



# Odyssey Tritoon Owners Manual

**Jessica J Manson**



## **Odyssey Tritoon Owners Manual:**

*APOPTOSIS INDUCTION/SUPPRESSION: A FEASIBLE APPROACH FOR NATURAL PRODUCTS TO TREATMENT OF DISEASES, 2nd Edition* Hong Zhang, Hailing Xin, Wei Peng, Khalid Rahman, 2025-01-23 This Research Topic is part of a series See also Apoptosis Induction Suppression A Feasible Approach for Natural Products to Treatment of Diseases Volume II Apoptosis is generally recognized as a form of programmed cell death which is beneficial for normal cell development organ growth and tissue homeostasis in multicellular organisms In normal conditions millions of cells would indeed die and proliferate every day in the human body However an imbalance between cell death and proliferation can lead to some serious diseases Two different case scenarios can be distinguished 1 uncontrolled cell proliferation and insufficient cell apoptosis would lead to various cancer types and autoimmune diseases e g rheumatoid arthritis lupus erythematosus etc 2 excessive apoptosis in normal cells e g neural cells or cardiomyocytes would result in neurodegenerative diseases Alzheimer s disease Parkinson s disease and Huntington s disease and ischemia injuries myocardial infarction stroke etc respectively Natural products from plants animals microorganisms and minerals are potentially important resources in the context of drug discovery for various diseases Importantly increasing scientific evidence has suggested that apoptosis induction or suppression might be one of the predominant molecular mechanisms whereby natural products could be used to treat diseases especially cancer rheumatoid arthritis Alzheimer s disease Parkinson s disease stroke etc However not much is known about the detailed molecular mechanisms underlying apoptosis induction or suppression including signaling pathways novel and key pharmacological targets as well as the action of specific active substances extracted from plants etc In addition lots of active natural products based on apoptosis regulation have already received drug regulatory approvals e g taxol camptothecin and sinomenine and been used as clinical drugs to treat diseases furthermore there are many other natural products in the stages of the clinical investigations However the related advance and update of the current drug development correlated to apoptosis induction or suppression systematic reviews or meta analysis of these clinical drugs or candidate drugs in clinical research stage are insufficient This Research Topic will provide an academic platform to discuss how natural products can be used to treat several types of diseases via apoptosis induction suppression We invite authors to contribute original research and review articles testing the action of natural bioactive products on various diseases through apoptosis regulation including induction and suppression We aim to particularly focus on the recent advances in the curative properties of natural products on cancers tumors autoimmune diseases neurodegenerative diseases and cardiac diseases through apoptosis regulation and new natural bioactive agents for controlling diseases via regulating apoptosis Potential topics will include but won t be limited to the following 1 Advance in curative properties of natural products on diseases via apoptosis regulation 2 Advance and update of the current drug development correlated to apoptosis induction or suppression 3 Novel natural products with curative activities via apoptosis induction in particular for cancers tumors rheumatoid arthritis and

lupus erythematosus 4 Novel natural products with curative activities via apoptosis suppression in particular for neurodegenerative diseases Alzheimer s disease Parkinson s disease and Huntington s disease and ischemia injury myocardial infarction stroke 5 Novel signal molecules for the apoptosis related signal pathway 6 Systematic reviews or meta analysis of the approved natural drugs or candidate natural drugs in clinical research stage with induction or suppression of apoptosis The four pillars of best practice in ethnopharmacology With these guidelines we define in detail what constitutes best practice for manuscripts submitted to Frontiers in Pharmacology Section Ethnopharmacology They provide a basis for the peer review and build on the general requirements of Frontiers in Pharmacology 1 Pharmacology a The manuscript MS must report a substantive body of ethnopharmacological research to be considered as an independent addition to the literature In general we expect that such studies are based on local traditional uses of plants or other natural substances which need to be spelled out clearly b For pharmacological studies the model used must be one which is either generally accepted in the field as valid or a credible alternative whose general development and application in the reported instance has been justified Specifically antioxidant activity must be based on a pharmacologically relevant in vivo or cell based model Simple in silico and pharmacologically irrelevant assays for antioxidant activity e g the DPPH assay FRAP Ferric Reducing Ability of Plasma ABTS 2 2 azinobis 3 ethylbenzothiazoline 6 sulfonic acid are not acceptable as a main tool for assessing an extract or a compound for activity c Similarly simple screening for anti microbial effects of crude extracts is no longer state of the art Authors must follow the widely accepted standards for microbiological testing cf Cos et al 2006 Anti infective potential of natural products How to develop a stronger in vitro proof of concept Journal of Ethnopharmacology 106 290 302 and subsequent methods papers Such research is only meaningful if it contributes to our mechanistic understanding of anti microbial effects its specificity or identifies novel leads d The dose ranges must be therapeutically relevant While it will be impossible to define an exact cut off the literature in the field is now replete with studies which test extracts at implausibly high doses Single dose studies will only be of relevance in exceptional circumstances e g in case of specific complex pharmacological models And of course positive and negative controls must be included e In order to establish therapeutic benefits selectivity data are essential How specific is the effect Many compounds have non selective in vitro effects and research on common compounds must be justified in terms of the potential therapeutic benefits While such research may be relevant and have potential applications authors will need to assess the specificity of a single compound or an extract rich in a well studied compound like rutin curcumin or quercetin and provide evidence for the relevance and novelty of the approach f Docking studies must be justified with affinity experiments or other well established experimental methods to support a proposed mechanism of action Algorithmic docking studies will not be accepted these indicate if a compound will fit into a binding site but do not indicate the binding affinity or the ability to induce a conformational change 2 Composition a Botanical The identification of the study material must be described well All species are fully validated using Kew MPNS

portal or The Plant List initiative or Plants of the World Online Of course full botanical documentation is essential i.e. a voucher specimen deposited in a recognised herbarium A scan of the voucher is welcome as supplementary material and encourage authors to include the coordinates of the location where the material had been collected b Chemical The composition of the study material must be described in sufficient detail Chromatograms with a characterisation of the dominating compound(s) are preferable If preparations are used which are available commercially quality parameters provided in pharmacopoeia must be provided The material under study must be characterised using the methods of the relevant monograph If pure compounds are used sufficient information on the level of purity must be included Especially in in vitro models the authors must be confident that the compounds are stable under the conditions used for example they do not degrade due to high concentrations of DMSO A critical aspect that should be considered is how these assays and extraction protocols are linked to local and traditional uses In this way variables such as the solubility of the compound in the traditional preparation and in the analytical extraction protocol should be taken into consideration All chemical line structures must be drawn using an internationally accepted structure drawing programme must be consistent and if possible and relevant the stereochemistry needs to be given c Multiherbal preparations Very often multiherbal preparations are used Full information on their composition in terms of the botanical drugs species included and information on the rationale for studying this preparation needs to be included It is essential that in these cases sufficient details are provided on the botanical 2a and chemical 2b characterisation 3 Basic requirements and research ethics Frontiers has very well developed guidelines relating to ethical aspects of a MS Specifically for Frontiers in Pharmacology Ethnopharmacology the following key requirements are essential a The objectives of the research reported must be spelled out clearly and in detail All MS must critically assess the scientific basis of the work and provide meaningful conclusions which are based on a clear hypothesis research question as defined in the introduction Ethnopharmacological research must assess whether a compound or plant extract has a certain effect and it cannot be about confirming an extract's or compound's effects or efficacy b Research must add new and scientifically substantive knowledge to our understanding of the pharmacology and use of medicinal plants A key basis for this is a review of literature relevant to the pharmacological activity already reported on the species including possibly related taxa or compounds This must be up to date and clearly demonstrate the substantive addition to the literature the MS submitted represents Simply using advanced measurements techniques protocols reproducing previous studies of the same plant product will only be accepted in exceptional circumstances e.g. previously unknown highly active components are discovered c Compliance with all international ethical standards is essential In the context of ethnopharmacology the Convention on Biological Diversity and most recently the Nagoya Protocol are of particular relevance <https://www.cbd.int/abs> d Research in ethnopharmacology is based on local and traditional knowledge often passed on orally over generations Ultimately research in this field must therefore benefit those populations who are or were the

original keeper of this knowledge e The use of animals must be justified in the context of novelty see also part 1 It is ethically not acceptable to have yet another in vivo study on an already well studied species demonstrating some common activity e g an anti inflammatory effect studied in the rat paw edema The same is true for species which are chemically very similar and generally are rich in common ingredient to ones already studied pharmacologically Such studies must meet s the standards of rigor we expect in ethnopharmacology as defined in the Frontiers guidelines 4 Other specific requirements a Studies focusing on local and traditional uses of plants ethnopharmacological field studies must be based on substantial original data The relevance of the MS in the context of previous studies in the geographical region must be spelled out clearly and it must contribute to the understanding of the therapeutic uses of plant species and inform experimental or clinical studies This includes an adequate presentation and discussion of the data Also social science centered studies e g ethnobotanical studies or health system research of local and traditional medical systems are welcome This journal subscribes to the ConSEFS standards including any updates b In case of reviews we expect clearly defined scientific aims objectives a comprehensive critical and specific assessment of the relevant information linking local and other medical uses to the biomedical and bioscientific evidence Reviews need to define future research needs and priorities It is essential that the scientific quality of the original articles cited is assessed If pharmacological studies are reviewed particular attention must be paid to assessing the quality of the studies c Food plants are commonly reported to have pharmacological effects Frontiers in Ethnopharmacology focuses on therapeutic benefits of such species and not on the general food nutritional properties

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