**TraFo Cooperation Proposal**

# RESEARCHER DETAILS

## PROPOSAL TITLE:

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## PROPOSAL SUBMISSION DATE:

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## CONTACT DETAILS (Principal researcher):

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| --- | --- |
| NAME: |  |
| JOB TITLE: |  |
| INSTITUTION ADDRESS: |  |
| TEL: |  |
| E-MAIL: |  |
| INVOLVEMENT IN GBG STUDIES: |  |

## CONTACT DETAILS (statistician – if applicable\*):

\*Typically, GBG will do the statistics due to data protection issues. If an external statistician is explicitly needed, please give a rationale and add the contact details.

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| --- | --- |
| NAME: |  |
| JOB TITLE: |  |
| INSTITUTION ADDRESS: |  |
| TEL: |  |
| E-MAIL: |  |
| INVOLVEMENT IN GBG STUDIES: |  |

## CONTACT DETAILS (Other Collaborator):

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| --- | --- |
| NAME: |  |
| JOB TITLE: |  |
| INSTITUTION ADDRESS: |  |
| TEL: |  |
| E-MAIL: |  |
| INVOLVEMENT IN GBG STUDIES: |  |

## CONTACT DETAILS (2nd Other Collaborator):

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| --- | --- |
| NAME: |  |
| JOB TITLE: |  |
| INSTITUTION ADDRESS: |  |
| TEL: |  |
| E-MAIL: |  |
| INVOLVEMENT IN GBG STUDIES: |  |

# PRELIMINARY INFORMATION

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| Specify the overall aim of the project (concise) |
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| Provide background on the research hypothesis (max. 300 words, including a max. of 5 high impact references supporting the need of your work). |
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| Since the GBGs intention is to deliver high quality research for improvement of breast cancer therapy, what does your project contribute to the current research landscape? |
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| Which journal is planned for publication of your project (please specify also journal impact factor)? |
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| What resources are needed from GBG to support the project  |
| Is biomaterial needed?[ ]  yes [ ]  noWho will perform statistical analysis?[ ]  GBG [ ]  applicantPlease specify who will write the publication:Klicken Sie hier, um Text einzugeben.Comment: Klicken Sie hier, um Text einzugeben. |

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| Please specify the study and cohort needed for the project  |
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| What is the type and estimated quantity of samples that are required?  |
| Specify the type of samples and estimated quantity:[ ]  quantity Whole blood, specify collection time point: Collection time point[ ]  quantity Blood serum, specify collection time point: Collection time point[ ]  quantity Blood plasma, specify collection time point: Collection time point[ ]  quantity Fresh Frozen Tissue[ ]  quantity Tissue in RNAlater [ ]  quantity FFPE Tissue[ ]  quantity Pre-therapeutic cores[ ]  quantity Residual tumor tissue (surgery)[ ]  quantity Tumor recurrence (metastasis)[ ]  quantity TMAs[ ]  quantity Other, please specify: Please click to enter text |

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| Does the proposal require the use of the clinical data collected during the study? |
| [ ]  Yes [ ]  No | if yes, specify: |
| Clinical data collected during the trial[ ]  only baseline/patient characteristics[ ]  treatment data[ ]  safety / adverse event data[ ]  response data (pCR)[ ]  long-term survival data (DFS, OS …)[ ]  biomarkers, please specify: Please click to enter text. |
| Please justify the need of the data according to your hypothesis (statistical analysis plan):  |

# Milestones, Responsibilities and Funding

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| Please provide the main milestones of the project and responsibilities within your team |
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| Do you cooperate with others profit and/or non-profit organizations within the project? If yes, please specify. |
| [ ]  yes [ ]  noPlease specify. |

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| How is the project funded? Please specify (e. g. industry, foundation grants, institutional funds) |
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| Apart from the ethical approval at study submission, is there any additional ethics requirement for the project? |
| [ ]  Necessary and done[ ]  Necessary but not done yet[ ]  Not necessary, please explain: Please specify |