**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  no  Who will perform statistical analysis?  GBG  applicant  Please 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  no  Please 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 |