**Fit** for **Health 2.0** Support to SMEs & Researchers in FP7 and Horizon 2020 health-oriented projects

# Fast track to Innovation: a new instrument in Horizon 2020



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29.01.2015 | Webinar

Ines Haberl | Austrian Research Promotion Agency





# Content



- 1. Concept
- 2. How to prepare an application?
- 3. Evaluation procedure & ranking & timelines
- 4. Further information & support
- 5. Recommendations

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Concept



- Only fully bottom-up measure in Horizon 2020 (within the "Societal Challenges" and "Leadership in Enabling and Industrial Technologies")
- Addresses close to the market innovation activities
- Reduces time from idea to market
- Increases participation of industry
- Stimulates private sector investment in research and innovation

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### Concept



- Time to market: 36 months (from project start)
- Open to all types of participants
- Consortium partners with complementary backgrounds, knowledge and skills

-> Support of trans-disciplinary and cross sector cooperation

 Transnational value-chains and EU-wide / global markets are addressed

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### Budget



- For 2015 and 2016:
  - 200 Mio € (100 + 100)
  - about 50-70 actions (projects) will be funded / year
  - Grants for up to 3 Mio € possible (maximum!)
  - -> approximately 20 actions per Cut-off to be funded for all themes (bottom-up!)
  - -> highest competition and very high oversubscription to be expected!

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### Funding



• Funding:

- 70% for private-for-profit legal entities (like for innovation actions)
- 100% for non-for-profit legal entities

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## Which activities will be funded?



Time to market: 36 months or less from action start

- Advanced and specific research and development
- Advanced performance testing
- Piloting
- Demonstration activities
- Final validation of a system in the operational environment
- Business model validation
- Marketing activities (not purely commercial)
- Activities for strategic commercial and technical relevance

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## Which activities will be funded?



#### Level of development:

- Technology readiness level (TRL)
- TRL 6 -> TRL 9 in very short time frame
- TRL 6 is minimum requirement (demonstration in a relevant environment):

#### **Examples**:

- Development of a new drug: should have already been tested in Phase 1 -> Phase 2 clinical studies will be funded
- Medical devices: Class III device safety has been demonstrated -> final product design validation and final prototype production and testing will be funded
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### Who can participate?



- Open to all types of participants from EU-member states or associated countries:
  - SMEs
  - Industry
  - First-time industry applicants
  - Universities
  - Research centers
  - Technological organisations
  - Clusters & Associations
  - Incubators
  - End-users
  - Public sector
    - Investors

#### IMPORTANT: Third country partners: only as subcontractors! www.fitforhealth.eu

### Who can participate?



- Consortia of min. 3 and max. 5 participants from at least 3 different countries
- Industry involvement:
  - is mandatory!
  - either at least 60% of overall budget of proposal allocated to industry partners in the consortium

#### or

- minimum number of industry participants must be
  - 2 in a consortium of 3 or 4 partners
  - and 3 in a consortium of 5 partners

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FTI

### **Comparison FTI and SME-instrument**



#### Fully bottom-up

- Consortial approach
- Not SME related
- Obligatory to be on market after 3 years
- No coaching support
- One-stage scheme

### **SME-Instrument**

- Topics
- Single participation possible
- Restricted to SMEs
- No obligation to be on market after phase 3
- Coaching support
- 3 phases

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# Content



- 1. Concept
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### **Application**



Your proposal consists of 2 parts:

- On-line administrative forms Part A
- Upload descriptive part Part B (2 pdf files)



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### Application



- Specific template is available on the participant portal: <u>http://ec.europa.eu/research/participants/data/ref/h2020/other/call\_ptef/pt/h2</u> <u>020-call-pt-ia-fti-pilot\_en.pdf</u>
- As soon as you are registered with an action you will be able to download the word documents to insert your text

#### **ATTENTION:**

- The proposal template deviates substantially from the standard proposal template for other innovation actions!
- No separate business plan needs to be submitted

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### **Application: Part A**

#### Structured Part A – online

### Section 1

- Title, acronym, objective etc.
- Keywords
- 2000 character proposal abstract
- Declarations

# Section 2 (one form per partner)

- Participant Identification Code (PIC)
- Department
- Contact information
- Budget
- Ethics issue table
- Call specific questions



Proposal ID 123456 Acronym ACRO					
General Inf	ormation				
Tapic	ABCD Type of action Action				
Call identifier	SEP-ABCD				
Proposal title	The title should be no longer than 200 characters (with spaces) and should be understandable to the non- specialist in your field.				
Duration in months	Insert the estimated duration of the project in full months.				
Ford keyword	Please select the keyward(s) that best characterise the subject of your proposal in order of priority.				
Free keywords	You may enter a number of keywards that you consider necessary to characterise the scope of your proposal. There is a limit of 200 characters.				
The abstract should their relevance to th process and in com and precise and sho characters. If the pr	provide the reader with a clear understanding of the objectives of the proposal, how they will be achieved, and the Work Programme. This summary will be used as the short description of the proposal in the evolution munications to the programme management committees and other interested parties. It must therefore be short will not contain confidential information. Please use pixel topat lend, avoiding formulae and other special expand is written to a forugues other than English, please include an English version of the proposal abstract in ork. There is a limit of 2000 characters (with spaces).				
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# **Application: Part B**



- **1. EXCELLENCE**
- 2. IMPACT
- **3. IMPLEMENTATION** –

#### Attention:

Page limit: cover page, sections 1, 2 & 3, together **not longer than 30 pages**! All tables in these sections must be included within this limit!

- 4. MEMBERS OF THE CONSORTIUM
- **5. ETHICS AND SECURITY**

not covered by the page limit!

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**Cover page** 



	COVER PAGE		
Title of Proposal			
List of participant	ts (min. 3, max. 5)		
Participant No *	Participant organisation <u>name</u>	Country	First Time Industry Applicant**
1 ( <u>Coordinator</u> )			Y/N
2			Y/N
3			Y/N
4			Y/N
5			Y/N

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# Application First time industry applicant:

- Legal for profit organisation
- Obtained a PIC (Participant Identification Code) for the very first time

registrered for the first time in the beneficiary register during the preparation of the proposal (this will be checked by the EC!)

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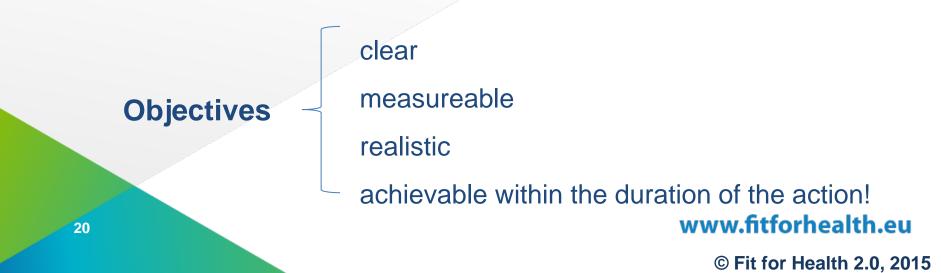


# **1. EXCELLENCE**

- 1.1 Objectives
- 1.2 Relation to the work programme
- 1.3 Concept and approach
- 1.4 Ambition

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- 1. EXCELLENCE
  - 1.1 Objectives
- Specific objectives
- Consistent with expected exploitation and impact
- Which industrial/economic/social challenge do you address?
- How will your solution meet this challenge?







#### 1. EXCELLENCE

- 1.2 Relation to the work programme
- What are the reasons why you fit into the Fast Track to Innovation Work Programme?
  - Business driven
  - Realistic potential for quick deployment and market take-up of your innovation
  - .....

-> See other main characteristics in the introduction part of this webinar!

#### Work programme:

http://ec.europa.eu/research/participants/portal/doc/call/h2020/common/1615113part\_18\_fti\_v2.0\_en.pdf

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# **1. EXCELLENCE**

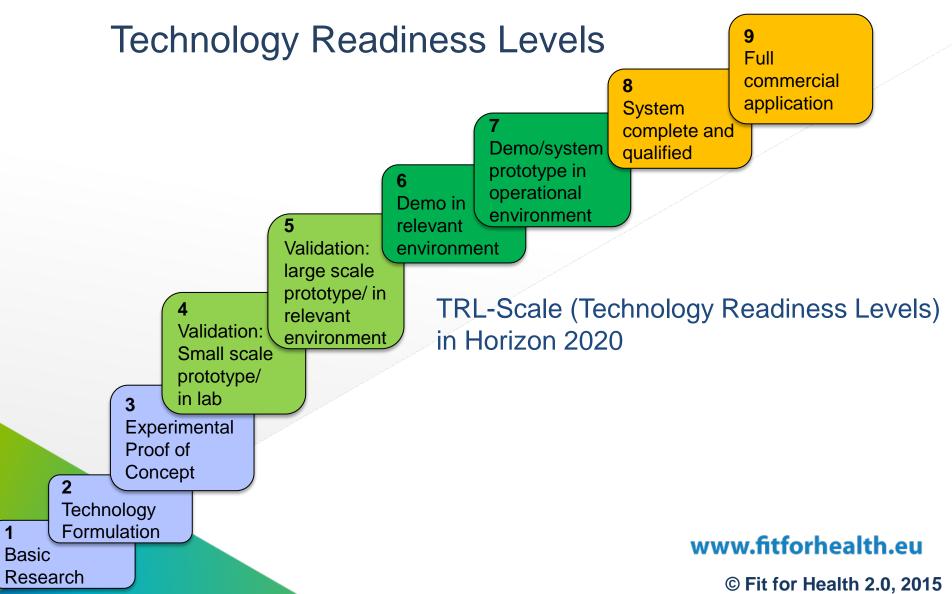
- 1.3 Concept and approach
- Overall concept, main ideas, models, assumptions, transdisciplinary considerations: which activities will you implement?
- Starting point and level of maturity: refer to TRLs (Technology Readiness Levels)

**ATTENTION: TRL of 6 or higher is expected!** 

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# **Technology Readiness Levels**

#### Examples:

- Scale-up and initiate validation of GMP manufacturing process
- Phase 2 clinical trials
- Validation of assays for manufacturing quality control and immunogenicity

Annex G of the Work Programme:

http://ec.europa.eu/research/participants/data/ref/h2020/wp/2014\_ 2015/annexes/h2020-wp1415-annex-g-trl\_en.pdf

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# **1. EXCELLENCE**

- 1.3 Concept and approach
- How will the concept and development lead to effective market take-up?
- Elaborate on the European dimension, what is the added value for Europe?
- Where relevant, describe how sex and/or gender analysis is taken into account in the action's content <u>http://ec.europa.eu/research/science-</u> <u>society/gendered-innovations/index\_en.cfm</u>







# **1. EXCELLENCE**

- 1.4 Ambition
- What is the innovation potential? What is the advantage beyond the state-of-the-art?
- What are expected key market applications extracted from the results what provides highest value added to potential customers?
- What is your **USP**?

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### 2. IMPACT

- 2.1 Expected Impacts
- 2.2 Measures to maximise impact
  - a) Dissemination and exploitation of results
  - b) Intellectual property, knowledge protection and regulatory issues
  - c) Communication

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### 2. IMPACT

- 2.1 Expected Impacts
- What are the needs of European / global markets?
- How will your action strength competitiveness and growth of all industrial partners in your consortium?
- Which user needs have been identified (outcompete current solutions?)
- Cost-benefit analysis (comparing to solutions offered by competitors)
- What is your target market? Estimation of available market size, growth rate, trends, main selling points, initial introduction, potential early adopters/users
- How will your innovation influence scientific, technological / other progresses?

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### 2. IMPACT

2.1 Expected Impacts

- How will the (industry) partners in the consortium grow? (turnover, market share, employment, long term sales, return on investment,...).
- Capital investment policy for the next 3 years (percentage/relevance of own funds, FTI funding, other external funds like loans, venture capital etc.)
- What are **barriers**, obstacles, framework conditions,...?

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### 2. IMPACT

• 2.2 Measures to maximise impact

### • a) Dissemination and exploitation of results

- Draft plan for dissemination and exploitation of action's results
- Plan needs to include full range of potential users (research, commercial, investment, social, environmental, policy making, setting standards, skills, educational trainings)
- Plan should be tailored in
  - Specific technical issues
  - Market issues
  - Organisational issues

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### 2. IMPACT

2.2 Measures to maximise impact

### • a) Dissemination and exploitation of results

- Which stakeholders should be involved?
- How will the consortium partners benefit in terms of competitiveness, innovation capacity (corporate strategies of industrial partners)
- o Time to market / deployment
- How will you **position yourself in the market** over time?
- Timeline for detailed planned marketing and sales efforts (distribution!)
- Possible further development strategies

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### 2. IMPACT

- 2.2 Measures to maximise impact
  - b) Intellectual property, knowledge protection and regulatory issues
- Who owns IPR? Which ways of protection did you choose?
- How is commercial exploitation ensured?
- What is the strategy for knowledge management, current IP status?
- Regulatory and/or standard requirements are to be fulfilled for the exploitation of the innovation
  - Where relevant: info how you will manage research data generated/collected and measures to provide open access (green or gold model)

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## 2. IMPACT

2.2 Measures to maximise impact

### c) Communication

- How will you promote the action?
- Consider the needs of various audiences (groups beyond the action's own community)
- Measures for public/social engagement

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# **3. IMPLEMENTATION**

- 3.1 Work plan Work Packages, deliverables and milestones
- 3.2 Management structure and procedures
- 3.3 Consortium as a whole
- 3.4 Resources to be committed

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# **3. IMPLEMENTATION**

- 3.1 Work plan Work Packages, deliverables and milestones
  - Briefly present the overall structure of the work plan
  - Timing of work packages (e.g. with Gannt chart)
  - Detailed work description with tables
  - Graphical presentation of inter-relation (e.g. Pert chart)

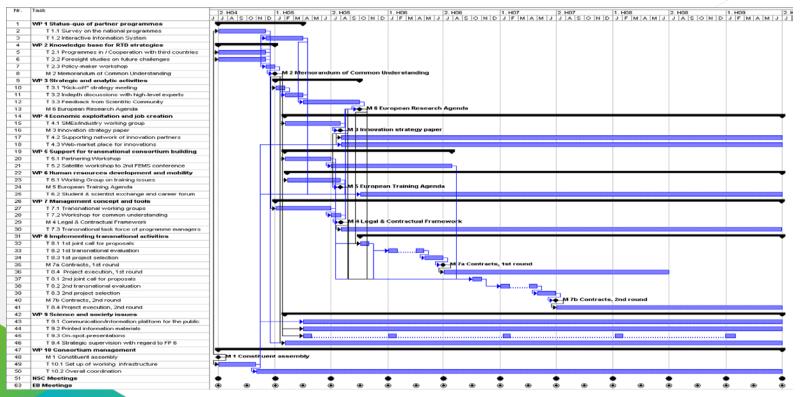
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### **3. IMPLEMENTATION**

#### **Example Gannt Chart**

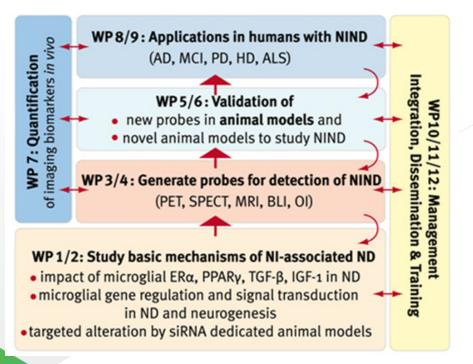


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# **3. IMPLEMENTATION**

### **Example Pert Chart**



INMiND project (<u>www.uni-muenster.de/InMind/</u>)



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# **Application**

### **3. IMPLEMENTATION**

Work package title Participant number Short name of participant Person/months per participant:						
Short name of participant						
Person/ <u>months</u> per participant:						
out at						
Objectives						
Description of work (where appropriate, I	oroken down i	nto tasks)	lead partr	ner and rol	e of partic	rinants
beschption of work (where appropriate, i	or oken downin	1100 (03/03/),	ices para		e or paren	
Deliverables (brief description and month	of delivery)					
	_	1	_	_	_	_

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#### Work Package (WP) description (table 3.1a)

- Give full details for the resources for each WP
- Number of WPs should reflect the complexity
- Always include a WP for 'management'
  - 'Dissemination' and 'exploitation' & 'Communication' as distinct tasks or distinct WPs
- Quantified information!

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## **3. IMPLEMENTATION**

#### Table 3.1b: List of Work packages

Work package No	Work Package Title	Lead Participant No	Lead Participant Short Name	Person- Months	Start <u>Month</u>	End month
				Fotal <u>months.</u>		

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#### **3. IMPLEMENTATION**

Deliverable name	Work package number	Short name of lead participant	Туре	Dissemination level	Delivery date
		Deliverable name package	Deliverable package of lead	Deliverable package of lead Type	Deliverable package of lead Type Dissemination



#### Different types (codes)

- R: document, report
- DEM: demonstrator, pilot, prototype
- DEC: website, patent filing
  - **OTHER:** software

#### **Dissemination level (Codes)**

- PU: public, fully open (web)
- CO: confidential, restricted
- CI: classified

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### **3. IMPLEMENTATION**

- 3.2 Management structure and procedures
- Organisational structure & the decision-making (appropriate to complexity and scale!) including *table 3.2a: list of milestones*
- How will you effectively address *innovation management*?
- What are *critical risks* and what are your *mitigating actions* (*table 3.2b*)?

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Means of

verification

Estimated date

### **3. IMPLEMENTATION**

List of milestones (table 3.2a)

- Control points
- Facilitate to chart the progress
- Could correspond to the completion of key deliverables

Milestone

number

Milestone

name

Related work package(s)

Critical decision points

Means of verification:

how will you confirm that a milestone has been attained?

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## **3. IMPLEMENTATION**

#### Critical risks for implementation (table 3.2b)

Description of risk	Work package(s) involved	Proposed risk-mitigation measures

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### **3. IMPLEMENTATION**

- 3.3 Consortium as a whole
- Describe the consortium:
  - o matching project's objectives
  - o complementing each other (cover the value chain)
  - o contribution to the project
  - o effectively work together

**ATTENTION:** detailed description of consortium members in section 4!

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# **3. IMPLEMENTATION**

3.4 Resources to be committed

Summary of staff effort table (table 3.4a)

- Showing the number of person months required
- person months for whole duration for each WP & participant

	WPn	WPn+1	WPn+2	Total Person/ Months per Participant
				montus per Participant
Participant				
Number/Short Name				
ParticipantNumber				
Short Name				
Participant Number/				
Short Name				
Total Person/Months				

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# **3. IMPLEMENTATION**

3.4 Resources to be committed

'Other direct cost' items (travel, equipment, other goods and

services, large research infrastructure) (table 3.4b)

 Showing other direct costs for participants where those costs exceeds 15% of the personnel costs

Participant Number/Short Name	Cost (€)	Justification
Travel		
Equipment		
Other goods and services		
Total		
Participant Number/Short Name	Cost (€)	Justification
Large research infrastructure		

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### 4. MEMBERS OF THE CONSORTIUM

### **4.1 Participants (applicants)**

- Description of the legal entity and main tasks
- Curriculum vitae or description of the profile of the persons, who will be primarily responsible for carrying out the proposed activities
- List of up to 5 relevant publications, and/or products, services
- List of up to 5 relevant previous projects or activities
  - Description of any **significant infrastructure** and/or any major items of technical equipment

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# 4. MEMBERS OF THE CONSORTIUM

### 4.2 Third parties involved in the action

Please complete, for each participant, the following table (or simply state "No third parties involved", if applicable)

Does the participant plan to subcontract certain tasks (please note that core tasks of the action should not be sub-contracted)	Y/N
If yes, please describe and justify the tasks to be subcontracted	
Does the participant envisage that part of its work is performed by linked third parties <sup>1</sup>	Y/N
If yes, please describe the third party, the link of the participant to the the describe and justify the foreseen tasks to be performed by the third part	
Does the participant envisage the use of contributions in kind provided by third parties (Articles 11 and 12 of the General Model Grant Agreement)	
If yes, please describe the third party and their contributions	-

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### \_ \_\_\_\_

### **5. ETHICS AND SECURITY**

**5.1 ETHICS** 

**Application** 

Ethics issues have to be entered in the **ethical issue table** in the administrative proposal forms:

- submit an ethics self-assessment
- provide the documents that you need under national law (if you already have them) e.g.
  - an ethics committee opinion
  - the document notifying activities raising ethical issues or authorising such activities

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# **5. ETHICS AND SECURITY**

# **5.2 SECURITY**

Please indicate if your project will involve:

- activities or results raising security issues: (YES/NO)
- 'EU-classified information' as background or results: (YES/NO)

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# **Evaluation Procedure**

- Thresholds:
  - Overall score of 12 out of 15
  - Score for 'Impact' at least 4 out of 5
- Evaluation occurs fully remote, no consensus meetings
- Final scores: arithmetic average of individual scores
- Evaluation summary reports with short feedback sent to you

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### Who are your evaluators?



- Each proposal will be evaluated by 4 different experts:
  - Experts with market knowledge, financial skills, business experience
  - Only 1 of them can be from the same country as the coordinator!

ATTENTION: Evaluators are not necessarily from your research area!

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# Ranking



- All proposals reaching the thresholds will be ranked
- Score for 'impact' will be multiplied with 1.5 before the ranking starts!
- Additional criteria for evaluation:
  - Value of 'impact' criterion
  - Size of budget allocated to SMEs
  - Number of first-time industry applicants
  - Number of industry applicants
  - Gender balance of consortium staff performing the action

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## **Timelines**



- Single permanently open Call since 6th January 2015 with Cut Off Dates
- Proposals can be submitted any time
- Evaluation starts for all proposals after the respective Cut off date
- 3 Cut off dates in 2015 and 2016
- First results from 1st Cut off: end July 2015
- Time to grant: 6 months
  - First grants signed: end October 2015

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Cut-off-dates 2016	TBD	TBD	TBD	
Cut-off dates 2015	29/4/2015	1/9/2015	1/12/2015	

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# **Further information and support**



Participant portal

http://ec.europa.eu/research/participants/portal/desktop/en/opportunities/h2020/topics/9096ftipilot-1-2015.html

http://ec.europa.eu/programmes/horizon2020/en/h2020-section/fast-track-innovation-pilot-2015-2016

Work Programme

http://ec.europa.eu/research/participants/data/ref/h2020/wp/2014\_2015/main/h2020wp1415-fast-track\_en.pdf

Horizon 2020 Online Manual:

http://ec.europa.eu/research/participants/portal/desktop/en/funding/guide.html

Information about EU-member states and associated countries

http://ec.europa.eu/research/participants/data/ref/h2020/grants\_manual/hi/3cpart/h2020-hilist-ac\_en.pdf?\_=58655886

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# **Further information and support**



### National Contact Points

http://ec.europa.eu/research/participants/portal/desktop/en/sup

port/national\_contact\_points.html

### IPR-Helpdesk

https://www.iprhelpdesk.eu/

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# Content



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Take home messages & recommendations



### Advantages and disadvantages of FTI:

- High competition -> high visibility for successful consortia -> increased market success
- Establishment of new value chains and networks
- Fast funding
- BUT

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- Take high competition into account!
- The FTI-programme is a pilot, new to all of us...

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### Thank you!

FFG

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