistance. Regulatory agencies, as well as learned societies, have accepted this approach, but it is acknowledged that it is frequently unknown whether combined use results in a better long-term outcome than consecutive use.

6.4.1. Combining conventional cytotoxic compounds

In the selection of patients with available alternative therapies, the documented activity of the individual components of the combination regimen should be taken into account.

The exploratory phase encompasses the determination of MTD and RP2D for the combination and a preliminary assessment of anti-tumour activity in terms of ORR and PFS/TTP. While the degree of anti-tumour activity for a new combination relies on assumptions, it is often possible to predict toxicity, based on the toxicities of the individual components. If relevant PK interactions can be excluded, and pending on the dose-response/toxicity profiles, dose-finding studies may be initiated at about 1/2 of the recommended mono-therapy dose for each compound. It might also be appropriate to start at the full recommended mono-therapy dose for one of the compounds and reduced dose (<50%) for the other compound. As the sequence of administration may be of importance with respect to potential PK interactions and anti-tumour activity, this has to be accounted for in the design of the studies.

There is no uniform way to balance dose intensity between components of a combination regimen to optimise benefit – risk. It is thus accepted that, e.g. priority in terms of dose intensity is given to the compound with the highest monotherapy activity.

If one of the components is regarded as an acceptable treatment regimen in monotherapy, a randomised phase II study comparing the monotherapy regimen with the combination is informative. For confirmatory studies a comparison with the best available, evidence-based reference regimen is expected.

6.4.2. Combinations involving a non-cytotoxic drug.

If there are no strong biological/pharmacological arguments to the contrary, the selected chemotherapy regimen to be combined with the non-cytotoxic should normally be “best available”. If the dose intensity/systemic exposure of the chemotherapy regimen is unaltered it can be assumed that all patients will receive appropriate therapy. Therefore there is no need to restrict the eligibility of patients from this perspective.

Whenever previous non-clinical and clinical experience has suggested that PD markers, etc. might be informative with regard to anti-tumour activity, they should be part of the experimental plan. This may include investigations whether the expression of the target for the non-cytotoxic compound is affected by treatment with cytotoxic agents and if appropriate *vice versa*.

Given the current status with respect to predictability of add-on activity in non-clinical models, randomised phase II studies comparing the experimental regimen with the chemotherapy-alone regimen are considered essential. For these studies, it is recommended that conventional anti-tumour activity data (ORR and TTP) are supplemented with tumour markers and sensitive measures of, e.g. tumour metabolic activity as appropriate.

When add-on activity of the non-cytotoxic compound to a chemotherapy regimen has been demonstrated, the need for further randomised phase II studies when new indications are studied may be dispensable. This, however, should be justified as the importance of target expression and inhibition thereof might differ between malignancies.

If the expression of the target for the non-cytotoxic compound may be differently affected by different chemotherapy regimens, it is advisable to study target expression during treatment with a new chemotherapy regimen prior to the conduct of add-on studies.

Research aiming at understanding the mechanisms and prerequisites for the add-on effects is encouraged, as it may allow for an improved characterisation of target populations in future studies.

It is conceivable that for some non-cytotoxic compounds, combinations are needed not only to optimise anti-tumour activity, but actually are required in order to obtain activity. For such compounds, e.g. target saturation in monotherapy and, importantly, non-clinical toxicity for the combination may be used to define suitable starting doses and schedules. Otherwise dose/schedule exploratory and therapeutic exploratory studies may proceed essentially as for a monotherapy regimen.

If supported by strong biological and/or pharmacological non-clinical and early proof-of-principle clinical data, two new compounds may be combined in a co-development program.

The following three scenarios are foreseeable:

Uni-enhancement refers to scenarios when one combination partner *B*, which has no or minimal anti-tumour activity per se, but enhances the anti-tumour activity of the other partner *A* (e.g. through prevention of resistance development). The contribution of *B* needs to be established by data from appropriate non-clinical models. In phase II the comparison to a reference treatment is encouraged, while Phase II monotherapy data for *B* may be considered dispensable. An appropriate phase II design would be a randomised three-arm study *AB vs. A vs. reference treatment*.

Co-enhancement is considered when both combination partners demonstrate (modest) anti-tumour activity per se and the anti-tumour activity of the combination is considerably increased. In phase II, the new combination should be compared to both combination partners as single agents at efficacious doses and preferably a reference treatment: *AB vs. A vs. B vs. reference treatment*. Depending on the phase II results one or both monotherapy arms may be dispensable in phase III.

In case the monotherapy arm of one combination partner (B) is part of phase III (*A+B vs. B vs. reference*) the same monotherapy may not need to be included in phase II (*A+B vs. A vs. reference treatment*).

Synthetic lethality refers to a scenario when both combination partners have no or minimal anti-tumour activity per se, but exhibit potent activity as a combination. If non-clinical and clinical studies indicate "inactivity" at dosages/exposure levels considerably above that of the combination and the combination is clearly active, the contribution of both partners may be dispensable for phase 2 and phase 3 studies.

As the same targets may have a different impact in different malignancies the necessity of both combination partners may need to be shown for new indications.

7. Phase III, confirmatory trials

Confirmatory trials should be designed with the aim to establish the benefit - risk profile of the experimental medicinal product, including supportive measures, in a well-characterised target population of relevance for clinical practice.

In the general part of this section (7.2 – 7.4), the aim of therapy, curative versus long term disease control vs. palliation and not the underlying disease has been used to structure the discussion.

For some malignancies where treatment is administered without curative intent, there are alternative, in clinical practise still well established regimens, showing major differences in anti-tumour activity. This reflects that selection of therapy in the clinic is guided by efficacy and safety. It is therefore of relevance in the planning phase to take into account the expected tolerability/toxicity profile of the experimental regimen compared with the selected reference regimen. It is fully acknowledged that safety data may be rather limited prior to the conduct of the first confirmatory trial, but main toxicities should normally have been identified and this should be sufficient for a rough estimate of the expected relative toxicity of the experimental regimen compared with alternative reference regimens.

Three categories are used in this document: Reduced or similar toxicity, increased toxicity and major increase in toxicity. No precise definition is given here due to heterogeneity of the conditions. "Major increase in toxicity", however, in most cases refers to a fear that the experimental regimen might be associated with an increase in treatment related deaths, irreversible adverse events with a long-term impact on QoL, or severe impairment to patient condition. Other issues to take into account include risk for secondary tumours. This categorisation is mainly meant for guidance in the planning of confirmatory studies and in order to provide advice on regulatory expectations with respect to study outcome measures in order to enable a proper benefit – risk assessment.

7.1. Design

7.1.1. Patient population

With respect to diagnosis, criteria for initiation of treatment, eligibility, response criteria and choice of reference therapy, a justification based on scientific evidence and/or generally acknowledged and

updated treatment guidelines are expected. While this is true in general, it is also expected that the exploratory studies through the judicious use of biomarkers provide guidance with respect to selection of patients in order to optimise benefit – risk, whether patient selection is in need for confirmation or not, in the planned phase III trials.

There is a general wish to reduce heterogeneity of study populations (performance status, co-morbidity, organ dysfunction, etc.) in order to increase the ability of the study to detect differences between study arms. This has to be balanced against the availability of patients for inclusion and the wish to enrol a clinically representative selection of patients. Therefore investigators should normally be encouraged to include patient's representative of those likely to be treated with the experimental compound in clinical practice. Restrictions as regards, e.g. performance status should be reflected in the SPC. With respect to studies with a non-inferiority efficacy objective, please refer to 7.7.3.

Patients are expected to be characterised by relevant tumour parameters, e.g. stage, grade, target expression, other biomarkers of importance for prognosis and/or tumour sensitivity, prior therapy (responsive/ resistant/refractory as appropriate), as well as performance status, co-morbidity, organ dysfunction, etc. Stratification based on important and well established prognostic covariates should be considered. In case adjusted analyses are to be undertaken for covariates other than those used for stratification, these factors should be pre-specified in the protocol or the statistical analysis plan (CPMP/EWP/2863/99).

If exploratory studies provide a basis for including/excluding certain patients based on tumour phenotype/genotype, this will be reflected in the labelling. As a corollary, if patients with tumours not expressing the target for activity are eligible, a restricted labelling may still be appropriate if it has not been demonstrated, e.g. by subgroup analyses, that target expression is irrelevant for anti-tumour activity.

If it is expected that a biomarker defining eligibility to the trial will be assessed locally or regionally in clinical practise, it is recommended that this is done also for the trial, complemented with central assessment of the biomarker to make feasible sensitivity analyses, etc.

As some of the conditions are rare, it is understood that the Sponsor might wish to define the target population using alternative criteria to those commonly employed. For example, in studies investigating the activity of a compound targeting a specific, molecularly well-defined structure assumed to be pivotal for the condition(s), it might be possible to enrol patients with formally different histological diagnosis, but expressing this target.

The pivotal role of the target in different histological diagnoses, however, must be demonstrated. This should be addressed in clinical studies, but it is accepted that formal testing with adequate statistical power of such a hypothesis cannot always be done. Possible consequences with respect to selection of proper reference therapy(ies) must be considered and the study should be designed so that it is possible, based on all available evidence, including non-clinical and pharmacological data, to conclude on the benefit – risk in the different subgroups of patients for which a claim is to be made. Prior to the initiation of confirmatory studies using non-conventional criteria for eligibility, EU scientific advice should be sought.

Some possible target indications comprise very small groups of patients, so small that “exceptional circumstances” might apply. Unless the target for activity is expressed only in these rare conditions, Sponsors are in general advised to undertake studies in these small patient groups in parallel to or when benefit – risk is established in indications allowing a more comprehensive evaluation, especially with respect to safety.

7.1.2. Reference therapy

The choice of reference regimen should be justified and normally this regimen should be selected from best available, evidence-based therapeutic options. In this context, “best available, evidence-based” should be read as a widely used, but not necessarily licensed regimen with a favourable benefit-risk convincingly documented through randomised trials and considered at least as good from a benefit/risk perspective as alternative, treatment options.

It is acknowledged that there are different, region-preferred standards. For superiority studies (test vs. reference) this should normally not constitute a problem as long as the reference is evidence-based as defined above. For add-on studies (reference + test vs. reference), it might also be possible to use a few, region-preferred references. Here a convincing clinical/pharmacological justification is needed, and EU scientific advice is recommended. Whenever more than one reference regimen is used, stratification is recommended and the overall superiority results should not be driven by the inferior results of one reference regimen.

If the aim is to demonstrate non-inferior efficacy, the selected reference regimen must enable a proper definition of the non-inferiority margin. In most cases, this would require that randomized well-controlled studies have shown the superiority of the selected reference vs. control. Please also refer to 7.6.3.

Amongst best available references, regimens with similar cycle lengths should be prioritised as it facilitates the identical scheduling of tumour assessments. If the objective is not to improve tolerability and toxicity, a regimen with similar expected toxicity to the experimental regimen is also preferred. This might also make the conduct of the study under double-blind conditions possible, a design recommended whenever adverse reactions do not make attempts to blind the study futile. In add-on studies (to an active reference or BSC), placebo is also recommended whenever meaningful.

In some cases there is no well documented reference regimen, even though patients in clinical practice are treated with certain regimens. Even though BSC is acceptable in these cases, an active comparator, documented e.g. in terms of response rate, is often preferable. If a single reference regimen cannot be defined, investigator’s best choice is an option. In these cases reference regimens with low toxicity are favoured and superiority in terms of patient relevant endpoints should be demonstrated.

The absence of evidence-based therapies often refers to patients who have failed several lines of therapy. In this situation, it might be more informative and also easier to obtain the data needed for marketing authorisation based on a properly conducted randomised study in less advanced patients, supported by “salvage” single arm studies, compared with conducting a last line, randomised BSC/investigator’s best choice comparative study.

7.1.2.1. *Single agent and combination therapies*

Whether the experimental agent is used as a single agent or in combination, the experimental regimen should be compared with the “best available” comparator again referring to benefit/risk, not only to efficacy.

If the experimental agent (A) is added to an established regimen (B), superiority of AB vs. B should be demonstrated and benefit-risk should be shown to be favourable. A discussion is expected based on available data as regards dose intensity of B and benefit risk. Traditionally, this type of studies does not include an A alone third arm, but this should be justified based on available exploratory study data.

In case of substitution studies, i.e. studies where a component (C) of an established regimen (BC) is replaced with an experimental agent (A) and if non-inferiority (BC vs. BA) is the aim, the contribution of C to the activity of BC has to be well defined (CPMP/EWP/2158/99).

Uncommonly, an entirely new combination AB is tested against a reference regimen. In these cases, solid non-clinical and clinical phase I/II data should support the need for both components in the experimental regimen.

7.1.3. Cross-over

In order to enable a qualified benefit – risk assessment, cross-over at time of progression should be undertaken only when detrimental effects on OS have been excluded (see Appendix 1).

7.1.4. Randomisation and blinding

Randomisation and stratification should adhere to the general principles laid down in current guidelines (CPMP/ICH/363/96). In many cases, a double-blind design is no option due to obvious differences in toxicity between study regimens or due to safety concerns. If the study has to be conducted open label, this has implications with respect to choice of study endpoints, independent review, conduct of sensitivity analyses and other measures to be undertaken to limit potential bias related to the open-label nature of the trial.

7.1.5. Endpoints

Confirmatory trials should demonstrate that the investigational product provides clinical benefit. There should thus be sufficient evidence available demonstrating that the chosen primary endpoint can provide a valid and reliable measure of clinical benefit in the patient population described by the inclusion criteria. In the following, superiority trials are the focus of the discussion.

Acceptable primary endpoints include cure rate, OS and PFS/DFS. Convincingly demonstrated favourable effects on survival are, from both a clinical and methodological perspective, the most persuasive outcome of a clinical trial. Prolonged PFS/DFS as such, however, is considered to be of benefit to the patient. The choice of primary endpoint should be guided by the relative toxicity of the experimental therapy, but e.g. expected survival after progression, available next-line therapies and the prevalence of the condition must also be taken into account. Irrespective of chosen primary endpoint, it is emphasised that it is the magnitude of the treatment effect on all relevant outcome measures that forms the basis in the benefit – risk assessment.

If PFS/DFS is the selected primary endpoint, OS should be reported as a secondary and *vice versa*.

When OS is reported as secondary endpoint, the estimated treatment effect on OS should ensure that there are no relevant negative effects on this endpoint, in most cases by showing trends towards superiority. In situations where there is a large effect on PFS, or if there is a long expected survival after progression, and/or a clearly favourable safety profile, precise estimates of OS may not be needed for approval.

When OS is reported as primary endpoint, consistency is expected as regards effects on PFS. If foreseen not to be the case, e.g. in case of certain immune modulating therapies, this should be made clear already in the study protocol.

For some conditions, events of progression will be observed at a slow rate making frequent assessments of events of progression a burden to the patients. Event rate at a pre-specified and justified fixed point in time might be used as primary outcome measure in these cases. When event rate at a single point in time is selected for the primary analysis, it is in most cases recommended that all patients should have been on study for that period of time. PFS, in a time to event analysis, and as assessed by the investigator should be reported as a secondary endpoint when a fixed time-point assessment is used as primary outcome measure.

For further methodological guidance as regards PFS, please refer to appendix 1.

It should be noticed that it is expected that the tumour's drug resistance profile is affected by therapy. This might be of relevance for the activity of next-line therapies. This is most obvious if maintenance/prolonged therapy is compared with no treatment or placebo such as in areas where a fixed number of cycles is the standard, for example, first-line ovarian cancer, NSCLC and some haematological conditions. The consequences of progression **on** maintenance therapy, signifying resistance at least to the maintenance regimen, might thus differ from progression **off** therapy. In principle, this applies to all comparisons, i.e. the degree of cross resistance as regards next-line therapy might differ between experimental and control regimens.

From a regulatory perspective, this concern has mainly been emphasised in settings where a new concept is introduced such as maintenance therapy or an increased number of "induction" cycles. If possible, these studies should therefore be designed with the aim to document patient benefit in terms of survival. If non-feasible, endpoints such as PFS on next-line therapy (PFS2) should be determined (see Appendix 1). This should ideally be done within the study so that agreed next line therapy(ies) is used after progression in the control and maintenance arms. In order to capture possible negative effects on next-line therapy and to outbalance tolerability and toxicity concerns related to maintenance therapy, it is expected that time from randomisation to PFS2 in the experimental arm is sufficiently superior to time from randomisation to PFS2 in the control arm. As the regulatory experience is limited and as methodological issues are foreseeable, EU scientific advice should be considered.

If the experimental compound used for maintenance therapy can be used as single agent also at time of recurrence, it is recommended that early treatment, i.e. maintenance, is compared with deferred therapy, i.e. treatment at time of progression.

It is accepted that it may not be feasible to define next-line therapy within the study protocol and to follow patients with scheduled assessments until PFS2. Time on next-line therapy might in these cases be used as a proxy for PFS2. The likely increased variability in the assessment of "PFS2" will be taken into account in the comparison $PFS2_{\text{control}}$ vs. $PFS2_{\text{exp}}$

It is also acknowledged that the choice of next-line therapy might reflect e.g. the patient's performance status at time of progression. As this is of relevance also for clinical practice, it is recommended that time on next-line therapy are captured in most studies, i.e. not only in studies introducing new concepts such as maintenance therapy. In these cases it might be informative if the CRF captures reasons for selecting a certain next line therapy.

A discussion on data maturity is warranted in all these cases as it is expected that, in general, early progression on or off therapy is related to more aggressive disease, i.e. biasing early PFS2 results in favour of the arm showing inferior PFS1 results.

Alternative primary endpoints, such as TTP or time to treatment failure (TTF) might uncommonly be appropriate. This has to be fully justified.

In patients with tumour-related symptoms at base line, symptom control, if related to anti-tumour effects, is a valid measure of therapeutic activity and may serve as primary endpoint in late line therapy studies, provided that sources of possible bias can be minimised. In certain cases, time to symptomatic tumour progression may also be an adequate primary measure of patient benefit.

There are also examples where tumour response-related activities, e.g. limb-saving surgery may be reasonable primary measures of patient benefit. Analyses of location- or cause-specific events, however, should in general be avoided as the focus may be drawn away from the main objective, namely the overall success of the treatment strategy in question.

Biomarkers convincingly demonstrated to reflect tumour burden can be used, in combination with other measures of tumour burden, to define tumour response and progression, an example being multiple myeloma and the M-component. For new classes of compounds, however, it has to be demonstrated that the marker is a valid measure of tumour burden and that no bias in the assessment is introduced, e.g. through differential suppression of the tumour marker.

7.1.5.1. Secondary endpoints and exploratory analyses

Irrespective of the choice of primary endpoint OS or PFS, ORR and rate of tumour stabilisation for, e.g. 3 or 6 months should be reported. Especially in the palliative setting, HRQoL/PRO using generally accepted instruments might be informative (Appendix 2)

7.2. Treatment administered with curative intent

The ultimate aim of developing new therapies, e.g., in patients with high grade lymphoma, germ cell tumours or in the adjuvant setting, is to improve cure rate and survival or to relevantly decrease toxicity without loss of efficacy. Nevertheless, in some cases and due to the complexity of administered therapies, e.g. in AML, the impact of a relevantly active experimental compound on these endpoints may be hard to demonstrate.

It is foreseen that the experimental compound rarely will be used as single agent therapy, but will be used as add-on to an established, perhaps modified regimen, or as substitution for a compound being part of the established regimen. In this context, maintenance therapy may be regarded as add-on therapy if maintenance therapy is considered non-established.

In the treatment of acute leukaemia, lack of achievement of CR, relapse and death without relapse are counted as events in an EFS analysis. Those patients who did not reach CR during the pre-specified induction phase will be considered as having an event at time 0.

In case EFS is found to be a justified primary endpoint, it is of importance that study data are analysed only when sufficiently mature, i.e. when it is foreseen that the EFS plateau is stable or when additional disease recurrence is rare.

In patients with high grade lymphoma or solid tumours, PFS may be used as outcome measure. Not achieving at least PR after a defined period/number of cycles may be regarded as treatment failure in some protocols and only those achieving at least PR continue on therapy. In the primary analysis it is recommended that patients not reaching PR are followed off or on next-line therapy until an event of progression or death is reached.

When improved cure rate is the objective of therapy, it is advised that disease-free survival at a pre-specified time point is used as outcome measure (see above with respect to timing).

7.2.1. Reduced or similar toxicity expected

In most cases, a substitution design is foreseen, meaning that A in an established regimen (AB) is replaced with the experimental agent X (XB). From a regulatory perspective, a non-inferiority design is acceptable and in most cases EFS or PFS, as appropriate, are acceptable primary endpoints.

In cases where induction is followed by consolidation and/or maintenance therapy, confounding effects of therapies administered after the end of experimental therapy may make endpoints other than PFS or EFS more appropriate. This means that CR (and CR + PR, if specifically justified) after end of experimental therapy could be an acceptable primary endpoint when further therapy is scheduled. In these cases, the possible influence of the experimental compound on the activity of consolidation

therapy should always be addressed and outcomes with respect to CR should be supported by EFS or PFS data.

It is recommended that CR is defined according to established clinical criteria, but supportive evidence in terms of Minimal Residual Disease (MRD) as defined, e.g. by molecular criteria should be sought when applicable. As for other biomarkers, intra- and inter- laboratory variability should be minimised through standardisation.

7.2.2. Increased toxicity expected

Substitution or add-on designs may apply. In most cases, superiority in terms of EFS, PFS, or OS as appropriate, should be demonstrated and the benefit in terms of prolonged time to event should be sufficiently large to balance increased toxicity.

A major increase in CR after induction therapy associated with trends in PFS or EFS, and survival, however, might be sufficient if scheduled treatments administered after the end of the experimental therapy are likely to confound overall outcome. This is of special relevance if the target population is small.

7.2.3. Major increase in toxicity expected

The aim should be to demonstrate increased cure rate or improved OS. In some cases, such as in small study populations, a major increase in EFS or PFS, as appropriate and supportive data compatible with a favourable trend on survival might be sufficient.

7.3. Treatment administered with the intent to achieve long-term disease control

Typical conditions include early lines of therapy in advanced breast cancer, colorectal cancer, low-grade lymphomas and the chronic leukaemias for which established reference therapies are available and next-line treatment options are likely to be meaningfully efficacious.

7.3.1. Reduced or similar toxicity expected

Substitution or single agent studies are foreseen. From a regulatory perspective, a non-inferiority design is acceptable and PFS is considered an appropriate primary endpoint. In case of relevantly reduced toxicity, mature survival data may be submitted post licensure if justified by study data.

7.3.2. Increased toxicity expected

The aim should be to demonstrate superiority at least in terms of PFS.

Survival data should be made available at the time of submission. It is acknowledged that mature survival data cannot be expected in all cases, though a justification explaining why this is the case should be provided. Post approval follow-up with respect to survival is expected in these cases. If absence of an increase in treatment-related mortality is not established with reasonable certainty, mature survival data should be available for the assessment of benefit – risk prior to licensure.

It is acknowledged that alternative endpoints may be more appropriate in certain situations, e.g. when maintenance therapy is investigated in areas where this has not established (Endpoints, 7.1.5). The aim may also be to enable a long treatment-free interval after intense induction therapy.

7.3.3. Major increase in toxicity expected

The principal objective should be to demonstrate improved survival.

In individual cases this might be non-achievable due to expected good prognosis with respect to survival and availability of several active next-line regimens, including experimental therapies, at the time of disease progression and a small target population. If PFS is the selected primary endpoint for the study, this requires a thorough justification. A careful discussion at the planning stage is also needed for the assessment of possibly therapy-related fatalities. Even though only a major benefit in terms of PFS prolongation would be acceptable, whenever possible the number of patients included should be sufficient to obtain an estimate on overall survival where a trend in a favourable direction is expected.

7.4. Palliative therapy

This mainly refers to last line settings where the prognosis for survival is poor and where it might be problematic to identify sufficiently documented reference therapies. In other cases, patients are considered not suitable for intensive, potentially curative therapy as defined by clear and as far as possible unambiguous criteria.

In cases where there is no established reference therapy, investigator's best choice or BSC with or without placebo are acceptable.

In a study conducted with BSC as reference therapy, the objective should be to demonstrate prolonged OS and/or globally improved symptom control or HRQoL. The latter requires that all efforts are undertaken to reduce possible bias (Appendix 2). Irrespective of aim, studies in this population requires that the treatment is well tolerated.

If the reference regimen is known to be active, but not established, superiority in terms of PFS might be acceptable. In these cases, the following will be taken into account in the benefit – risk assessment: the evidence showing activity of the reference therapy, the magnitude of the PFS benefit over the reference regimen, the tolerability/toxicity profiles, survival after progression and the prevalence of the condition.

It is acknowledged that patients may be considered suitable only for palliative therapy at baseline due to, e.g. poor performance status, but may respond so well that further therapy can be administered with curative intent, including, e.g. reduced intensity HSCT. How to handle these patients should be defined in the analysis plan.

7.5. Special considerations

7.5.1. Haematopoietic stem cell transplantation, methodological considerations

If allogeneic haematopoietic stem cell transplantation (HSCT) is a foreseeable treatment option, it is of importance to define how transplantation should be handled in the analysis plan. It is fully acknowledged that criteria for HSCT (e.g. patient eligibility, HLA matching, conditioning regimen, graft versus host disease prevention, etc) vary between institutions and regions. Nevertheless, these criteria should be defined as far as possible in the protocol and reasons for performing or not performing HSCT should be captured by the CRF.

Even though transplant related mortality is an issue and long-term benefit needs prolonged follow-up, it is normally expected that patients undergoing HSCT are followed for OS and EFS as randomised. Patients may be censored at time of conditioning for HSCT as a sensitivity analysis.

As treatment administered prior to transplantation might affect outcome of HSCT, proportion of patients undergoing HSCT is not considered to be a suitable primary outcome measure even if all patients responding sufficiently well to treatment are scheduled for transplantation.

Autologous stem cell transplantation constitutes less of a concern from an assessment perspective and may be viewed as intensified consolidation therapy where the consequences on short-term mortality and possible long-term benefit are less pronounced than after HSCT. Nevertheless, heterogeneity in the conduct of autologous transplantation should be avoided as far as possible, and censoring should not be undertaken.

With respect to drug development specifically in relation to HSCT, please refer to Appendix 4.

7.5.2. (Neo)adjuvant therapy

In the adjuvant setting, the ultimate aim is to increase cure rate. While effects on DFS are considered relevant to the individual patient, it is of importance to consider in the planning of the study whether it is at all possible to demonstrate a favourable effect on cure rate, i.e. in analyses conducted when recurrence rates have reached an apparent plateau.

As the use of adjuvant therapy may limit therapeutic options at time of recurrence, OS data should be reported. For established areas of adjuvant therapy, e.g. breast and colorectal cancer, and if benefit-risk is considered favourable for the experimental regimen based on DFS and available safety and survival data, including PFS on next-line therapy following recurrence of the disease, mature survival data may be reported post-licensing. In some cases and due to major toxicity concerns, favourable effects on OS have to be demonstrated.

The objectives of neoadjuvant therapy may include improved overall outcome (OS, DFS/PFS), enabling surgery and organ preservation (e.g. more conservative surgery). If organ preservation is the main objective, at least non-inferior DFS/PFS should be documented. As for adjuvant therapy, a defined number of cycles is frequently administered. Pending on the objectives of the study it is accepted that treatment is withdrawn if tumour shrinkage is not observed after a defined treatment period.

When pathological CR at time of surgery is reported as secondary endpoint, patients withdrawn should be considered as non-responders.

7.5.3. Drug resistance modifiers, chemoprotective agents and radio/chemo sensitizers

In principle, the design of confirmatory studies for experimental drug resistance modifying agents and radio/chemo sensitizers (A) is straight forward; AB should be demonstrated to be more active than an established regimen (B) in terms of anti-tumour activity and the benefit – risk for the combination should be shown to be favourable. If there are PK interactions, or dynamic interactions not related to anti-tumour activity, dose adjustments of B in the combination arm might be needed in order to make the comparison AB vs. B at similar overall toxicity. If the full effects of the PK interaction is captured by changes in the plasma levels of B (e.g. no changes in distribution), however, dose adjustments of B in order to compare AB vs. B at similar exposure of B is preferred.

For a chemoprotective agent, it has to be shown that normal tissues are more protected from toxicity than tumour tissue. For most cytotoxic compounds, it is, however, easier to detect dose-related differences in toxicity than in efficacy. This means that in many cases very large studies are needed with tight confidence intervals around measures of anti-tumour activity in order to prove that normal tissue protection is achieved without loss of anti-tumour activity. Co-primary endpoints are thus needed, testing the hypotheses of improved safety and non-inferior anti-tumour activity. In some cases, it might actually be easier to convincingly demonstrate differential tissue protection by increasing the dose of the cytotoxic compound in the experimental arm aiming to show enhanced anti-tumour activity without increased toxicity.

However, if it can be shown conclusively that there is no PK interaction and that the chemoprotective compound cannot interact with the tumour, e.g. by absence of target in tumour cells, it might be acceptable only to show reduced toxicity without formal non-inferiority testing of tumour protection.

7.5.4. Tumour Prevention

Regulatory experience is limited, but conceptually the situation is rather similar to the adjuvant setting. Thus individuals at risk should be defined so that the observed risk reduction in tumour incidence outweighs the side effects of therapy. As tumour prevention may select for tumours with altered biological behaviour, comparative data on tumour pheno/genotype are expected and data on tumour response to therapy or OS may be needed. In the planning of these studies, regulatory scientific advice is recommended.

7.6. Methodological considerations

Frequently, only one single study is foreseen for a specific indication. Licensing based on one pivotal study, however, requires demonstration of efficacy at levels beyond standard criteria for statistical significance (CPMP/EWP/2330/99). This is of special relevance in non-inferiority trials, in trials with PFS as primary endpoint and in a comparison with BSC/investigator's best choice. It is acknowledged that supportive evidence from confirmatory studies conducted in other indications should be taken into account in the assessment. The supportive value of these studies might vary and a discussion is expected as regards the relevance of these findings in relation to the application for the new indication.

7.6.1. Adaptive Design

If a phase II/III study is designed only to address a single and non-complex question in phase II of the trial, such as proper dose for the confirmatory stage, adaptive design might increase the efficiency of drug development (CHMP/EWP/2459/02).

Whenever more complex issues are to be addressed, e.g. involving defining the proper target population, or multiple issues, e.g. sample size re-estimation and cut-offs for biomarker positive tumour samples, etc. it is questioned whether adaptive design approaches are advantageous and scientific advice should be considered. The need for independent supportive efficacy/safety studies as part of the application for marketing authorisation should also be considered (CPMP/EWP/2330/99).

7.6.2. Interim analyses

Interim analyses are frequently undertaken in Phase III trials, but early stopping whether for futility or superiority is a sensitive issue. Early stopping for superiority requires that the treatment effect in patients with rapidly progressing tumours ("early events") is similar to that in less aggressive tumours ("late events") in the absence of data actually demonstrating that this is the case.

If a clear majority of the total number of expected events in the long term has been observed and a difference has been documented, this is normally accepted as an indicator that the study is reasonably mature and that the study results will remain stable over prolonged follow-up. The interpretation of interim analyses conducted on a less mature data set is problematic.

In cases where the treatment effect has been underestimated in the planning of the study, this may create a dilemma if statistically convincing effects in terms of overall survival have been demonstrated before a representative and mature dataset is available. Other monitoring committee decisions might be investigated in this instance such as restricting the continuation of the trial to the under-represented subsets to which the observed effect cannot be extrapolated. Analyses according to

stratification factors of major importance for prognosis might provide insights as well as similar analyses with respect to PFS.

In general, interim analyses based on PFS data other than for futility are not encouraged (Appendix 1).

7.6.3. Time to event analyses and assessment of response and progression

For studies with PFS/DFS as primary endpoint, symmetry with respect to imaging and study visits is pivotal and adherence to protocol-defined schedules is essential and deviations should be reported (Appendix 1).

As discussed above (Exploratory trials with time-related endpoints), a comparison in terms of PFS between a predominantly tumour shrinking compound and a predominantly growth inhibiting compound may “favour” the latter compound with respect to tumour burden at time of progression. Until now, there is no regulatory experience with respect to comparisons with clearly discordant outcomes in terms of ORR and PFS and there are no established ways to adjust for this. If exploratory studies indicate that this might become the case, alternative endpoints such as OS should be considered.

Differences in mode of action between the experimental and reference therapy might generate problems in relation to measurements of tumour burden and anti-tumour activity, one example being early tumour swelling as discussed previously. Whenever such problems are foreseen, which may require deviation from standard approaches (RECIST, WHO), it is recommended that agreement is reached with regulatory agencies prior to the initiation of pivotal trials. Similarly, if tumour assessment techniques cannot be used that allow for independent adjudication, it is advisable to discuss available alternatives with regulatory agencies.

Pseudo-response should always be considered a possibility when tumour related oedema is an issue such as in high grade gliomas. Updated response and progression criteria taking this into account should be applied when available. If such criteria has not yet been established, scientific advice is recommended in order to discuss alternative ways forward.

7.6.4. Non-inferiority studies

Guidance of design, conduct and analysis of non-inferiority studies is given in other regulatory guidance documents (Choice of a Non-Inferiority Margin CPMP/EWP/2158/99), but some topics deserve particular attention in the oncology setting. For a PFS endpoint, which can be considered a composite endpoint, the discussion of a non-inferiority margin should consider the effect of the reference treatment overall but inference should also include a discussion on each type of events (death, new metastases, progression of target lesions, clinical progression) including description of the effect of the reference regimen on each component when available. If differences in the profiles of progressive disease might be expected, this should be accounted for in the planning stage with a suitably conservative margin and appropriate sample size to obtain the required number of events for reliable inference.

Given the importance of study sensitivity (i.e. the ability of a trial to detect differences) for the assessment of non-inferiority trials, where similar activity is assumed for test and reference, it is of importance to plan in advance for a subgroup analysis, e.g. excluding patients with poor prognostic factors at baseline such as poor PS, co-morbidities, etc. as in these patients it might be harder to detect a difference in activity between treatment regimens, if there were one. Similarly a per protocol analysis set should be defined so that protocol violations, compliance problems, etc. do not reduce the possibility to detect a difference. These analyses are expected to be undertaken with the aim to show consistency.

7.6.5. Analyses based on a grouping of patients on an outcome of treatment

Comparisons of time-to-event variables (like OS, or PFS) by grouping patients on a post-randomisation outcome of treatment are problematic. Since outcomes like tumour response, dose intensity, toxicity, or compliance represent an interaction between therapy, patient and tumour the contribution of therapy cannot be disentangled. Nevertheless, certain unexpected outcomes such as clearly improved survival despite dose-reduction due to toxicity, or absence of prolonged survival in responding patients might be informative. A search for unexpected findings constitutes a rationale for conducting these exploratory analyses.

Response duration comparing groups of patient on different therapies may be regarded as informative. Data should be reported with confidence intervals for the individual study arms, but significance testing comparing duration of response between study arms should not be undertaken as the comparison refers to groups that are not fully randomised. "Time in response" where patients without response are assigned a duration of zero enables a statistical comparison between study groups.

7.6.6. Studies in small study populations, very rare tumours

For some truly rare tumours or very narrow indications, whether due to tumour phenotype or restrictions related to target expression, it is simply not possible to recruit a sufficiently large number of patients to conduct reasonably powered, randomised studies in order to detect clearly relevant differences in anti-tumour activity. In some cases a small, randomised, reference controlled study is the best option, in other cases a within-patient TTP/PFS analysis (or the combination) might be a better alternative. In the latter case, TTP on last prior therapy is compared with time to progression or death on the experimental therapy. This would require that the clinical appropriateness of the last administered therapy prior to study therapy and progression on prior therapy is independently adjudicated and that the study protocol clearly defines the proper conditions for the analysis. Superiority should be demonstrated.

Problems related to studies in small populations are further discussed in the Guideline on clinical trials in small populations (CPMP/EWP/83561/2005). In these small target populations all evidence with respect to efficacy and safety must be taken into account. This encompasses clinical response rate, duration of response as well as outcome measures such as HSCT rate, use of minimal residual disease (MRD) to define response rate and recurrence of disease, as appropriate. Mature time to event endpoints such as PFS and OS should be reported even though it is acknowledged that formal statistical significance cannot always be expected, even if the experimental compound is relevantly more efficacious.

As there is no general solution to the problem of how to document benefit – risk in these cases, scientific advice is recommended.

7.6.7. Use of external control

The use of external control (including historical control) is discussed in ICH Topic E10 (CHMP/ICH/364/96) and it is concluded that "the inability to control bias restricts use of the external control design to situations where the treatment effect is dramatic and the usual course of the disease highly predictable".

Dramatic effects are uncommonly documented in the treatment of malignancies, but it is acknowledged that such effects, obvious to any qualified observer, are seen occasionally. In these cases, prospective confirmation in randomized, reference-controlled studies is not only unacceptable to investigators, patients and ethics committees, but also unnecessary.

7.7. Special populations

7.7.1. Elderly and frail patients

Whenever elderly patients are expected to be treated with the new medicinal product in clinical practise, the clinical studies program should enrol a sufficiently large number of elderly, including those with co-morbidities, to enable a benefit – risk assessment. It is acknowledged that for some products, the safety of the drug needs to be established in otherwise healthy patients prior to enrolment of less fit elderly in confirmatory studies, but a justification is expected in these cases. Of note, eligibility criteria per se is frequently not the hurdle, in order to accomplish a fair representation of elderly, investigators need specific encouragement and support to enrol these patients.

It is expected that all reasonable efforts are undertaken to provide informative data in the MAA, however, if benefit – risk cannot be assessed with reasonable certainty in elderly patients or those with prevalent co-morbidities in the target population, this should be reflected in the labelling and post approval studies may need to be undertaken. In this context it is noticed that also well-planned cohort studies may provide valuable information.

Data from elderly patients should be available for pharmacokinetic analyses, e.g. as part of population pharmacokinetic analyses. Description of the safety profile should include aspects of severity of the adverse events profile and consequences, e.g. dose reduction, dose delay or initiation of concomitant treatment. An evaluation of the consistency of treatment effects and safety profile in elderly population, including age groups as appropriate, with the younger population(s) is expected.

Some compounds may be specifically suitable for the treatment of elderly, e.g. due to PK properties such as low sensitivity to impaired organ function. In these cases, dedicated studies in the elderly are encouraged. It is acknowledged that it may be hard to identify appropriate reference therapies in some of these cases and that other outcome measures than PFS/OS might become more relevant. In these cases it is advisable to seek regulatory agreement on the development program.

Frail patients, whether elderly or not, with clearly impaired performance status (PS) constitute a vulnerable group of patients rarely included in conventional studies. Clinical studies in this group of patients are supported from a regulatory perspective.

7.7.2. Children

See *Addendum* (CPMP/EWP/569/02 under revision).

7.7.3. Gender

For some tumours and/or therapies, a difference in antitumour activity related to gender has been reported. Where a priori it is likely that there may be a treatment by gender interaction, this should be taken into account in the design of the study. Otherwise it is expected that the proportion of females and males reflects the prevalence of the disease and that the sponsor provides exploratory subgroup analyses (efficacy and safety) by gender.

7.7.4. Patients with impaired organ function

Please refer to Section 4, Pharmacokinetics.

7.8. Safety

In addition to standard reporting of adverse events, it is expected that effects of preventive measures, such as anti-emetics or use of growth factors are reported. Acute, sub-acute, chronic and late toxicities

should be described. Safety in special populations, as detailed above, should be summarised from the full studies programme.

For common events, safety in relation to treatment cycle, first, second, third etc., is of value. Similarly, timing and duration, including grade, of some events such as diarrhoea, mucositis, or cytopenias should be reported.

Monitoring of frequency and type (viral, bacterial, fungal) of possible, probable or proven infections should be undertaken in patients undergoing more intensive cytotoxic/immunosuppressive therapy. For compounds known or suspected to cause long term immunodeficiency, monitoring for opportunistic infections for up to one year after the end of therapy should be considered.

Cumulative toxicity should always be investigated.

If cure is the objective, long term follow up for toxicity is highly relevant. Late toxicity includes secondary malignancies and certain organ toxicities (e.g. CNS, cardiovascular). The number of patients suffering from late toxicities may increase over time and is therefore an objective for post licensure pharmacovigilance activities.

As radiation therapy is a standard treatment option in malignant tumours, it is foreseeable that patients will be receiving radiation therapy, e.g. for symptom palliation, concomitantly with or in a time frame close to administration of the medicinal agent. Safety information on concomitant or sequential use of the medicinal agent with radiotherapy should therefore be collected throughout the entire study programme, including data on "radiation recall". Subjects requiring radiation therapy while enrolled in a trial of a novel agent or combination of agents should normally be withdrawn from study therapy. If safety and tolerability information with radiation is desired, well-designed dedicated studies should be conducted in the setting of concomitant radiation, with dosing schedules guided by preclinical toxicology studies.

In haematological malignancies, bone marrow failure is often a presenting symptom and is frequently aggravated by treatment. In these cases, dose reduction for haematological toxicity may not be indicated.

If the aims of the study include demonstration of improved safety, the protocol should specify how this should be accomplished. It is not acceptable to focus on one toxic effect only. The outcome measure(s) should provide unbiased information on overall toxicity and tolerability, perhaps in addition to a specific item such as neuropathy where a clinically relevant improvement is expected. As there is limited experience with this type of studies, EU regulatory advice should be considered.

Where appropriate, pharmacogenomics may be used to identify patients at increased risk for severe toxicities.

8. Definitions and Abbreviations

Chemoprotectant: A compound which counteracts the activity of anti-tumour compounds on normal tissue without (or clearly less) affecting the anti-tumour activity.

Chemotherapeutic (or drug resistance modifier): A compound without own anti-tumour activity which increases the activity through pharmacodynamic interaction with anti-tumour compound(s).

Cytostatic: Anticancer compound shown to inhibit cell division without direct effects on tumour cell viability in non-clinical studies.

Cytotoxic: Anticancer compounds inducing irreversible lethal lesions through interference with DNA replication, mitosis, etc. following short term exposure in non-clinical studies.

Data maturity: A clinical study is considered mature if the distribution of events over time (early – late) makes it feasible to estimate the treatment effect in the full study population. This refers to the assumption that there is a biological difference between e.g. tumours progressing early and late and that the treatment effect might differ. The number of late events should therefore be large enough for study data to be stable. In practice, if a treatment difference has been established and a clear majority of events expected over long term have occurred, the study may in most cases be regarded as “mature”.

Non-cytotoxic: Anticancer compounds not belonging to the class of cytotoxic compounds.

Primary (innate) resistance: Progression without prior objective response or growth inhibition.

Randomised phase II trial: Randomised exploratory study designed to provide data of importance for the design of Phase III confirmatory studies, e.g. with respect an estimate of the possible magnitude of the effect using a clinically relevant measure of activity and/or biomarkers.

Refractory: Progression on therapy or within a short period of time after last cycle of therapy.

Resistance: Progression within a defined timeframe after end of therapy.

Secondary resistance: Progression after documented objective response or period of growth inhibition.

Window of opportunity: Under certain well-defined conditions it is acceptable to conduct a clinical study with an experimental compound in settings (line of therapy, stage, etc.) where available data for this compound normally would be regarded as too limited. The conditions for conducting such a study must be set rigorously so that the interest of the patient is guaranteed. Circumstances to take into account include benefit-risk of available therapies, available safety/activity data for the experimental compound, tumour-related symptoms (in most cases absent), expected evolution of the disease if left untreated or treated with available therapies, ease of frequent monitoring of tumour evolution (including use of biomarkers), planned intervention post chemotherapy, etc.

ADCC: Antibody dependent cellular cytotoxicity

ANC: Absolute neutrophil count

BSA: Body surface area

BSC: Best supportive care – include antibiotics, nutritional support, correction of metabolic disorders, optimal symptom control and pain management (including radiotherapy), etc. but does not include tumour specific therapy

CBR: Clinical benefit response rate. CR or PR or prolonged SD. “Prolonged SD” is defined condition specific, for breast cancer normally ≥ 24 weeks.

CR: Complete response

CRF: Case report form

DFS: Disease-free survival (time from randomisation to recurrence or death from any cause)

DLT: Dose limiting toxicities

EFS: Event-free survival in this guideline refers to lack of achievement of CR, relapse and death without relapse are counted as events in an EFS analysis. Those patients who did not reach CR during the pre-specified induction phase will be considered as having an event at time 0.

HRQoL: Health related quality of life

MoAb: Monoclonal antibody

MTD: Maximum tolerated dose, often defined by dose-limiting toxicity occurring in at least 2 of 6 patients so that further dose-escalation is not undertaken.

ORR: Objective response rate (the proportion of patients in whom a CR or PR was observed)

OS: Overall survival (time from randomisation to death from any cause)

PD: Pharmacodynamics

PK: Pharmacokinetics

PR: Partial response

PRO: Patient reported outcome

PFS: Progression-free survival (time from randomisation to objective tumour progression or death from any cause)

PFS2: Time from randomisation to objective tumour progression on next-line treatment or death from any cause. In some cases, time on next line therapy may be used as proxy for PFS.

RP2D: Recommended phase 2 dose

SD: Stable disease

TTF: Time to treatment failure (time from randomisation to discontinuation of therapy for any reason including death, progression, toxicity or add-on of new anti-cancer therapy)

TTP: Time to tumour progression (time from randomisation to observed tumour progression, censoring for death not related to the underlying malignancy)