medartis®

# Terms and Conditions

You are applying for a research grant ("Grant") to support your research project ("Project") at Medartis AG, Basel, Switzerland ("Medartis"). Please note that your application will be reviewed in the first line by our clinical research group as well as by other departments at Medartis' sole discretion.

In case of questions or uncertainties, feel free to contact us:

APTUS.research@medartis.com (clinical research upper and lower extremities)
MODUS.research@medartis.com (clinical research CMF)
general.research@medartis.com (other research)

By submitting your application, you accept the following terms and conditions:

1. Your Project will be evaluated for scientific and clinical merits, relevance, originality of hypothesis, as well as cost-effectiveness.

2. Medartis only supports Projects which are compliant with all applicable laws and regulations, in particular, but not limited to, the Declaration of Helsinki, ISO 14155 / ICH-GCP (in case of prospective clinical studies), as well as applicable data protection regulations (e.g. GDPR or HIPAA).

3. Grants will be awarded only to applicants affiliated with a hospital / academic institution. Grants will only be paid to accounts of hospitals / academic institutions. The account holder must be the hospital / academic institution at which the Project will be carried out.

4. Your application and any information related to your Project will be treated confidentially and in accordance with applicable data protection regulations.

5 . Within 3 months' time you will be notified about our decision or whether more information is required.

6. There is no obligation for Medartis to support your Project. The decision is at the sole discretion of Medartis and does not require justification. The decision is final and non-appealable.

7. In case your Project is accepted and support will be granted a cooperation agreement — to be signed by both parties — will be negotiated. This agreement compulsively includes the specification of deliverables by the applicant (e.g. detailed study protocol, ethics committee decision, GCP certificates of all co-applicants, reports, etc.), as well as the responsibilities of both parties.

8. Medartis requests the submission of supported Projects to a public database (e. g. clinicaltrials.gov).

9. Any unspent Grant has to be returned to Medartis.

10. Medartis requests the correct and complete disclosure and declaration of conflicts of interests.

11. Medartis does not intervene with or monitor the content of your work and will not act as a sponsor (as defined in the European MDR) for clinical research. The full responsibility for the design, execution and management of the Project, as well as for compliance with all regulatory and legal requirements for or related to the Project, remains with you. The role of Medartis is limited to providing support in order to promote the quality of research and of the resulting publications.

12. By submitting your application, you agree that Medartis has the right to use, at its own discretion, the results of your Project and any information arising from the scientific exchange between you and Medartis, e. g., in the process of development, to integrate it into the "Medartis Learning Management System" or for promotional activities.

13. Any potential contribution to your Project is not in any way connected to any of your past, present or future purchase, recommendation or use of Medartis services or products or their respective pricing. Scientific support is solely based on the merits of your project and not associated with or conditioned on any purchase or use of Medartis products.

14. By submitting and signing the application form you acknowledge to have read and understood the above terms and conditions and confirm that you will adhere to them.

These Terms and Conditions may be changed by Medartis at any time without notification. Please refer to this website for the actual version thereof.