

## Novel targets for drug development



Join us in developing new therapeutic approaches for diseases with high unmet medical needs.

Apply for funding to promote your research and explore novel targets for drug development.

**Submission ends August 31st, 2017**

**SUBMIT PROPOSAL**



## WHAT WE ARE FUNDING

We are offering grants for researchers investigating novel drug targets in the fields of Oncology, Gynecology, Cardiology and Hematology to promote innovative therapeutic ideas from basic research into novel drugs.

We provide financial support to test your hypothesis, start or extend your present research activities.

### WHAT WE ARE FUNDING

### YOUR BENEFITS

**We are looking for novel targets for the indications of**

**APPLICATION PROCESS** ➤ Oncology, Gynecology, Cardiology and Hematology

### SUBMISSION

Oncology

Focus on oncogenic signaling.

Gynecology  
FAQ

Focus on novel treatment options for endometriosis (incl. adenomyosis, uterine endometriosis interna), uterine fibroids (uterine leiomyoma) and polycystic ovary syndrome (PCOS).

Contact Us

(/home/pharma/faq/index.php)

(scripts/components/forms/contact.php)

Pharmaceuticals

Cardiology ▼



Focus on novel approaches to the care of chronic and/or acute pulmonary hypertension & adjacent lung indications, atrial fibrillation, heart failure, acute coronary syndrome, ischemic stroke, peripheral arterial occlusive disease, acute lung injury/adult respiratory distress syndrome, cardiorenal syndrome and including kidney and lung diseases.



(<http://www.bayer.com/>)

#### Hematology

Focus on novel, innovative targets for hemophilia.

- Target must be a nucleic acid or a protein (e.g. an enzyme, a receptor) whose activity can be modified by a drug

The drug can be a small-molecular-weight chemical compound or a biological, such as an antibody or a recombinant protein.

Target should have shown to be effective/ mechanistically involved in the disease by relevant in vitro or in vivo models.

Target is disease-modifying and/or has a proven function in the pathophysiology of a disease.



## YOUR BENEFITS

### Financial Support

There are two types of grants, that will be allocated by Bayer depending on the target, the scientific data provided and the maturity of the proposal.

#### Support grants (€ 5,000 - €10,000)\*

For druggable targets that are at a very early stage of discovery.

#### Focus grants (€ 10,000 - €125,000)\*

For more mature ideas, e.g. to address specific aspects of a target as a first step towards transferring it to the drug discovery process.

\* The size of the individual grant will depend on the target specifics and the phase of development and validation.

#### WHAT WE ARE FUNDING

##### Conditions

- The decision on grant allocation is made at the sole and absolute discretion of Bayer.
- The applicant will retain ownership of any intellectual property he or she develops.
- Special conditions may apply for proprietary targets and the in-licensing of targets.

#### APPLICATION PROCESS

#### SUBMISSION

Grants may be complemented by

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FAQ

(</home/pharma/faq/index.php>)



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Pharmaceuticals



Small Molecule Tool  
Compounds





## Antibodies

Bayer's Antibody Discovery unit may be able to support your target validation work. Recombinant antibody engineering technology can be used to generate suitable antibodies for your validation needs.

Please specify which in vitro/in vivo models you'll be using and the requirements potential tool antibodies should fulfill.



## Technology platforms

We can offer access to different technologies to assist in your target validation, e.g. gene expression profiling or other state-of-the-art research technologies. Please detail your research interest for the technology platforms in the online submission.



## Further Collaboration

Successful projects supported by the grant may lead to further collaboration.

WHAT WE ARE FUNDING



YOUR BENEFITS

### APPLICATION PROCESS

APPLICATION PROCESS

SUBMISSION

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FAQ

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### 1. CREATE AN ACCOUNT



Pharmaceuticals





## 2. SUBMIT YOUR TARGET IDEA

**Deadline 31st of August 2017**

You'll need to submit an abstract describing your idea and target, its characteristics and a separate description of its therapeutic potential.

Please note that only non-confidential information should be provided.



## 3. EVALUATION OF PROPOSALS

**6 - 8 weeks after deadline**

The information provided will be evaluated by Bayer scientific experts. The decision on amount of funding is dependent on the specific target and research plan proposed.



## 4. PROJECT STARTS

**From November 2017**

Successful applicants will be notified by Bayer and will receive further instructions concerning the grant. A Bayer caretaker will be in contact with you during the grant period.



## READY, SET, SUBMIT

### WHAT WE ARE FUNDING

### YOUR BENEFITS

To submit, you will be redirected to ScholarOne™ online submission platform. All the information provided has to be non-confidential to allow a first evaluation of the

### APPLICATION PROCESS

proposal.

### SUBMISSION

**Please include the following information in your submission:**



FAQ

Proposal abstract (max. 2000 characters)



Contact Us

(/home/pharma/faq/index.php)

Describe your idea and target, its characteristics (indication and treatment paradigm, the target's druggability, and patent status).

(/scripts/components/forms/contact.php)



- + Description of therapeutic potential of the target
- + Upload CV and relevant publications

#### GO TO SUBMISSION

(<http://mc.manuscriptcentral.com/grants4targets>)



### WORDS FROM OUR PARTNERS

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### PUBLICATIONS

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- ↗ Gashaw I., et al., 2011. What makes a good drug target?  
(<http://www.ncbi.nlm.nih.gov/pubmed?term=21945861>)
- ↗ Kortylewicz Z., et al., 2012. Targeted molecular radiotherapy of cancer: Synthesis and Biological Evaluation (<http://www.ncbi.nlm.nih.gov/pubmed/22339166>)
- ↗ Lessl M., et al., 2011. Crowd sourcing in drug discovery  
(<http://www.nature.com/nrd/journal/v10/n4/full/nrd3412.html>)
- ↗ Stebbing J, et al., 2013. LMTK3 is implicated in endocrine resistance via multiple signaling pathways Oncogene (<http://www.ncbi.nlm.nih.gov/pubmed/22869149>)
- ↗ Suo G., et al., 2014. Telomerase expression abrogates rapamycin-induced irreversible growth arrest of uterine fibroid smooth muscle cells.  
(<http://www.ncbi.nlm.nih.gov/pubmed/24784716>)
- ↗ Shahraz A., et al., 2015. Anti-inflammatory activity of low molecular weight polysialic acid on human macrophages.  
(<http://www.ncbi.nlm.nih.gov/pubmed/26582367>)

WHAT WE ARE FUNDING

YOUR BENEFITS

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**MORE OPEN INNOVATION BY BAYER**

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