



Version 21/04/2016

COST Action BM1406 (2015-03-31– 2019-03-30)

IONCHAN-IMMUNRESPON: ION CHANNELS and IMMUNE RESPONSE Toward a global understanding of immune cell physiology And for new therapeutic approaches

PROGRESS REPORT AT MONTH 24

This report is submitted by the MC Chair on behalf of the Management Committee.

Confidentiality: This report, other than section II.D, is non-confidential. Section II.D is confidential to the Management Committee and the COST Association (including the Committee of Senior Officials, Scientific Committee and Administration).

Executive summary of the Progress Report:

The activity of the COST Action BM1406 entitled “Ion Channels and Immune Response toward a global understanding of immune cell physiology and for new therapeutic approaches (IONCHAN-IMMUNRESPON)” during the 24 Month period was characterised by 5 meetings and workshops including the kick off meeting, and 9 STSM.

The aim of all the meetings was to merge in a single location the participants in order to spur new collaborative ideas and projects. In this term, the activity was very successful. Of particular interest, the outcome of the meeting was the beginning of collaborative exchange on topics such as strategies to modulate ion channels in immune cells, thanks to antibodies, drugs and siRNA. The Third Meeting in Lisbon, in which we focused on the WG3, clearly demonstrated that the siRNA approach is not yet available for therapies through the ion channels. Chemical drugs and therapeutic antibodies seem to be a better trail to investigate ion channel modulation. Each meeting gave the opportunity to the participants to exchange scientific expertise and to have fruitful discussions. The STSMs of young researchers inside the Action increased the knowledge of skills of each laboratory that allows the collaborations and material exchanges. In this, our objective, which is to build working relationships within European research groups, is reached in full.

The outcomes of the Action could be appreciated thank to the numbers of submitted projects on the national and European levels, the numbers of publications in which the Action has been acknowledged, the collaborative projects supported or not by the Action, the number of ITC partners involved into these projects. The important scientific advance has been reported in the paper of Prof. di Virgilio and Pablo Pelegrin concerning the role of the nlrp3/P2X7 in cancer cells. It is a major concern for this type of cancers to understand which inflammation is induced and why the immune system does not attack against the cancer cells. The Action has allowed also the study of very particular channels such as aquaporin, which has lead to the publication of three articles.

The Action has increased the individual scientific foresight and a large number of us have been invited to the other laboratories for conferences and seminars in each partners countries. We paid a large attention to the young researchers and PhD. Fellow for those STSM are priority **dedicated**.

We have a nice success story with the creation of a start ‘up by our colleagues from Switzerland in Geneva that participate to the industrial task of Europe. Nevertheless, we observed that the relation with industries in the Action is less efficient to start collaborative projects and so we decided for the next GP to increase the number of STSM for the industrial partners. To sum up, this first period of activity allows us to establish scientific relation in a common scientific area, a to build strategies to study the role of ion channels in immune cells, and evaluate the appropriate trail to module the immune cell activity.



COST is supported by
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I. Progress Report

I.A. COST Action Profile

Objective/ Aim

The main objective of the Action is to use modulation of ion channels for therapeutic approaches in immune diseases by identification of the pertinent targets, the development of animal models and a better understanding of their role in immune cell functionality.

Details

MoU:	084/4	Start of Action:	2015-03-31
CSO approval date:	2014-11-13	End of Action:	2019-03-30

COST Member Countries and Cooperating State having accepted the MoU

Country	Date	Status
Austria	20/01/2015	Confirmed
Belgium	06/03/2017	Confirmed
Bosnia and Herzegovina	21/10/2015	Confirmed
Croatia	03/02/2015	Confirmed
Czech Republic	26/03/2015	Confirmed
Denmark	05/12/2014	Confirmed
Estonia	11/02/2015	Confirmed
France	27/11/2014	Confirmed
fYR Macedonia	30/01/2015	Confirmed
Germany	15/12/2014	Confirmed
Greece	16/12/2014	Confirmed
Hungary	17/12/2014	Confirmed
Ireland	25/07/2016	Confirmed
Israel	30/11/2014	Confirmed
Italy	29/01/2015	Confirmed
Latvia	27/03/2015	Confirmed
Luxembourg	01/12/2014	Confirmed
Norway	11/02/2015	Confirmed
Poland	05/12/2014	Confirmed
Portugal	16/02/2015	Confirmed
Serbia	04/03/2016	Confirmed
Slovenia	18/03/2015	Confirmed
Spain	21/11/2014	Confirmed
Sweden	11/02/2015	Confirmed
Switzerland	16/01/2015	Confirmed
Turkey	24/08/2015	Confirmed
United Kingdom	18/11/2014	Confirmed

Total: 27

Intentions to Accept the MoU

0

Other participants:

Institution Name	Country
New York University School of Medicine Department of Pathology and Cancer Institute	USA
Washington State University, Neuroscience Department of Integrative Physiology and Neuroscience	USA

Contacts

Chair/ Vice Chair

Position	Name	Contact details	Country	Date of PhD:	Gender
Chair:	VELGE-ROUSSEL Florence	University of Tours, UFR Medicine 10, Bvd Tonnellé, 37032 TOURS France, Tel.+33247366058 Fax +33247366146 velge@univ-tours.fr	France	1989	F
Vice Chair:	PELEGRIN Pablo	Murcia's Biomedical Research Institute (IMIB-Arrixaca) Edificio LAIB Carretera Buenavista s/n 30120 Murcia Spain +34868885038 pablo.pelegrin@ffis.es	Spain	2003	M

Working Group Leaders

WG#	WG Title	WG Leader	Country	Date of PhD:	Gender	Number of participants
1	Identification and Characterization of ion channels in immune cells	Dr Ruth Murrell-Lagnado	UK	1988	F	60
2	Role of ion channels in immune pathologies	Prof Francesco Di Virgilio	IT	?	M	40
3	Ion channels as new targets in therapy and diagnosis	Prof Friedrich Koch-Nolte	DE	?	M	30

Other positions if applicable (STSM Coordinator, WG Vice Leader, Task Force Leader...)

Position	Name	Country	Date of PhD:	Gender
Representative of Early-Stage researchers (ESRs)	Ellegaard M.	DK	2013	F
Representative of SMEs	Toomas Neuman	EE	?	M
Coordinator of STSMs	Barbara Niemeyer	DE	?	F
Dissemination Coordinator	Antoine El Chemaly	CH	?	M

Action website: <http://costbm1406.univ-tours.fr/home/>



I.B. Progress with MoU objectives and deliverables and additional outputs

MoU objectives

MoU objective	Achieved Yes/ Partially/ No	Evidence of (partial) achievement including hyperlink to enable assessment of the achievement ¹ . Justification if full achievement is not foreseen
OBJ1. Available data managing and validation strategies consensus for identified targets.	Partially	Each scientific article represents a step in the knowledge about each target identified Each meeting allowed achievements and collaborations i.e. scientific reports of all meetings http://costbm1406.univ-tours.fr/home/
Obj2. Identification of target diseases.	Partially	WG2 meeting in Belgrade focused on the thematic and some candidates have been evidenced
Obj3. Translating OBJ1 and 2 into preclinical animal models.	Partially	Exchanges between partners during the Zagreb meeting allow the exchange of mouse models
OBJ4. Therapeutics	Partially	We explored the three strategies explained into the MoU in the Lisbon meeting with expert's advices. See Lisbon report (http://costbm1406.univ-tours.fr/home/)

MoU deliverables

MoU deliverable	Level of progress ¹	Evidence of (partial) delivery achievement including hyperlink to enable assessment of the delivery ¹ . Justification if full achievement is not foreseen
Building of BM1406 website to communicate with all members	Achieved	http://costbm1406.univ-tours.fr/home/ https://www.facebook.com/BM1406?fref=ts
Position paper of F di Virgilio, Involvement of the P2X7-NLRP3 axis in leukemic cell proliferation and death. Sci. Rep. 6, 26280	Achieved	http://www.nature.com.gate2.inist.fr/articles/srep26280
Publication of periodic newsletter send to all members of BM1406 Action	Partially achieved	Two newsletters per year http://costbm1406.univ-tours.fr/newsletters/
SOPs in microscopy for calcium image	Achieved	http://kurser.ku.dk/course/nscphd1123/
Position paper Special issue in Current Opinion in Immunology will be elaborated by the BM1406 members	Partially achieved	http://www.current-opinion.com/journals/current-opinion-in-immunology/

Co-authored publications and FP7/ H2020 proposals

The co-authored publications and FP7/ H2020 proposals/ projects resulting from the Action are listed on the page following the "Additional outputs and achievements" section

Additional outputs and achievements

Please describe any other outputs and achievements that have resulted or are in progress, focusing in particular on those that contribute to the COST mission of "COST enables break-through scientific developments leading to new concepts and products and thereby contributes to strengthen Europe's research and innovation capacities."
Collaboration between Croatia and Portugal members open the new thematic with the identification of the role of aquaporins in immune cells (ITC, EWP)

¹ The links to the outputs and deliverables will be used by the Action Rapporteur in assessing the progress.



Determination of STIM1 role in depression (Majeswski et al.2016)
Demonstration of the pole of MLRP3 and P2X7 axis in the cancer cells.



Co-authored publications

Enter in the table below only publications on the topic of the Action, co-authored by at least two Action participants from two different countries participating in the Action and for which the Action networking added value. A maximum of ten publications may be entered. If the Action has more than ten such publications the Core Group should select the ten most significant ones to include in the table below.

NO.	Bibliographic data (including: Title, Authors, Title of the periodical or the series, Issue number or volume, Publisher, Year of publication, Relevant pages)	Main author	Number of authors	Action participants listed among the authors (Name, country and role ²)	WGs involved in publication	Date of submission (must be after Action start date)	Expected date of publication (if not already published)	Persistent link to publicly available version of the paper (if available) or the abstract	Is/Will open access ³ provided to this publication?	Is/ will COST be cited/ acknowledged in the publication?	Are/ will COST funds (be) implicated in this publication	Relevance to H2020 Societal Challenges ⁴ ?	Is it peer-reviewed?	Was the added value of the Action Networking necessary for the publication	Impact Factor (if applicable)
1	Immature human dendritic cells enhance their migration through KCa3.1 channel activation. David Crottès, Romain Félix, Daniel Meley, Stéphanie Chadet, Florence Herr, Cindy Audiger, Olivier Soriani, Christophe Vandier, Sébastien Rogerb, Denis Angoulvant, Florence Velge-Roussel. <i>Cell Calcium</i> 59 (2016) 198–207.	David Crottès	11	C. Vandier, FR WG1, S. Roger, FR WG1, F. Velge-Roussel, FR Chair	WG1	17/11/2015	Accepted 15/02/2016	http://www.sciencedirect.com/science/article/pii/S0143416016300112	no	no	no	Yes	Yes	USA partner pre-reviewed the article	3,43
2	M1 and M2 Functional Imprinting of Primary Microglia: Role of P2X7 Activation and miR-125b. Mediators of Inflammation Volume 2016, Article ID 2989548, 9 pages http://dx .	Chiara Parisi	4	Chiara Parisi, Italy, STSM recipient, Pablo Pelegrin, Spain V Chair, and Cinzia Volonté Italy, MC substitute	WG1	23 /09/2016	Accepted 24 November 2016	doi.org/10.1155/2016/2989548	no	Yes	Yes	Yes	Yes	STSM of Parisi	3.418
3	Involvement of the P2X7-NLRP3 axis in leukemic cell proliferation and death. <i>Sci. Rep.</i> 6, 26280	Erica Salaro	16	Alba Clara Sarti, Italy STSM recipient, Pablo Pelegrin, Spain, V Chair, Francesco Di Virgilio, Italy, WGL2	WG2	30/07/2015	Accepted: 25 April 2016	doi: 10.1038/srep26280 (2016)..	no	Yes	No	Yes	Yes	STSM Sarti	5.578
4	UCP2 up-regulation within the course of autoimmune encephalomyelitis correlates with T-lymphocyte activation. <i>Biochimica et Biophysica Acta</i> 1863 (2017) 1002–1012.	Alina Smorodchenko	8	Anne Rupprecht Italy, WG2, Elena E. Pohl, Austria WG2	WG2	09/10/2016	Accepted 23 January 2017	http://www.sciencedirect.com/science/article/pii/S0925443917300327	no	Yes	No	Yes	Yes	Meeting exchanges	3.66
5	Rat Aquaporin-5 is pH-Gated Induced by Phosphorylation and is Implicated in Oxidative Stress. <i>Int. J. Mol. Sci.</i> 17: 2090, 2016.	Claudia Rodrigues	8	C. Rodrigues, Portugal STSM recipient, Ana Gasparovic, Croatia WG1, G. Soveral, Portugal, WG1	WG1	28/09/2016	06/12/2016	doi:10.3390/ijms17122090	no	Yes	Yes	Yes	yes	STSM Rodrigues	3.25
6	Yeast aquaporin regulation by 4-hydroxynonenal is implicated in oxidative stress response. <i>IUBMB Life.</i> 2017 Mar 24.	Claudia Rodrigues	5	C. Rodrigues, Portugal STSM recipient, Ana Gasparovic, Croatia WG1, G. Soveral, Portugal, WG1	WG1	09/01/2016	03/03/2017	doi: 10.1002/iub.1624.	no	Yes	Yes	Yes	Yes	STSM Rodrigues	2.65
7	NLR Proteins, <i>Methods in Molecular Biology</i> 1416, Springer ed.	Francesco Di Virgilio Pablo Pelegrin	55	Anna Rubartelli, WG2, Francesco di Virgilio, WG2 Pablo Pelegrin WG1, Vice Chair	WG1 /WG 2	16/10/2015	15/05/2016	http://www.springer.com/series/7651	no	no	no	Yes	Yes	Meeting exchanges	0.79

² MC Member/ MC Substitute/ MC Observer/ WG Member/ Training School Trainee/ STSM Recipient/ Other Action Participant

³ Open Access is defined as free of charge access for anyone via Internet. Please answer "yes" if the open access to the publication is already established and also if the embargo period for open access is not yet over but you intend to establish open access afterwards.

⁴ H2020 Societal Challenges are "Health, demographic change and wellbeing"; "Food security, sustainable agriculture and forestry, marine and maritime and inland water research, and the Bioeconomy"; "Secure, clean and efficient energy"; "Smart, green and integrated transport"; "Climate action, environment, resource efficiency and raw materials"; "Europe in a changing world - inclusive, innovative and reflective societies"; "Secure societies - protecting freedom and security of Europe and its citizens"

8	<i>P2 receptors in cancer progression and metastatic spreading</i> <i>Current Opinion in Pharmacology</i> 2016, 29:17–25	Francesco di Virgilio	4	Francesco di Virgilio, WG2	WG2	31 March 2016	20 June 2016	http://dx.doi.org/10.1016/j.coph.2016.05.001	no	Yes	no	Yes	Yes	Meeting exchanges	4.742
9	<i>Extracellular purines, purinergic receptors and tumor growth</i> , <i>Oncogene</i> (2017) 36, 293–303	Francesco di Virgilio	2	Francesco di Virgilio, WG2	WG2	31 March 2016	2 May 2016	doi:10.1038/onc.2016.206	no	Yes	no	Yes	Yes	Meeting exchanges	7.932
10	<i>Overexpression of STIM1 in neurons in mouse brain improves contextual learning and impairs long-term depression</i> . <i>Biochim Biophys Acta</i> . 2016 Nov 29. pii: S0167-4889(16)30320-2.	Luckas Majewski	8	Luckas Majewski, WG1 Jack Kuznicki WG1	WG1	28 Sept 2016	26 Nov 2016	doi: 10.1016/j.bbamcr.2016.11.025 .	Yes	Yes	Yes (STSM)	Yes	Yes	STSM Iga Wasilewska	5.343
11	<i>Purinergic signaling in the immune system, Autonomic Neuroscience: Basic and Clinical</i> 191 (2015) 117–123	Francesco Di Virgilio	2	Francesco di Virgilio, WG2	WG2	02/2015	23 Nov 2015	doi: 10.1016/j.autneu.2015.04.011	Yes	Yes	no	Yes	Yes	Meeting exchanges	1.620

FP7/ H2020 Proposals and projects

This table contains FP7/ H2020 proposals/ projects spinning off from Action activities and including in the proposing consortium at least three Action participants from at least three different countries participating in the Action.

NO.	Title	Name and country of main proposer	Number of proposers	Action participants listed among the proposers (Name, country, role ³ in the Action)	Funding agency submitted to	Date submitted	Date results expected	Result	Call identifier	Relevance to H2020 Societal Challenges ⁴ ?	Was the added value of the Action Networking necessary for the proposal / project?
Projects											
1	• PHARMPROT teaming, GINOP-2.3.2-15-2016-00044 project.	Hungary	1	Prof. G Panyi Hungary WG1,	co-financed by the European Union and the European Regional Development Fund.		2016	accepted	ERDF	Yes Health, demographic change and wellbeing	
2	3-year project grant application	Ruth Murrell-Lagnado (UK)	2	Ruth Murrell-Lagnado (UK) Sebastien Roger (France)	Breast Cancer Now	June 2016		refused			
Proposals											
	Ion Transport Remodelling in Pancreatic Cancer IonPaC	G. Panyi, Hungary	2	G. Panyi, Hungary WG1, I. Novak, Denmark, WG1	MARIE SKŁODOWSKA-CURIE ACTIONS,	02/2017	08/2017		International Training Network	Yes, Health, demographic change and wellbeing	Open to European grants
	3-year project grant application	Ruth Murrell-Lagnado (UK)	2	Ruth Murrell-Lagnado (UK) Sebastien Roger (France)	World Wide Cancer Research (UK)	April 17	September 17			Yes, Health, demographic change and wellbeing	Building of Collaboration inside of the Action



I.C. Networking

Added value of the Networking

Joined publications

Numerous joined publications between Cost Countries

New Collaboration started

- 1, Collaboration with Nace Zidar Univerza v Ljubljani F
- 2, Collaboration with Iva Bozic, Irena Lavrnja, Univerz
- 3, Collaboration of Katja Ester Croatia with Jost Ludwig
- 4, Collaboration of Luckas Majesky, Poland with Barab
- 5, Collaboration of Ruth Murrell-Lagnado, UK with Coli
- 6, Collaboration of Ruth Murrell-Lagnado, UK with Seb
- 7, Collaboration of Ruth Murrell-Lagnado, UK with Rair
- 8, Collaboration of Pablo Pelegrin (Spain) with Frances
- Hafner (Slo), Cinzia Volonté (Italy), Sebastien Roger (F
- 9, Collaboration of Graca Soveral (Portugal) with Ana C
- 10, Collaboration of N. Demaurex (Swiss) with Christo

on STIM/ORAI

New projects submitted at national level

- Hungarian National Science Fund, K119417 F
- tumours, obtained
- 3-year project grant applications with R. Murre
- Cancer Now (June 2016) refused
- one 3-year project recently to World Wide Can
- One project from N. Demaurex (Swiss) obtaine
- Scientifique en 2016 (SNF 31003A_16949)
- 2016, "Regulacion del inflamasona nlrp3: impl
- Programa Estatal de Fomento de la Investigac
- Ministry of Economy and Competitiveness. Na
- 3-year project grant application Barbara Nieme
- from the DFG (FOR2289, 2016-2018)
- 4 year project grant application Barbara Nieme
- immune cell function by the DFG (SFB1027, 2

New thematic

Role of Aquaporins in immune cells (Croatia, Poland)
Neuronal inflammatory diseases (Belgrade, Croatia)

Extent of the networking

- Interface with BM1305 (A FACTT)
 - We contacted Dr Eva M MARTINEZ-C
 - Accelerate Cell-based Tolerance-induc
 - participants
- Interaction with BM 1307 ((PROTEOSTASIS)
 - We contacted BMBS 1307 thanks to the C
 - people interested by the immune cell them
- Meetings in ITC countries
 - At this date, we organized 4 meetings
 - in March 2016, Croatia in September 2
- Invitation of a local speaker in each meeting
 - We have the use to invite one local sp
 - which welcome us (see scientific progr



- STSM for the ESR
 - 7 applicants for the STSM came from ITC, 2 from Cost Member Country.
 - 4 STSM are from ITC to CMC,
 - 2 from CMC to ITC
 - 2 from ITC to ITC
 - 2 in the CMC

- Gender balance in the MC, number of ITC vs CMC
 - MC gender balance 31 women / 56 members = 50%
 - The great majority of the STSM have done by women (7 out 9 STSM from GP1 and GP2)
 - 6 out of 9 STSM have been done from people from ITC to a CMC.

I.D. Impacts

The impacts that have resulted, or might result from the Action are described in the following table.

Description of the impact	Type of impact ⁵	Timing of impact ⁶
Creating the ERA network <ul style="list-style-type: none"> • 66 MC members from 26 countries (12 ICT/26 countries) • Trans-disciplinarily network with Biophysics, chemists, immunologists, clinicians 	Scientific	Foreseen 5-10 years
New resources and tools According the scientific collaborations, new tools have been shared	Scientific	Foreseen within 2 years
Fruitful collaborations New collaborations have been build thank to the Action listed in I.C. networking Joined scientific publications	Scientific	Foreseen within 2 years
Enabling young scientist to work in excellent European partner laboratories	Scientific, Soft skill and professional advancement	immediate

I.E Dissemination and exploitation of Action results

Describe the Action's dissemination and exploitation approach as well as all activities undertaken to ensure dissemination and exploitation of Action results and the effectiveness of these activities.

Add description here

Item/ activity	Target audience	Result	Hyperlink
Newsletters	Scientists	¼ on website	http://costbm1406.univ-tours.fr/newsletters/
Dissemination Meetings	Scientists of the same area	1 in Lecce, Italy	Slides?

⁵ Scientific/ technological, Economic, Societal

⁶ Achieved/ Foreseen within 2 years/ Foreseen 2-5 years/ Foreseen 5-10 years/ Foreseen 10+ years

I.F. Action success(es)

COST regularly communicates the successes of Actions. At this point in time what aspect(s) (outcomes and/ or impacts, rather than activities) of this Action is/ are the most suitable for communication?

<p>The research on Hv1 proton channels financed by COST resulted in the creation of a company (Hplus therapeuDCs) of which Karl-Heinz Krause (Swi) is founder and Nicolas Demareux (Swi) the advisor scientific. http://ge.ch/hrcintapp/externalCompanyReport.action?companyOfrcld13=CH-660-0246017-1&ofrcLanguage=1</p>	<p>Dimension of the success</p> <ul style="list-style-type: none"> ■ Breakthrough: technological ■ Policy implementation (specify which policy) ■ Capacity building

II. Management Report

II.A. Overview of expenditure

Insert below in the yellow cells the summary of figures from the Yearly Financial Reports (YFRs) of completed Grant Periods and an IFR of any incomplete Grant Period – the Totals (non-yellow cells) will automatically sum.

	Grant Period 1	Grant Period 2	Grant Period 3	TOTAL
GP start and end dates	(01/06/2015-31/05/2016)	(01/05/2016-30/04/2017)	(01/05/2017-30/04/2018)	
Grant Holder institution	GH institution name (country code)	GH institution name (country code)	GH institution name (country code)	
Meetings	EUR 53 697,49	EUR 53 506,29	-	EUR 107 203,78
Training Schools	EUR -	EUR -	-	EUR -
STSMs	EUR 13 250,00	EUR 4 800,00	-	EUR 18 050,00
Dissemination	EUR -	EUR 1 325,00	-	EUR 1 325,00
OERSA ¹	EUR -	EUR -	-	EUR -
Total Scientific Expenditure	EUR 66 947,49	EUR 59 631,29	-	EUR 126 578,78
FSAC ²	EUR 10 042,12	EUR 8 922,69	-	EUR 18 964,81
TOTAL	EUR 76 989,61	68 553,98 EUR	-	EUR 145 543,59

¹ OERSA = Other Expenses Related to Scientific Expenditure (e.g. bank charges)

² FSAC = Amount received by Grant Holder for Financial Scientific and Administrative Coordination

II.B. Budget and Participation management

II.B.1 Budget spent in relation to individuals/ institutions outside participating COST countries					
STSMs from or to institutions from countries other than Participating COST countries					
We had no activity in this case.					
Grantee		Host		Date	Topic and value added to the Action
Institution	Country	Institution	Country		
Add home institution and country		Add host institution and country		Date	Describe topic of the STSM and the added value to the Action
Add home institution and country		Add host institution and country		Date	Describe topic of the STSM and the added value to the Action
Add home institution and country		Add host institution and country		Date	Describe topic of the STSM and the added value to the Action
<i>Invited Speakers</i>					
The table below highlights the added value of Invited Speakers from COST countries that have not accepted the MoU and/ or non-participating NNC, IPC or Specific Organisations whose participation at a meeting or Training School was reimbursed by the Action.					
Participant name	Institution	Country	Event date	Topic and added value to the Action	
Thomas de Coursey	Department of Physiology & Biophysics, Rush Medical College, 1750 W. Harrison St. Suite 1245 Chicago, IL 60612	USA	9-11 th March 2016	BM1406 Lisbon Oral presentation invited by N. Demareux (Swiss) https://www.rushu.rush.edu/research/departamental-research/physiology-and-biophysics-research/laboratory-tom-decoursey-phd	
Stefan Feske	Department of Pathology and Cancer Institute NYU School of Medicine, Smilow Research Building 550 First Avenue, New York, NY 10016, USA	USA	23-24 th March 2017	BM1406 meeting in Belgrade, conference on Channelopathies invited by Ruth Murrell-Lagnado (UK) Event Facebook https://med.nyu.edu/faculty/stefan-feske	
<i>Dissemination meetings</i>					
The table below highlights the added value of Dissemination Meetings financed from Action funds.					
Participant name	Role	Country	Date	Location	Topic and added value to the Action
Dr. Anna Rubartelli	speaker	Italy	4-7 October 2016	Lecce, Italy	Unconventional proteins and membrane traffic, Presentation on COST Action (slides)

II.C. Participants

Management Committee		
Name	Country	Email address
Prof Christoph ROMANIN	Austria	christoph.romanin@jku.at
Prof Elena E. POHL	Austria	Elena.Pohl@vetmeduni.ac.at
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Prof Fotini PALIOGIANNI	Greece	fpal@upatras.gr
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Prof Gyorgy PANYI	Hungary	panyig@gmail.com
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Ms Zaiga NORA-KRUKLE	Latvia	Zaiga.Nora@rsu.lv
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II.D. Specific matters

This section is confidential to the Management Committee, and the COST Association (Administration, Scientific Committee and Committee of Senior Officials); and is not included in the version of the report that is made publicly available.

The Action encountered the following particular difficulties in the implementation of the Action (e.g. imbalances of participation across the Working Groups, inactive country representatives).
Describe the issue(s) here or write "no particular difficulties encountered".
no particular difficulties encountered
Write explanation here

Annex 1

Definitions:

COST Action Challenge (main aim)	“The research question addressed by the COST Action targeting scientific, technological, and / or socioeconomic problems”
COST Action Innovation	“The creation and / or development of new or improved concepts, products, processes, services, and / or technologies that are made available to markets, governments and society”
COST Action objectives	“COST Action objectives are the results that an Action needs to achieve in order to respond to meet its challenge. These are SMART (Specific, Measurable, Achievable, Relevant, Timely) and twofold: research coordination objectives and capacity building objectives.”
COST Action research coordination objectives	“Achieving these objectives turns COST Actions from initially scattered teams into one transnational team and leverages the existing funded research. These objectives entail the distribution of tasks, sharing of knowledge and know-how, and the creation of synergies among Action participants to achieve specific outputs.”
COST Action capacity building objectives	“Achieving these objectives entail building critical mass to drive scientific progress, thereby strengthening the European Research Area. They can be achieved by the delivery of specific outputs and / or through network features or types and levels of participation.”
COST Action networking activities	“any activities organised by the COST Action (whether or not directly funded by COST) in order to achieve research coordination and capacity building objectives.”
COST Action networking tools	“instruments through which eligible activities can be funded”
COST Action outputs	“direct results from the COST Action activities. These can be codified knowledge, tacit knowledge, technology, and societal applications.”
COST Action impact	“the short- to long-term scientific, technological, and / or socioeconomic changes produced by a COST Action, directly or indirectly, intended or unintended.”
COST Action deliverable	“a distinct, expected and tangible output of the Action, meaningful in terms of the Action’s overall objectives such as a report, a document, a technical diagram, a software etc. Action deliverables are used to measure its progress and success.”
COST Action milestones	“Control points in the Action that help to chart progress. They are also needed at intermediary points so that, if problems have arisen, corrective measures can be taken. A milestone may be a critical decision point in the Action where, for example, the MC must decide which of several technologies to adopt for further development (e.g. core group and MC meetings, mid-term reviews)”
Inclusiveness Target Country (ITC):	Current COST Member Countries targeted by the COST inclusiveness Policy (“Inclusiveness Target Countries” (ITC)): EU 13 (Bulgaria, Cyprus, Czech Republic, Estonia, Croatia , Hungary, Lithuania, Latvia, Malta, Poland , Romania, Slovenia, Slovakia), EU candidate countries (the former Yugoslav Republic of Macedonia, Montenegro, Republic of Serbia, Turkey) and potential EU candidate countries (Bosnia and Herzegovina). In addition, to comply with the EC criteria for ‘Spreading Excellence and Widening Participation’, Portugal and Luxemburg are included.