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Interview with new President-Elect

ECCO EpiCom Survey – Registries across Europe Page 20-21

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European Crohn's and Colitis Organisation

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- Quarterly ECCO News The society's magazine
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- Guidelines, ECCO Fellowships, Grants and Travel Awards
- Access to ECCO UR-CARE United Registries for Clinical Assessment and Research

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### ECCO NEWS

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# Dear ECCO Friends,

Summer is a wonderful period of the year: daily activities slow down a little, giving us the time to think bigger. The ECCO Committees and Governing Board also take this healthy approach to the life of our Organisation, meeting for two or three days in Vienna in June and July, in a constructive atmosphere, to set the strategies and plans to fulfill our mission. I hope that all of you enjoyed some time off during the summer and were able to "think big" about all aspects of life.

Now we have returned to our usual activities, but not to the old routine, because, as you will see in this issue of ECCO News, there are a lot of exciting items of news and stimulating future activities await.



Julián Panés © ECCO

One of the highlights is the *programme for our ECCO Congress 2017*, with an overview of the Scientific Programme and the Educational Programme. Many of the educational activities in the Congress are

booked well in advance, and you should have a look right now and secure your registration if you wish to attend. You will also find the Call for Abstracts: start planning the original work that you wish to submit to the ECCO Congress.

I want to draw your attention to the information on the activities that N-ECCO is organising at ECCO '17. This is relevant not only to nurses: physicians should also be aware of these European educational activities, and actively promote and facilitate the participation of nurses involved in IBD care. Nurses are at the centre of any well-organised IBD unit and we should all strive for their promotion and recognition.

*In terms of quality improvement*, you will find in this issue interesting information on quality indicators for running a paediatric Inflammatory Bowel Disease Centre.

Finally, on a more personal note, you will find an *interview with our ECCO President-Elect*, Silvio Danese, in which he reveals his vision for the organisation and the opportunities that he sees to improve education and research and, as a result, the care or our IBD patients.

Enjoy ECCO News!

JULIÁN PANÉS ECCO President

# Call for Abstracts for the 12<sup>th</sup> Congress of ECC

To submit an abstract for the 12<sup>th</sup> Congress of ECCO, please use our online abstract submission system. Please also view important information on the submission process and the guidelines for abstract submission. **Presentation format** 

• The 38 best abstracts (up from 30 in 2016) will receive an oral presentation slot in the scientific programme of the 12<sup>th</sup> Congress of ECCO.

• The next best ~90 abstracts will be digital oral presentations, with a 5 minute oral presentation on either Thursday, February 16, 2017 from 17:45-18:45 or on Friday, February 17, 2017 from 17:20-18:20.

• The remaining accepted abstracts will be displayed as hard copy posters throughout the Congress. Please find further details in the guidelines for presentation.

### Important note

There will be NO late-breaking abstracts, so please aim to get your abstract in on time! We look forward to welcoming you to the ECCO Congress in Barcelona, Spain on February 15-18, 2017!

Kind regards,

### SILVIO DANESE, PETER LAKATOS, CHARLIE LEES, LAURENT PEYRIN-BIROULET On behalf of the ECCO'17 Barcelona Organising Committee

JULIÁN PANÉS ECCO President and Chair of the Organising Committee

Key dates

August 17, 2016 November 3, 2016 December 1, 2016 February 1, 2017 February 15-18, 2017

Opening of abstract submission Deadline for early registration (midnight, CET) Deadline for abstract submission (midnight, CET) By December 23, 2016 Notification of abstract acceptance/rejection Deadline for late registration (after that date onsite registration only) 12<sup>th</sup> Congress of ECCO, Barcelona, Spain

# Scientific Programme at ECCO'17 ECCO'17 Theme: "Advancing Knowledge, Improving Care"

### as of September 23, 2016

Prelimina	ry program	me: Thursday, February 16, 2017			
	10:45 - 11:15	Top tips for chairs (closed session)		16:00 - 16:10	Oral presentation 6
11:30 - 12:30		Laurence Egan, Galway, Ireland nsored satellite symposia 1a & 1b		16:10 - 16:30	Personalised medicine: Dream or reality? Iris Dotan, Tel Aviv, Israel
	Welcome	insoled satellite symposia ia & ib		Scientific see	ssion 3: Vulnerable patients: 10 burning
12:45 - 12:50		t, Madrid, Spain	16:30 - 17:30	questions in	IBD
12:50 - 13:00	<b>Opening</b> Julián Panés,	Barcelona, Spain			tti, Lausanne, Switzerland ón, Barcelona, Spain
13:00 - 14:30	Scientific ses everyday po Stephan Vavr	sion 1: Pathways of environmental factors or isons and IBD icka, Zurich, Switzerland e Andersen, Copenhagen, Denmark		16:30 - 16:42	The child with IBD: Should anti-TFN be used as a mono-therapy in children (safety risk)? Who should (be allowed) to operate on the children in IBD? Dan Turner, Jerusalem, Israel
	13:00 - 13:20	Nutrition and Food additives Charlie Lees, Edinburgh, United Kingdom			Treating the pregnant woman with IBD: Should anti-TFN be stopped during
	13:20 - 13:30	Oral presentation 1		16:42 - 16:54	pregnancy? Should Vedolizumab be stopped during pregnancy?
	13:30 - 13:50	Everyday poisons: Smoking, pollution, stress, sedentary lifestyle Stephan Vavricka, Zurich, Switzerland			Janneke van der Woude, Rotterdam, The Netherlands
	13:50 - 14:00	Oral presentation 2			The elderly patient with IBD. Should Azathiorprine ever be used in the elderly?
	14:00 - 14:10	Oral presentation 3		16:54 - 17:06	Should the type of surgery be different in
	14:10 - 14:30	Environmental factors as therapy (diet modification; EEN feeding) Pierre Ellul, Sliema, Malta			an elderly patient? Jean-Frédéric Colombel, New York, United States
		Arie Levine, Tel Aviv, Israel			The use of IBD medicine in patients with cancer. Which treatment should
14:30 - 15:00	Coffee break			17:06 - 17:18	be stopped when cancer is diagnosed? Which treatment to use in a patient with
15:00 - 16:30	2017	ssion 2: Evolving therapeutic algorithms in rs, Adelaide, Australia			a prior diagnosis with cancer? Laurent Beaugerie, Paris, France
		kuś, Warsaw, Poland			The patient with psychological comorbidities; How to wean off opiates
	15:00 - 15:20	Placing new molecules in the treatment pathway Gerhard Rogler, Zurich, Switzerland		17:18 - 17:30	in these patients? Is there any preference in drugs? Jane Andrews, Adelaide, Australia
	15:20 - 15:30	Oral presentation 4	17:45 - 18:45	Industry spo	onsored satellite symposia 2a & 2b
	15:30 - 15:50	Maximizing the bang for your buck: How to manage IBD on a minimum budget Peter Irving, London, United Kingdom	17:45 - 18:45	Digital Oral I	Presentations (sessions 1-5)
	15:50 - 16:00	Oral presentation 5			

Prelimina	ry program	me: Friday, February 17, 2017			
07:15 - 08:15	Industry spo	nsored satellite symposia 3a & 3b		14:00 - 14:20	Microscopic colitis
		sion 4: Clinical trials versus real life		14:20 - 14:30	Oral presentation 17
08:30 - 10:30		der Woude, Rotterdam, The Netherlands at, Lausanne, Switzerland		14:30 - 14:40	Oral presentation 18
	08:30 - 08:50	Trial endpoint vs. therapeutic objectives		14:40 - 15:00	Eosinophilic enteritis
		Filip Baert, Roeselare, Belgium	EITHER:		
	08:50 - 09:00	Oral presentation 7	15:00 - 15:40	Scientific ses	sion 7: ECCO Fellowships & Grants and, Berlin, Germany
	09:00 - 09:10	Oral presentation 8	13.00 13.10	Gionata Fiori	no, Milan, Italy
	09:10 - 09:20	Oral presentation 9		15:00 - 15:07	Outcomes from the 2015-6 Fellowships Ferdinando Bonfiglio, Stockholm, Sweden
	09:20 - 09:40	The long-term efficacy outside clinical trials Stefan Schreiber, Kiel, Germany		15:07 - 15:14	Outcomes from the 2015-6 Fellowships
	09:40 - 09:50	Oral presentation 10			Jesús Cosín, Zurich, Switzerland Announcement of ECCO Fellowships &
	09:50 - 10:00	Oral presentation 11		15:14 - 15:20	Grants 2017
	10:00 - 10:10	Oral presentation 12		15:20 - 15:30	Oral presentation 19
	10:10 - 10:30	The long-term safety outside clinical trials Gert van Assche, Leuven, Belgium		15:30 - 15:40	Oral presentation 20
10:30 - 11:00	Coffee break		15:40 - 16:10	Coffee break	ζ.
11:00 - 12:30	Surgical or m	ision 5: Managing bowel damage in IBD: nedical treatment? nelli, Milan, Italy	16:10 - 17:10	Bruce Sands,	s <b>sion 8: IBD Horizons</b> New York, United States Innsbruck, Austria
	Edyta Zagóro	wicz, Warsaw, Poland		16:10 - 16:20	Oral presentation 21
	11:00 - 11:20	Are we truly changing disease course with current strategies?		16:20 - 16:30	Oral presentation 22
	11.00 11.20	Yves Panis, Clichy, France		16:30 - 16:40	Oral presentation 23
	11:20 - 11:30	Oral presentation 13		16:40 - 16:50	Oral presentation 24
	11.20 11.00	Combined approach for intestinal sparing		16:50 - 17:00	Oral presentation 25
	11:30 - 11:50	<b>in CD?</b> Paulo Kotze, Curitiba, Brazil		17:00 - 17:10	Oral presentation 26
	11:50 - 12:00	Oral presentation 14	OR:		
	12:00 - 12:10	Oral presentation 15 The ileorectal anastomosis in UC: Time for	15:00 - 17:10	Arthur Kaser,	cambridge, United Kingdom
	12:10 - 12:30	a comeback?			is, Barcelona, Spain
	Lunchbrook	Oded Zmora, Tel Aviv, Israel and guided Poster Session in the			IL-23 and beyond
12:30 - 13:30	exhibition ha	all		15:20 - 15:30	Oral presentation 27
12:40 - 13:20	Industry spo	nsored educational lunchtime satellite		15:30 - 15:40	Oral presentation28
	symposia LS		15:40 - 16:10	Coffee break	
	<b>always IBD</b> Gijs van den	ssion 6: Intestinal inflammation – not Brink, Amsterdam, The Netherlands		16:10 - 16:30	Genetics and function in IBD: The missing link? Sebastian Zeissig, Dresden, Germany
13:30 - 15:00	Carousel: (ea	, Tel Áviv, Israel ch topic has these four people presenting		16:30 - 16:40	Oral presentation 29
15.50 15.00	2-3 slides each) Epidemiologist: Vito Annese, Florence, Italy			16:40 - 16:50	Oral presentation 30
	Basic Science Pathologist: F	Basic Science: Britta Siegmund, Berlin, Germany Pathologist: Roger Feakins, London, United Kingdom Physician: James Lindsay, London, United Kingdom		16:50 - 17:10	Epigenetics in complex diseases Manel Esteller, Barcelona, Spain
	13:30 - 13:50	IBDU	17:20 - 18:20	Digital oral p	presentations (sessions 6-10)
		Oral presentation 16	18:30 - 19:30	Industry spo	nsored satellite symposia 4a & 4b
	.5.55 11.00	2.2. p. cochadon io			

Preliminary programme: Saturday, February 18, 2017				
07:15 - 08:15	Industry spo	Industry sponsored satellite symposia 5a & 5b		
08:30 - 10:20	of IBD John Mansfie	Scientific session 10: New compounds in the treatment of IBD John Mansfield, Newcastle upon Tyne, United Kingdom Pieter Hindryckx, Ghent, Belgium		
	08:30 - 08:50	<b>JAK-inhibitors</b> Séverine Vermeire, Leuven, Belgium		
	08:50 - 09:00	Oral presentation 31		
	09:00 - 09:10	Oral presentation 32		
	09:10 - 09:30	<b>S1P</b> Arthur Kaser, Cambridge, United Kingdom		
	09:30 - 09:40	Oral presentation 33		12:20 -
	09:40 - 09:50	Oral presentation 34		
	09:50 - 10:00	Oral presentation 35		
	10:00 - 10:20	Anti IL12/IL23 compounds Bruce Sands, New York, United States		
10:20 - 10:50	Coffee break			12:50
10:50 - 12:20	just bacteria Isabelle Cleyr	Scientific session 11: Gut luminal content – more than		

	10:50 - 11:10	The forgotten populations in the gut Xavier Ramnik, Boston, United States
	11:10 - 11:20	Oral presentation 36
	11:20 - 11:40	Metabolic activity Dirk Haller, Freising, Germany
	11:40 - 11:50	Oral presentation 37
	11:50 - 12:00	Oral presentation 38
	12:00 - 12:20	Microbiome reality check – looking beyond the hype Jeroen Raes, Gent, Belgium
12:20 - 12:50		<b>sion 12: ECCO Lecture</b> Barcelona, Spain Milan, Italy
	12:20 - 12:50	Best individualised IBD care – optimising drugs, psychology, diet and microbes Michael Kamm, Melbourne, Australia
12:50 - 12:55		c <b>losing remarks</b> Barcelona, Spain
12:55 - 13:00	The ECCO Film	m 2017

# Educational Programme at ECCO'17 as of September 23, 2016

he educational programme of the 12<sup>th</sup> Congress of ECCO starts prior to the official start of the ECCO Congress and courses take place from February 15-17, 2017. These activities are targeted towards ECCO's different interest groups including young gastroenterologists, surgeons, paediatricians, IBD nurses and allied health professionals and scientists.

An overview of these activities can be found below. Please note that some of these courses/ workshops run in parallel and that some have a limited capacity – please do register at your earliest convenience.

We look forward to seeing you in Barcelona!

	nesday y 15, 2017	Thurs February 1		Friday February 17, 2017		Saturday, February 18, 2017	
Morning	Afternoon	Morning	Afternoon	Morning	Afternioon	Morning	
15 <sup>n</sup> IBD Intensive Advanced Course	10 <sup>h</sup> Y-ECCO Careir Workshop	15" IBD Intensive Advanced Course			Programme exhibition		
8º N-EO	CO School	4° ECCO Ultrasound Workshop		Industry	exhibition		
2 <sup>nd</sup> School for Clinical Trialists	4" N-ECCO Research Forum	4º P-ECCO Educational Course	Digital Oral Presentation Sessions 1-5	2 <sup>se</sup> D-ECCO Workshop	Digital Oral Presentation Sessions 6-10		
	3rd Y-BCCO Basic Science Workshop	AOCC-ECCO Forum	2"H-ECCO IB	D Masterclass	ECCO Interaction Hearts & Minds		
	3 <sup>rd</sup> Advanced ECCO: EduCational COurse for Industry	11" N-ECCO Net	work Meeting			*:	
	2 <sup>rd</sup> ECCO Endoscopy Workshop	6* S ECCO IBD					
		5º SciCom Workshop Methodology on Research					
		EC	CO Business Meeting	р			

### Preliminary programme: 15<sup>th</sup> IBD Intensive Advanced Course Wednesday, February 15, 2017

07:45 - 08:00		distribution of voting pads	12:30 - 14:00	Session 3: Se Part I: Specia	eminars al clinical situations
08:00 - 08:10	Welcome Julián Panés, Barcelona, Spain Gerassimos Mantzaris, Athens, Greece			. aren opeen	EITHER: I.a. Managing IBD and pregnancy Janneke van der Woude, Rotterdam,
08:10 - 08:30	Pre-course test Peter Irving, London, United Kingdom			12.20 12.15	The Netherlands Iris Dotan, Tel Aviv, Israel
08:30 - 09:30	Session 1: Pathogenesis Lead discussant: James Lindsay, London, United Kingdom			12:30 - 13:15	OR: I.b. Managing extraintestinal manifestations of IBD
	08:30 - 08:50	<b>Exposome</b> Jonas Halfvarsson, Örebrö, Sweden			Stephan Vavricka, Zürich, Switzerland Peter Lakatos, Budapest, Hungary
	08:50 - 09:10	<b>Genetics</b> Miles Parkes, Cambridge, United Kingdom			EITHER: I.a. Managing IBD and pregnancy Janneke van der Woude, Rotterdam, The Netherlands
	09:10 - 09:30	<b>Inflammatory pathways</b> Yehuda Chowers, Haifa, Israel		13:15 - 14:00 Iris Dotan, Tel Aviv, Israel OR:	
09:30 - 09:50	Coffee break				I.b. Managing extraintestinal manifestations of IBD
09:50 - 11:10	Part I: Conve	rug management sessions entional drugs ant: Pascal Juillerat, Bern, Switzerland	14:00 - 14:15	Stephan Vavricka, Zürich, Switzerland Peter Lakatos, Budapest, Hungary Coffee break	
	09:50 - 10:10	5-ASA compounds Gerhard Rogler, Zürich, Switzerland	14:15 - 15:45	Session 3: Seminars	
	10:10 - 10:30	Thiopurines Peter Irving, London, United Kingdom		Part II: Long	term management EITHER: II.a. Managing complications associated
	10:30 - 1 0:50	Methotrexate Pascal Juillerat, Bern, Switzerland			with anti-TNF therapy Shomron Ben-Horin, Ramat Gan, Israel
	10:50 - 1 1:10	<b>Steroids</b> Stephan Vavricka, Zürich, Switzerland		14:15 - 15:00 OR: II.b. Perform endoscopy and IBD in	
11:10 - 12:00	Part II: Biolo	rug management sessions gics ant: Peter Irving, London, United Kingdom			<b>chromo-endoscopy, bálloon dilatation</b> Pierre Michetti, Lausanne, Switzerland Marc Ferrante, Leuven, Belgium
	11:10 - 11:30	Anti-TNF agents Filip Baert, Roeselare, Belgium			EITHER: II.a. Managing complications associated with anti-TNF therapy
	11:30 - 11:45	Vedolizumab James Lindsay, London, United Kingdom		15:00 - 15:45	Shomron Ben-Horin, Ŕamat Gan, Israel OR:
	11:45 - 12:00	<b>Ustekinumab</b> Marc Ferrante, Leuven, Belgium			II.b. Perform endoscopy and IBD incl. chromo-endoscopy, balloon dilatation Pierre Michetti, Lausanne, Switzerland
12:00 - 12:30	Lunch break	ζ			Marc Ferrante, Leuven, Belgium

# Preliminary programme: 15<sup>th</sup> IBD Intensive Advanced Course Thursday, February 16, 2017

08:00 - 10:00		teractive case discussions ant: Peter Irving, London, United Kingdom	10:20 - 12:15		<b>pecial cases scenarios</b> ant: Pascal Juillerat, Bern, Switzerland
	Case-based discussion: Fistulising Crohn's Disease: Medical and surgical approaches			10:20 - 10:40	Pre- and postoperative management of Crohn's Disease Glen Doherty, Dublin, Ireland
		Case presentation: Antonio López- Sanromán, Madrid, Spain Discussion: Paulo Kotze, Curitiba, Brasil			Case-based discussion: Investigation and management of mild/moderate Crohn's Disease
	08:50 - 09:00	Discussion		10:40 - 11:25	Case presentation: Timothy Raine,
		Case-based discussion: The patient with severe inflammatory Crohn's Disease			Cambridge, United Kingdom Discussion: Jane Andrews, Adelaide, Australia
	09:00 - 09:50	Case presentation: Pieter Hindryckx, Ghent, Belgium		11:25 - 11:45	Management of refractory pouchitis Iris Dotan, Tel Aviv, Israel
		Discussion: Laurence Egan, Galway, Ireland		11:45 - 12:15	Medical management of acute severe Ulcerative Colitis
	09:50 - 10:50	Discussion			Charlie Lees, Edinburgh, United Kingdom
10:00 - 10:20	Coffee break	(	12:15 - 12:30	Peter Irving,	<b>nd closing remarks</b> London, United Kingdom at, Bern, Switzerland
Responsible Committee: EduCom Farget audience: Junior gastroenterologists			ECCO Members Registration fee	hip 2017 requ	at, Bern, Switzerland iired: Regular/Y-ECCO Member

Target audience: Junior gastroenterologists Registration: Upon invitation, please see official call on page 6

# Preliminary programme: 8<sup>th</sup> N-ECCO School Wednesday, February 15, 2017

08:30 - 08:45	Welcome and introduction Usha Chauhan, Hamilton, Canada		13:15 - 14:50	Kay Grevesor	ase studies – Disease management n, London, United Kingdom
08:45 - 12:15	Session 1: Diagnosis and assessment Usha Chauhan, Hamilton, Canada Nicolette Wierdsma, Amsterdam, The Netherlands			13:15 - 14:05	erdsma, Amsterdam, The Netherlands Workshop 1 – UC Management (Group A) Andreas Sturm, Berlin, Germany
	08:45 - 09:30	Diagnosis, anatomy and physiology in IBD			Workshop 2 – CD Management (Group B Pieter Hindryckx, Ghent, Belgium
		Marc Ferrante, Leuven, Belgium			Workshop 1 – UC Management (Group B) Andreas Sturm, Berlin, Germany
	09:30 - 10:00	Psychosocial implications of living with IBD Kay Greveson, London, United Kingdom		14:05 - 14:50	Workshop 2 – CD Management (Group A Pieter Hindryckx, Ghent, Belgium
		Nutritional assessment in IBD	14:50 - 15:05	Coffee brea	k
	10:00 - 10:30	Konstantinos Gerasimidis, Glasgow, United Kingdom	15:05 - 16:05	Session 3: Multidisciplinary management in IBD Kay Greveson, London, United Kingdom	
10:30 - 10:45	Coffee breal	<		NICOlette VVie	erdsma, Amsterdam, The Netherlands
	10:45 - 11:15	<b>Surgery in IBD</b> Janindra Warusavitarne, London, United		15:05 - 15:35	Nutritional management in IBD Rotem Sigall-Boneh, Tel Aviv, Israel
		Kingdom		15:35 - 16:05	Nursing roles in IBD management Kathleen Sugrue, Cork, Ireland
	11:15 - 11:45	Medical treatment Konstantinos Katsanos, Ioannina, Greece	16:05 - 16:15	Closing rem	arks
	11:45 - 12:15	<b>Adherence</b> Palle Bager, Aarhus, Denmark	10.03 - 10.13	Usha Chauh	an, Hamilton, Canada
12:15 - 13:15	Lunch break	· · · · · · · · · · · · · · · · · · ·			
Responsible C Farget audien Registration: U	<b>ce:</b> IBD nurses	<ul> <li>new to the specialty, Dietitians</li> </ul>	ECCO Members Registration fe		uired: IBD nurse Member, Affiliate Member

### 2<sup>nd</sup> School for Clinical Trialists – Understanding the different types of clinical trials Wednesday, February 15, 2017

weanesday	, February	15, 2017			
08:00 - 08:15	Welcome and introduction Alessandro Armuzzi, Rome, Italy		10:00 - 11:20	Session 2 Javier Gisbert	, Madrid, Spain
08:15 - 09:30	<b>Session 1</b> Edyta Zagorowicz, Warsaw, Poland			10:00 - 10:30	Setting up and running large nationwide IBD trials
	08:15 - 09:00	<b>Clinical trial design and sample size</b> Timothy Raine, Cambridge, United Kingdom		10:30 - 11:00	Laurent Beaugerie, Paris, France Tips & tricks for the IBD clinical research team
	09:00 - 09:30	How to optimise recruitment to clinical trials in IBD		10.50 - 11.00	Liese Gijbels, Leuven, Belgium Karen Rans, Leuven, Belgium
09:30 - 10:00	Coffee break	Charlie Lees, Edinburgh, United Kingdom		11:00 - 11:20	What does the future hold for IBD clinical trials? Silvio Danese, Milan, Italy
			11:20 - 11:30	Summary & John Mansfie	<b>closing remarks</b> Id, Newcastle upon Tyne, United Kingdom
Responsible Committee: ClinCom Target audience: Clinical trial coordinators, IBD nurses and Allied health professionals				nip 2017 require	on • <b>d:</b> Regular/Y-ECCO/IBD nurse/Affiliate Member . 21% Spanish VAT

### 4<sup>th</sup> N-ECCO Research Forum Wednesday, February 15, 2017

13:00 - 13:10		<b>d introduction</b> Aarhus, Denmark	14:00 - 14:30	International IBD nursing research projects
	J .	ethodology: Examples from current	14:30 - 15:00	Coffee break
13:10 - 14:00	research	thousings. Examples nom current	15:00 - 16:25	Workshop: International research projects – 'pre investigators meetings'
	13:10 - 13:25	Systematic review		pre investigators meetings
	13.10 13.23	Dawn Farrell, Cork, Ireland	16:25 - 16:45	Short presentation of status from the workshops
	13:25 - 13:45	<b>Development of a questionnaire</b> Wladzia Czuber-Dochan, London, United Kingdom	16:45 - 17:00	<b>Learning from today: How to proceed?</b> Palle Bager, Aarhus, Denmark
	13:45 - 14:00	Measurement of conditions Randi Opheim, Oslo, Norway		
Responsible Committee: N-ECCO Target audience: IBD nurses and Allied health professionals			hip 2017 required: IBD nurse Member, Affiliate Member e: EUR 15,- incl. 21% Spanish VAT	

**Registration:** Online registration

### 2<sup>nd</sup> ECCO Endoscopy Workshop Wednesday, February 15, 2017

13:00 - 13:15	Welcome and introduction Julián Panés, Barcelona, Spain Gerassimos Mantzaris, Athens, Greece Pre-course test
13:15 - 14:15	Session 1: Assessment of endoscopic activity: Clinicaltrials and routine practiceChairs:Alessandro Armuzzi, Rome, Italy Konstantinos Katsanos, Ioannina, GreeceSpeaker:James Lindsay, London, United Kingdom
14:15 - 15:15	Session 2: Endoscopic Surveillance for IBD associatedcolorectal cancerChairs:Antonio López-Sanromán, Madrid, Spain James Lindsay, London, United KingdomSpeaker:Vito Annese, Florence, Italy
Target audiend	ommittee: EduCom <b>:e:</b> Physicians, Surgeons, Paediatricians )nline registration (max. 50 participants)

15:15 - 1	5.45	Coffee break					
15:45 - 1		Session 3: Endoscopic therapeutic intervention in IBD Chairs: Peter Irving, London, United Kingdom Marc Ferrante, Leuven, Belgium Speaker: Nik Ding, London, United Kingdom					
16:45 - 1	17:45	Session 4: Small bowel endoscopy: Capsule vs.         balloon enteroscopy         Chairs:       Peter Lakatos, Budapest, Hungary         Edyta Zagorowicz, Warsaw, Poland         Speaker:       Shomron Ben-Horin, Ramat-Gan, Israel					
17:45 - 1	17:45 - 18:00 Post-course test & concluding remarks Gerassimos Mantzaris, Athens, Greece						
	ECCO Membership 2017 required: Regular/Y-ECCO Member Registration fee: € 80 (half price for Y-ECCO and IBD nurse Members) –						

incl. 21% Spanish VAT

Panel discussion:

Panel discussion:

**Closing remarks** 

Silvio Danese, Milan, Italy

Non-Corporate Members: EUR 600.- incl. 21% Spanish VAT

Corporate Members: EUR 400.- incl. 21% Spanish VAT

(Pfizer)

16:15 - 17:05

17:05 - 17:55

17:55 - 18:00

**Registration fee:** 

Session 3: Learning for GI from other IMIDs Bruce Sands, New York, United States

Chris Gasink, Malvern, United States (Janssen) Wojciech Niezychowski, Collegeville, United States

Elmer Schabel, Bonn, Germany (EMA) Session 4: Treatment, strategies, trials Peter Irving, London, United Kingdom

Gerd-Rüdiger Burmester, Berlin, Germany (EULAR)

John Mansfield, Newcastle upon Tyne, United Kingdom Ana Lacerda, Chicago, United States (AbbVie) Rebecca Curtis, London, United Kingdom (Takeda)

### 3<sup>rd</sup> Advanced ECCO: EduCational COurse for Industry Wednesday, February 15, 2017

14:00 - 14:05	<b>Welcome</b> Julián Panés, Barcelona, Spain
14:05 - 14:55	Session 1: Preclinical models in IBD drug development Silvio Danese, Milan, Italy Panel discussion: Gerhard Rogler, Zürich, Switzerland Andy Whitney, Foster City, United States (Gilead) Philippe Clement-Lacroix, Romainville, France (Galapagos)
14:55 - 15:45	Session 2: Proof of concept studies Séverine Vermeire, Leuven, Belgium Panel discussion: Stefan Schreiber, Kiel, Germany Allan Olson, San Diego, United States (Celgene) Miguel Forte, Valbonne-Sophia Antipolis, France (TxCell)
15:45 - 16:15	Coffee break

Responsible Committee: ClinCom

Target audience: Corporate Members & Non-Corporate Members Registration: Please contact the ECCO Office at ecco17@ecco-ibd.eu ECCO Membership 2017 required: n.a.

### 3<sup>rd</sup> Y-ECCO Basic Science Workshop Wednesday, February 15, 2017

, reallestate	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	13/ 2017				
12:30 - 12:35	Introduction Isabelle Cleynen, Leuven, Belgium		14:10 - 15:25	Session 2: Identifying novel targets in IBD using patient samples		
12:35 - 13:50	Session 1: Genetics in IBD – moving beyond GWAS Miles Parkes, Cambridge, United Kingdom		11.10 13.25	Gijs van den Brink, Amsterdam, The Netherlands Nik John Ding, London, United Kingdom		
	Dominik Bet	tenworth, Münster, Germany		14:10 - 14:40	Gijs van den Brink, Amsterdam, The Netherlands	
	12:35 - 13:05	Miles Parkes, Cambridge, United			The Netherlands	
	12.55 - 15.05	Kingdom		14:40 - 14:55	Selected oral presentation 4	
	13:05 - 13:20	Selected oral presentation 1		14:55 - 15:10	Selected oral presentation 5	
	13:20 - 13:35	Selected oral presentation 2		15:10 - 15:25	Selected oral presentation 6	
	13:35 - 13:50	Selected oral presentation 3	15:25 - 15:30		arks and basic science abstract awards	
13:50 - 14:10	Meet the sp	eakers break		Isabelle Cley	nen, Leuven, Belgium	
Responsible Committee: Y-ECCO Target audience: Physicians, Paediatricians, Surgeons, IBD nurses		ECCO Membership 2017 required: Regular/Y-ECCO/IBD nurse/Affiliate Member Registration fee: EUR 80 (half price for Y-ECCO Affiliate and IBD nurse)				
Registration: Online registration Member			Members) – incl	Members) – incl. 21% Spanish VAT		

### 10<sup>th</sup> Y-ECCO Career Workshop – Successful grant writing skills Wednesday, February 15, 2017

,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,					
Welcome and introduction         Nuha Yassin, London, United Kingdom         Session 1: Grant Writing – How to get started         Isabelle Cleynen, Leuven, Belgium         Dominik Bettenworth, Münster, Germany				17:00 - 17:35	Feedback on projects and announcement of winning project
					Séverine Vermeire, Leuven, Belgium Gerhard Rogler, Zürich, Switzerland Helen Terry, St. Albans, United Kingdom
16:10 - 16:25 <b>Designing the ideal project</b> Séverine Vermeire, Leuven, Belgium			17:35 - 18:00	Session 3: Grant Writing – Examples of successful projects Nik John Ding, London, United Kingdom	
16:25 - 16:40	Grant Writing – Tips and Tricks Gerhard Rogler, Zürich, Switzerland				Examples of funded IBD projects by charities
Session 2: Gr	ion 2: Group Session – Designing a successful			17.55 17.50	Helen Terry, St. Albans, United Kingdom
project Timothy Raine, Cambridge, United Kingdom Dominik Bettenworth, Münster, Germany Isabelle Cleynen, Leuven, Belgium				17:50 - 18:00	Thank you and Prizes Nuha Yassin, London, United Kingdom Timothy Raine, Cambridge, United Kingdom
16:40 - 17:00	Group session – Design project suitable		18:00	Close - Y-ECC	CO Networking
5					
Responsible Committee: Y-ECCO Target audience: Physicians, Paediatricians, Surgeons, IBD nurses Registration: Online registration		F	ECCO Membership 2017 required: Regular/Y-ECCO/IBD nurse/Affiliate Memb Registration fee: EUR 80 (half price for Y-ECCO, IBD nurse and Affilia Members) - incl. 21% Spanish VAT		
	Welcome an Nuha Yassin, Session 1: Gr Isabelle Cley Dominik Bet 16:10 - 16:25 16:25 - 16:40 Session 2: Gr project Timothy Rair Dominik Bet Isabelle Cley Nuha Yassin, 16:40 - 17:00 ommittee: Y-Fice: Physicians,	Nuha Yassin, London, United Kingdom         Session 1: Grant Writing – How to get started         Isabelle Cleynen, Leuven, Belgium         Dominik Bettenworth, Münster, Germany         16:10 - 16:25       Designing the ideal project Séverine Vermeire, Leuven, Belgium         16:25 - 16:40       Grant Writing – Tips and Tricks Gerhard Rogler, Zürich, Switzerland         Session 2: Group Session – Designing a successful project         Timothy Raine, Cambridge, United Kingdom         Dominik Bettenworth, Münster, Germany         Isabelle Cleynen, Leuven, Belgium         Nuha Yassin, London, United Kingdom         16:40 - 17:00       Group session – Design project suitable for funding application         ommittee: Y-ECCO         ce: Physicians, Paediatricians, Surgeons, IBD nurses	Welcome and introduction Nuha Yassin, London, United Kingdom         Session 1: Grant Writing – How to get started Isabelle Cleynen, Leuven, Belgium Dominik Bettenworth, Münster, Germany         16:10 - 16:25       Designing the ideal project Séverine Vermeire, Leuven, Belgium         16:25 - 16:40       Grant Writing – Tips and Tricks Gerhard Rogler, Zürich, Switzerland         Session 2: Group Session – Designing a successful project         Timothy Raine, Cambridge, United Kingdom Dominik Bettenworth, Münster, Germany Isabelle Cleynen, Leuven, Belgium Nuha Yassin, London, United Kingdom         16:40 - 17:00       Group session – Design project suitable for funding application         ommittee: Y-ECCO       E	Welcome and introduction Nuha Yassin, London, United Kingdom         Session 1: Grant Writing – How to get started Isabelle Cleynen, Leuven, Belgium Dominik Bettenworth, Münster, Germany         16:10 - 16:25       Designing the ideal project Séverine Vermeire, Leuven, Belgium         16:25 - 16:40       Grant Writing – Tips and Tricks Gerhard Rogler, Zürich, Switzerland         Session 2: Group Session – Designing a successful project Timothy Raine, Cambridge, United Kingdom Dominik Bettenworth, Münster, Germany Isabelle Cleynen, Leuven, Belgium Nuha Yassin, London, United Kingdom         16:40 - 17:00       Group session – Design project suitable for funding application         00       Bis:00	Welcome and introduction Nuha Yassin, London, United Kingdom17:00 - 17:35Session 1: Grant Writing – How to get started Isabelle Cleynen, Leuven, Belgium Dominik Bettenworth, Münster, Germany17:35 - 18:0017:35 - 18:0016:10 - 16:25Designing the ideal project Séverine Vermeire, Leuven, Belgium Grant Writing – Tips and Tricks Gerhard Rogler, Zürich, Switzerland17:35 - 18:00Session 3: Gr projects Nik John Dim16:25 - 16:40Grant Writing – Tips and Tricks Gerhard Rogler, Zürich, Switzerland17:35 - 18:00I1:35 - 17:50Session 2: Group Session – Designing a successful project Timothy Raine, Cambridge, United Kingdom Dominik Bettenworth, Münster, Germany Isabelle Cleynen, Leuven, Belgium Nuha Yassin, London, United Kingdom 16:40 - 17:00Group session – Design project suitable for funding application18:00Close - Y-ECCce: Physicians, Paediatricians, Surgeons, IBD nursesECCO Membership 2017 required Registration fee: EUR 80 (had

# 6<sup>th</sup> S-ECCO IBD Masterclass in collaboration with ESCP – Difficult decision making in IBD Thursday, February 16, 2017

	February 16				
08:00 - 08:10		, Tel Aviv, Israel	13:30 - 14:00	Debate 4: Ar patients	nal fissure and haemorrhoids in Crohn's
08:10 - 09:50	Session 1: Difficult decision making in Ulcerative Colitis Fabrizio Michelassi, New York, United States			13:30 - 13:40	Hands off Peter Irving, London, United Kingdom
	Yehuda Chov	wers, Haifa, Israel		13:40 - 13:50	<b>Take the knife</b> Paulo Kotze, Curitiba, Brazil
08:10 - 08:40	Debate 1: Tir	ning of surgery in Acute Severe Colitis		13:50 - 14:00	Discussion
	08:10 - 08:20	Save the colon – Medical treatments Silvio Danese, Milan, Italy	14:00 - 14:30	Debate 5: Th	e ileocolonic anastomosis
	08:20 - 08:30	Save the patient – Early surgery Antonino Spinelli, Milan, Italy		14:00 - 14:10	Side to side: Isoperistaltic vs. antiperistaltic Gianluca Sampietro, Milan, Italy
	08:30 - 08:40	Discussion			Hand-assist vs. single port
08:40 - 09:10	Debate 2: Sur	gery for Ulcerative Colitis revisited		14:10 - 14:20	Christianne Buskens, Amsterdam, The Netherlands
	08:40 - 08:50	lleorectal anastomosis is an attractive option		14:20 - 14:30	Discussion
		Ýves Panis, Clichy, France	14:30 - 15:00	Video conte	st: How do I do lap ileocolic resection? eoperative and postoperative management)
	08:50 - 09:00	Ileoanal pouch is universally best André D'Hoore, Leuven, Belgium		14:30 - 14:37	My technique
	09:00 - 09:10				Oded Zmora, Tel Aviv, Israel My technique
09:10 - 09:40	The techniqu			14:37 - 14:44	Ny technique Nuha Yassin, London, United Kingdom
	09:10 - 09:20	<b>Top down pouch surgery</b> Tom Øresland, Oslo, Norway		14:44 - 14:51	<b>My technique</b> Monica Millan, Barcelona, Spain
	09:20 - 09:30	<b>Bottom up pouch surgery</b> Janindra Warusavitarne, London, United		14:51 - 15:00	Discussion
		Kingdom	15:00 - 15:30	Coffee breal	κ
	09:30 - 09:40		15:30 - 15:55		u <mark>tting edge topics in IBD</mark> inelli, Milan, Italy
09:40 - 09:50	Video: TAMIS pouch	s total proctocolectomy and ileoanal	10.00-10.00		Curitiba, Brazil
09:50 - 10:20	Anders Tøttru Coffee break	up, Aarhus, Denmark <b>K</b>		15:30 - 15:40	Stem cells for surgeons Daan Hommes, Los Angeles, United States
10:20 - 12:00	André D'Hoc	<b>e art and science of IBD surgery</b> ore, Leuven, Belgium , Tel Aviv, Israel		15:40 - 15:55	IBD surgery: Last resort or another option Willem Bemelman, Amsterdam,
	10:20 - 11:20	3 Free papers 3 Trial updates		Socion 5: D	The Netherlands
	10:20 - 11:40	Imaging in the peri-operative management Yoram Bouhnik, Clichy, France	15:55 - 16:55	Consultants' Alessandro A	<b>corner:</b> vrmuzzi, Rome, Italy nelassi, New York, United States
	11:40 - 12:00	Surgical techniques to reduce recurrence rates in Crohn's Disease			se, Budapest, Hungary
	11.40 - 12.00	Fabrizio Michelassi, New York, United States		15:55 - 16:15	Perianal fistula Hagit Tulchinsky, Tel Aviv, Israel
12:00 - 13:00	Lunch break			16:15 - 16:35	Crohn's pancolitis Christoph Isbert, Würzburg, Germany
13:00 - 15:00	Yves Panis, C	fficult decision making in Crohn's Disease lichy, France rusavitarne, London, United Kingdom		16:35 - 16:55	Acute Colitis Giuseppe Sica, Rome, Italy
13:00 - 13:30		ost successful percutaneous drainage of	16:55 - 17:00	<b>Closing rem</b> Janindra War	<b>arks</b> usavitarne, London, United Kingdom
	13:00 - 13:10	Surgery for all Michel Adamina, Winterthur, Switzerland			
	13:10 - 13:20	<b>Still place for medical therapy</b> Shomron Ben-Horin, Ramat Gan, Israel			
	13:20 - 13:30	Discussion			
	<b>ce:</b> Surgeons, F	ECCO in collaboration with ESCP Physicians, IBD nurses ion		e: EUR 150 (ha	ired: Regular/Y-ECCO/IBD nurse Member If price for Y-ECCO and IBD nurse Members)

### 4<sup>th</sup> ECCO Ultrasound Workshop Thursday, February 16, 2017

	· ·				
07:30 - 07:45	Welcome an Speaker (tba)	d introduction	10:15 - 10:30	Coffee break	<
07:45 - 10:15	Speaker (tba) Speaker (tba)		10:30 - 12:00	Session 2 Speaker (tba)	
	07:45 - 08:00	Lecture: Ultrasound findings in Crohn's Disease and Ulcerative Colitis: Active vs.		10:30 - 10:45	Lecture: Role of bowel ultrasound in the postoperative Crohn's patient Speaker (tba)
		chronic Speaker (tba)		10:45 - 11:00	Lecture: The role of contrast enhanced ultrasound and sonoelastography in IBE
	08:00 - 09:00	Part 1: Interactive video presentation of ultrasound cases of IBD: Active or chronic? Speaker (tba)		11:00 - 12:00	Speaker (tba) Part 3: Interactive video presentation of ultrasound cases of IBD: Postoperative Crobac patient
	09:00 - 09:15	Lecture: Mural and extramural complications of Crohn's Disease			<b>Crohn's patient</b> Speaker (tba)
	09.00 - 09.15	Speaker (tba)	12:00 - 12:25	Post-course	test
		Part 2: Interactive video presentation of ultrasound cases of IBD: Mural and	12:25 - 12:30	Final remarks Speaker (tba)	
	09:15 - 10:15	extramural complications of Crohn's Disease Speaker (tba)			
Target audieno	esponsible Committee: EduCom in collaboration with ESGAR rget audience: Physicians with routine experience in intestinal ultrasound			ship	quired: Regular/Y-ECCO Membership o
ILIC) and narticip	aants of the int		Domistration for	• FUD 00 (bal	forico for VECCO Mombors

(IUS) and participants of the intestinal ultrasound training programme **Registration:** Online registration

Registration fee: EUR 80.- (half price for Y-ECCO Members) incl. 21% Spanish VAT

### 2<sup>nd</sup> ECCO-AOCC Forum: Learning from the Masters Thursday, February 16, 2017

· · · · · · · · · · · · · · · · · · ·					
09:00 - 10:15	<b>Session 1</b> Julián Panés, Barcelona, Spain Mamoru Watanabe, Tokyo, Japan		10:45 - 12:00	<b>Session 2</b> Marc Ferrante, Leuven, Belgium Jiaming Qian, Beijing, China	
	09:00 - 09:15	Welcome and introduction Julián Panés, Barcelona, Spain			Topic: Magnetic Resonance, Balloon Enteroscopy or Capsule endoscopy
	09:15 - 09:45	<b>Topic: Surveillance colonoscopy for CRC</b> <b>in IBD</b> Speaker (tba)		10:45 - 11:25	for Examination of the small bowel in Crohn's disease – Tandem Talk Jeong-Sik Byeon, Seoul, Korea Rami Eliakim, Tel Aviy, Israel
	09:45 - 10:15	Endoscopic and imaging features of CD, intestinal TB, and Bechet disease: Is instantaneous diagnosis feasible? Speaker (tba)		11:25 - 11:55	Topic: Endoscopic balloon dilatation for stricturing CD Martin Gôtz, Tübingen, Germany
10:15 - 10:45	Coffee brea			11:55 - 12:00	<b>Closing remarks</b> Suk-Kyun Yang, Seoul, Korea
Responsible Committee: ECCO Governing Board Target audience: Physicians, Surgeons				21% Spanish VAT nts is limited. Registration will be on a first	

come, first served basis.

**Registration:** Online registration

ECCO Membership 2017 required: Regular/Y-ECCO Member; AOCC Member

### 5<sup>th</sup> SciCom Workshop: Methodology on Research Thursday, February 16, 2017

09:00 - 10:20	<b>Session 1</b> Charlie Lees, Edinburgh, United Kingdom Florian Rieder, Cleveland, United States			10:45 - 12:00	<b>Session 2</b> Britta Siegmund, Berlin, Germany Gijs van den Brink, Amsterdam, The Netherlands	
	09:00 - 09:05	Welcome and introduction Charlie Lees, Edinburgh, United Kingdom			10:40 - 11:05	Transgenic animal models in IBD – Tricks you cannot read in the manuscripts
	00.05 00.00	How to set-up a successful translational IBD unit – Providing the frame for future			10110 11103	Arthur Kaser, Cambridge, United Kingdom
	09:05 - 09:30	<b>discovery</b> Britta Siegmund, Berlin, Germany		11:05 - 11:30	Recreating the intestine ex vivo – Where are we on the way to the artificial gut	
	09:30 - 09:55	Studying the microbiota – Bacteria and	11.05 11.50	Gijs van den Brink, Amsterdam, The Netherlands		
		Dirk Haller, Freising, Germany			11.00 11.55	Studying drug response – Translation of
	09:55 - 10:20	Omics analysis – Can we see the forest 5 - 10:20 for the trees		11:30 - 11:55	system biology to systems medicine Stefan Schreiber, Kiel, Germany	
	Jeff Barrett, Cambridge, United Kingdom		11:55 - 12:00	<b>Closing remarks</b> Florian Rieder, Cleveland, United States		
10:15 - 10:45	Coffee brea	K				FIOHALI MEGEL, CIEVEIALIO, UTILEO STATES

Responsible Committee: SciCom

Target audience: Basic scientists and interested clinicians Registration: Online registration

ECCO Membership 2017 required: Regular/Y-ECCO/IBD nurse Member Registration fee: EUR 80.- (half price for Y-ECCO and IBD nurse Members) - incl. 21% Spanish VAT

### 11<sup>th</sup> N-ECCO Network Meeting Thursday, February 16, 2017

Thursday, I	February 16	, 2017				
09:00 - 09:15	Welcome and introduction Karen Kemp, Manchester, United Kingdom		12:30 - 14:00	Lunch breal	< (Self Guided Poster Round)	
	Session 1: Quality care in IBD		12:45 - 13:45	Industry spo	onsored satellite symposium	
09:15 - 10:30	Kay Grevesor	n, London, United Kingdom o Barrero, Barcelona, Spain	14:00 - 14:45		Manchester, United Kingdom	
	09:15 - 09:45	Quality indicators in IBD Marian O'Connor, London, United Kingdom			Aarhus, Denmark Abstract 1	
		Remote monitoring in IBD		14:15 - 14:30	Abstract 2	
	09:45 - 10:00	Kathleen Sugrue, Cork, Ireland		14:30 - 14:45	Abstract 3	
	Patient Reported Outcomes (PRO'S) 10:00 - 10:30 Karen Kemp, Manchester, United		14:45 - 15:15	Coffee break		
	10.00 - 10.30	Kingdom		Session 4: M	: Medical management of IBD	
10:30 - 11:00	Coffee break		15:15 - 16:30	Kathleen Sugrue, Cork, Ireland Ester Navarro Correal, Barcelona, Spain		
11:00 - 12:30	Usha Chauha	<b>pecial considerations in IBD</b> In, Hamilton, Canada rciano Gonzalo, Barcelona, Spain		15:15 - 15:45	Therapeutic algorithms for IBD management Axel Dignass, Frankfurt, Germany	
	11:00 - 11:30	<b>Fertility and pregnancy in IBD</b> Janneke van der Woude, Rotterdam, The Netherlands		15:45 - 16:30	Mono vs. combo therapy Laurent Peyrin-Biroulet, Vandoeuvre-Lès- Nancy, France	
	11:30 - 12:00	Travel vaccines in IBD Kay Greveson, London, United Kingdom	46.20 47.00	Closing rem	Charlie Lees, Edinburgh, United Kingdom Closing remarks, N-ECCO in 2017 and beyond	
	12:00 - 12:30	Low Fodmap in IBD Kevin Whelan, London, United Kingdom	16:30 - 17:00		, Manchester, United Kingdom	
Target audien	Responsible Committee: N-ECCO Target audience: IBD nurses – advanced level Registration: Online registration				<b>iired:</b> IBD nurse Member . 21% Spanish VAT	

### 4<sup>th</sup> P-ECCO Educational Course 2017 – Clinical challenges in paediatric IBD Thursday, February 16, 2017

10:00 - 12:00	Salvatore Cu	<b>lenges in paediatric IBD</b> cchiara, Rome, Italy erusalem, Israel		10:50 - 11:15	Bone health in IBD – Does it concern paediatricians? Dan Turner, Jerusalem, Israel
	10:00 - 10:05	10:05 Welcome and introduction Dan Turner, Jerusalem, Israel		11:15 - 11:40	New treatments targeting the microbiota (FMT, antibiotics) –
		Treat to target in paediatric IBD – Same as in adults?	_	11.10	<b>A case-based discussion</b> Jarosław Kierkuś, Warsaw, Poland
	10:05 - 10:25	Patrick van Rheenen, Groningen, The Netherlands		11:40 - 12:00	Vedolizumab in children and other biologics after TNF – What can we extrapulate? A case-based discussion Hankje Escher, Rotterdam, The Netherlands
	10:25 -10:50	<b>Cannabis in adolescents – Any role?</b> Richard Russell, Glasgow, United Kingdom			
Responsible C	ommittee: P-l	ECCO	ECCO Membersh	ip 2017 require	d: Regular/Y-ECCO/IBD nurse/Affiliate Member
•		ans, Physicians, Surgeons, IBD nurses	Registration fee	<b>:</b> EUR 80 (ha	If price for Y-ECCO Affiliate and IBD nurse
Registration: (	Online registrat	tion	Members) - incl. 2	21% Spanish V/	AT

2 <sup>nd</sup> H-ECCO IBD Masterclass Thursday, February 16, 2017			Friday, Febr	Friday, February 17, 2017		
13:30 - 13:35	Welcome and introduction Roger Feakins, London, United Kingdom		08:00 - 09:40	Session 3: Recent advances Vincenzo Villanacci, Brescia, Italy Fernando Magro, Porto, Portugal		
13:35 - 15:00		k, Paris, France ay, London, United Kingdom		08:00 - 08:25	Pathologists' role in clinical trials (Tandem talk) Fernando Magro, Porto, Portugal	
	13:35 - 13:50	What clinicians expect from pathologists James Lindsay, London, United Kingdom		00.25 00.55	Paula Borralho, Lisbon, Portugal Hot topics	
	13:50 - 14:00	What pathologists expect from clinicians Paula Borralho, Lisbon, Portugal		08:25 - 08:55	Magali Svrcek, Paris, France Pathogenesis of fibrosis and fistulas in IBD	
	14:00 - 14:15	<b>IBD – endoscopic features</b> Gionata Fiorino, Rozzano, Italy		00.33 05.10	Roger Feakins, London, United Kingdom Prognostic factors for IBD – clinical and	
	14:15 - 14:35	IBD in biopsies Roger Feakins, London, United Kingdom		09:10 - 09:40	pathological (Tandem talk) Paula Borralho, Lisbon, Portugal Laurent Peyrin-Biroulet, Vandoeuvre-lès-	
	14:35 - 15:00	Pathology of IBD in resections Cord Langner, Graz, Austria	09:40 - 10:10	Coffee brea	Nancy, France	
15:00 - 15:30 15:30 - 17:10	Coffee breal Session 2: M Paula Borralh		10:10 - 11:35	Roger Feakir	<b>utside the colon</b> 1s, London, United Kingdom London, United Kingdom	
	15:30 - 15:45	Clinical mimics of IBD Laurent Beaugerie, Paris, France		10:10 - 10:30	Extraintestinal manifestations of IBD Peter Irving, London, United Kingdom	
	15:45 - 16:05	Relationships between diverticulosis and IBD		10:30 - 10:50	<b>Upper GI pathology in IBD</b> Paula Borralho, Lisbon, Portugal	
	16:05 - 16:25	Vincenzo Villanacci, Brescia, Italy Microscopic colitis and IBD		10:50 - 11:10	Ileal pathology in IBD and related disease Vincenzo Villanacci, Brescia, Italy	
	16:25 - 16:45	Cord Langner, Graz, Austria Infection, drugs, and other pathological mimics of IBD		11:10 - 11:25	Pathology of IBD in primary sclerosing cholangitis Magali Svrcek, Paris, France	
	16.45 17.10	Roger Feakins, London, United Kingdom Dysplasia (with interactive cases)		11:25 - 11:35	The ideal IBD pathology report Roger Feakins, London, United Kingdom	
	16:45 - 17:10	Magali Svrcek, Paris, France Vincenzo Villanacci, Brescia, Italy	11:35 - 11:40	Closing rem Roger Feakin	<b>arks</b> ıs, London, United Kingdom	
Target audiend	Responsible Committee: H-ECCO Working Group Target audience: Histopathologists; Clinicians Registration: Online registration			e: EUR 80 (hal	<b>d:</b> Regular/Y-ECCO/IBD nurse/Affiliate Member f price for Y-ECCO Affiliate and IBD nurse AT	

### 2<sup>nd</sup> D-ECCO Workshop Friday, February 17, 2017

08:30 - 08:35	Session 1: Nutritional science in IBD			10:20 - 10:40	Enteral nutrition in adults: Has anything changed?
08:35 - 09:40				10:40 - 11:00	Walter Reinisch, Vienna, Austria Case presentation – Dietitian's view Nicolette Wierdsma, Amsterdam,
	08:35 - 08:55	<b>Biology of SCFA</b> Konstantinos Gerasimidis, Glasgow, United Kingdom		11:00 - 11:05	The Netherlands Q&A
		Food additives – A role in IBD?	11:05 - 11:20	Coffee breal	k
	08:55 - 09:15	Arie Levine, Tel Aviv, Israel		Session 3: Di	iet and nutritional aspects of IBD
	09:15 - 09:35	<b>Diet and microbiome</b> Jeroen Raes, Ghent, Belgium	11:20 - 12:25		ner, London, United Kingdom s Gerasimidis, Glasgow, United Kingdom
	09:35 - 09:40			11:20 - 11:40	<b>Calcium, Vitamin K2 and Vitamin D</b> Tibor Hlavaty, Bratislava, Slovakia
09:40 - 10:00	Coffee breal	κ			Role of dietitian in IBD teams across
10:00 - 11:05		<b>Session 2: Clinical nutrition in IBD</b> Kevin Whelan, London, United Kingdom		11:40 - 12:00	Europe Rotem Sigall-Boneh, Tel Aviv, Israel
	Arie Levine, T 10:00 - 10:20			12:00 - 12:20	Food related quality of life and the patient's perspective on diet Kevin Whelan, London, United Kingdom
	10.00 10.20	Miranda Lomer, London, United Kingdom		12:20 - 12:25	Q&A
			12:25 - 12:30	Closing rem Konstantino	<b>arks</b> s Gerasimidis, Glasgow, United Kingdom
Responsible C Farget audien Registration: (	<b>ce:</b> Dietitians, l				d: Regular/Y-ECCO/IBD nurse/Affiliate Member cl. 21% Spanish VAT

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# Interview with the new President-Elect



Silvio Danese © Silvio Danese

### Silvio Danese

Position:	President-Elect, 2016–2018
Nationality:	Italian
Date of birth:	January 10, 1975
Past ECCO Positions	SciCom Member,
	Secretary

# What was your motivation for taking up the position of President-Elect?

Right from the beginning of my GI fellowship I was very intrigued by IBD, and I soon started to commit to this field with the intention of helping patients struggling with these diseases. In my professional life there are three major activities: Clinical practice, research and education (as an academic gastroenterologist). This fits exactly with the three major goals of ECCO. This is why, for me, applying for the ECCO Presidency means once again making a large commitment to the field of IBD.

# As a future ECCO President, what are your visions for ECCO in the short and long term and what are the next steps in the evolution of ECCO?

It is true that in recent years ECCO has evolved a lot. It was initially a sort of club of friends, and I still remember when I started as a Y-ECCO Member, with new friends like Séverine, Simon, Daan and many others. That spirit of friendship has been contagious. As a result, ECCO has become THE global IBD society and now has 3,215 members. One of my main goals during my terms of office is to foster further enlargement of ECCO on a global scale, with greater involvement of our colleagues in the United States and Asia through more proactive interactions with various ECCO Initiatives. Though already in place, these mechanisms should be boosted much more, and this would tremendously advance the field of IBD, particularly in terms of research activities. As a member of IOIBD and having done most of my training in the United States, I firmly believe in international cooperation, as collaboration rather than competition is obviously the key to progress within the IBD field, the promotion of which is our ultimate duty as ECCO Members.

### What concrete actions/activities will you carry out in order to achieve the main objectives during your terms of office?

I truly believe that an even greater focus on young members should be considered, as Y-ECCO is the future of ECCO. Active involvement of Y-ECCO is already a reality, but I am convinced that young members should be even more actively involved in all the scientific activities. After all, investing in Y-ECCO is truly investing in the future of ECCO and I will make more efforts to further integrate Y-ECCO into the ECCO Activities. During my terms of office I will try to engage Y-ECCO Members in clinical practice, research and education:

### **Clinical practice**

The aim is to develop pathways of care across Europe despite the economic challenges. Guidelines have been and will remain instrumental. But we need to move to the next level, developing both tools able to measure quality of care and pathways of care that overcome the fragmentation that may occur from GP to tertiary referral centre. To achieve these goals, more cooperation between EFCCA and ECCO is needed.

### Research

This is a strategic topic. ECCO until now has facilitated "non-sponsored, academic" clinical studies and supported research with grants. However, it is obvious that much more can be done with respect to fund raising campaigns, lobbying at EU level, working more closely with patients and donors as well as with our sister societies. In addition, our long-term relations with EMA foster exchange of knowhow and aim at faster development processes of new drugs. We need more awareness for IBD and this would lead to more resources.

### Education

Here I would make two simple points: (1) we must ensure the spreading of educational activities in multiple countries, particularly where fewer resources are available; (2)

e-Learning needs to be further implemented and educational activities should be expanded to all those areas that are of strategic significance for multidisciplinarity of IBD care.

# What are the things that, in your opinion, make ECCO special/distinguish it from other associations?

Even though we now have well over 3,200 members, the element of friendship is still contagious and the ECCO Spirit of collaboration in the IBD field is very strong among all members.

# Who was the person who most influenced you in your career, and how?

Claudio Fiocchi, my mentor. He taught me that while young fellows need guidance, they must also be allowed the freedom to make mistakes and to learn from those mistakes in order to grow stronger!

In the lab it happens all the time. During my postdoc I had a lot of fun because Claudio felt able to allow me to take forward my enthusiasm for research without limitations.

# What, in your eyes, have been your biggest professional achievements thus far?

Recognising the vascular involvement in intestinal inflammation, and how its blockade in multiple ways can be beneficial for IBD therapy. Today we have the first drug that blocks leukocyte-endothelial interactions and many others will arrive soon.

## What are your strongest points, and what is your weakest?

I am stubborn and emotional. Those qualities can be strengths and weaknesses, depending on the timing!

# If you had not become a doctor, what might you have been doing today instead?

I was in love with literature, so perhaps that is the reason I keep writing... albeit in medicine.

# What do you do for recreation and fun and what is your favourite dish?

I love eating...that is why I also run, trying to burn off a few calories. I try to spend as much time as I can with my three little kids (two twin girls, who are 6 years old, and one boy, aged 5 years) and am always running behind them! My favourite dish is pizza....what else, being Italian?!

> PIETER HINDRYCKX ECCO News Associate Editor

# I-CARE Study - Ibd CAncer and seRious infections in Europe

Congratulations from the I-CARE team! Over 1,700 patients have been included in 14 countries in I-CARE!



As you know, the I-CARE European prospective study, endorsed by ECCO, will evaluate for the first time the effectiveness and safety of biologics (anti-TNF and vedolizumab) alone or in combination therapy in a large cohort of IBD patients. After CESAME with thiopurines, I-CARE will provide unique information on biologics currently used in our clinical practice.

It takes only 15 minutes to include a patient in the I-CARE cohort study and then you will

I-CARE Map © I-CARE only need to check the data entered by your patient on a yearly basis – that's all!

We have very good news: 84% of the patients have confirmed their participation! However, the number of active investigators in I-CARE is still too low (128, which equates to just 21% of registered investigators)! As a reminder, we need 800 investigators to reach our objective (22 patients per investigator for a final number of 17,600 patients)



### WE ARE COUNTING ON YOU!

Please contact your National Coordinator or the I-CARE team (*icare@getaid.org*) if you wish to be part of this great adventure or have any questions.

### All the best,



L. Peyrin-Biroulet (President of I-CARE), L. Beaugerie (Head of SciCom & Head of Cancer Committee), F. Baert (General Secretary), J.F. Rahier (Head of Infection Committee) M.J. Bertin (Project Director), P. Rabiéga (Project Manager), L. Jerber (French Study Coordinator)

# ECCO Fellowship Study Synopses

Oral nutritional supplement therapy improves bone structure and body composition in Paediatric Crohn's Disease: a controlled trial

### Aim of research

Growth failure is a frequent and serious complication of Paediatric Inflammatory Bowel Disease (PIBD). Muscle mass, bone density and geometry are adversely affected in PIBD. Growth does not normalise during therapy (with either exclusive enteral nutrition or biologicals). In a previous study we showed that 8 weeks of exclusive enteral nutrition therapy improves muscle mass and bone health towards normal within 3 months, but that it deteriorates after discontinuation of the treatment. Few data are available on the effects of an oral nutritional supplement (ONS) in children with Crohn's Disease (CD) on the different outcome parameters, and no study has been published with respect to bone quality. We hypothesise that a supplementary enteral nutrition providing 25% of daily energy and nutrient needs, in addition to maintenance drug treatment, will improve growth, muscle mass and bone health,

compared to a control group without ONS. With this controlled trial we aim to investigate the short- and long-term efficacy of ONS on height development, muscle mass, bone metabolism, bone structure, body composition, metabolomics, intestinal microbiota and clinical course of Paediatric CD (PCD). If the benefits of ONS can be proven, it could potentially become a long-term treatment option for PCD patients, justifying the costs.

### Methodology and proposed timing

Participants have been recruited since April 2016 and they return for follow-up visits at 3-month intervals up to one year. Each visit includes anthropometric measurement, physical examination with grip strength assessment, questionnaire administration, blood collection for inflammatory markers, targeted metabolomic profile and bone parameters, and stool and urine sample collection. Peripheral quantitative computed tomography is used at the nondominant forearm to evaluate bone and muscle parameters. Body composition is calculated precisely by air displacement plethysmography. The metabolomic and microbiota profile will be assessed and analysed after study completion (planned for August 2017).



Klara Frivolt © G Frivolt

KLARA FRIVOLT ECCO-Nestlé Health Science Nutrition Fellow 2016

### The interleukin-1 pathway as a putative new target in pediatric inflammatory bowel disease

Inflammatory Bowel Disease (IBD) is driven by uncontrolled T-cell responses. The disease cannot be cured and, despite the current treatments, 40%–50% of patients suffer from frequent relapses or continuous inflammation. Currently, there is a great need to identify biomarkers for classification of IBD patients that will predict therapy responsiveness. Given the fact that the current immunosuppressive strategy fails in a subgroup of patients, we need to identify biomarkers that differentiate responders from non-responders and to define novel targets for immunosuppressive therapy. Our preliminary studies revealed that the production of the pro-inflammatory cytokine IL-1 $\beta$  by antigen-presenting cells (APC) is enhanced in a subgroup of IBD patients. Based on these findings, we hypothesise that IL-1ß drives inflammation in a subgroup of IBD patients and may be a novel target for more

effective treatment. The aim of this project is to determine the role of IL-1 $\beta$  in promoting inflammatory T-cell responses and intestinal inflammation in paediatric IBD patients.

Together with Warren Leonard (Chief, Laboratory of Molecular Immunology and Director, Immunology Center, National Institutes of Health, Bethesda, USA), who is one of the leading investigators in the molecular regulation of cytokine signalling, we will identify the molecular signature of APC from paediatric IBD patients with dominant IL-1 $\beta$  expression using high-throughput RNA sequencing. This will yield novel biomarkers that will subsequently be used, in the second half of the project, to assess the effectiveness of existing IL-1 blocking agents and new therapeutics in inhibiting APC-driven T cell-mediated immunity using an in vitro assay. Besides delivering

major advances in understanding the T celldriven immunopathology in IBD, knowing the mechanism by which IL-1 stimulates inflammatory T cells will yield parameters to classify IBD patients. This knowledge on the IL-1 pathway will allow us to identify patients who will benefit from anti-IL-1 therapy.



Sharon Veenbergen © Veenbergen

SHARON VEENBERGEN ECCO-Nestlé Health Science Nutrition Fellow 2016

### Post-genomic application in general, and the establishment of computational approaches relevant to traslational medicine

### Aim of the research project

There is a great need to improve IBD diagnosis and treatment, and finally establish novel criteria for tailor-made therapy in individual patients. Genetic discoveries have had a remarkable impact on our understanding of IBD pathogenesis, though their translation to boost the development of novel therapeutic strategies has been limited. A promising way to exploit this knowledge is the identification of alternative indications for already approved drugs (drug repositioning). It has been shown that predisposing variants at IBD risk loci primarily exert their functional effects through alteration of gene expression (eQTLs). Therapeutic effects may be obtained through attempts to restore physiological expression of risk IBD genes in directions opposite to those induced by the risk variants. The aims of this research are to develop an in silico pipeline using computational methods for drug repositioning

in IBD and to establish personalised treatment predictors integrating GWAS/eQTL data and drug-induced gene expression signatures from the LINCS database.

### Methodology

eQTL IBD risk variants will be compiled into a catalogue of genotype-driven risk expression signatures. Based on the available genotype data, we will then assign specific signatures to individual patients (n=200) for tailored computational drug screening. Patients' signatures will be matched against the LINCS drug-induced gene expression database (*www. lincsproject.org*) in order to obtain, for each patient, a list of top-scoring drugs predicted to counteract specific risk eQTL profiles. These anti-correlated drugs represent putative new therapeutic candidates (repositioning). As a downstream proof of principle, we aim to perform correlation analyses connecting predicted personalised treatment profiles (individually repurposed top-scoring drugs) to relevant patients' clinical phenotypes (location, activity, response to therapy and need for surgery). The aim is to verify whether IBD patients cluster into discernable groups according to their computationally predicted treatment modalities.



Ferdinando Bonfiglio © Bonfiglio

FERDINANDO BONFIGLIO ECCO Fellow 2016

### Hypoxia, autophagy and inflammasome

Inflammatory Bowel Disease (IBD) is a chronic intestinal pathology caused by various complex interactions between the environment, genetics and gut microbiota. Despite all the research that has been conducted to date, its aetiology is still not well understood. In recent years, a huge number of genetic studies have strongly pointed to a defective autophagy in these patients due to the presence of several genetic alterations in autophagy-related genes, such as ATG16L1, NOD2 and IRGM. Furthermore, it has recently been reported that this process is defective in these patients. On the other hand, another important feature of this pathology is the presence of hypoxia, which is increased in chronic inflammation, inducing both autophagy and inflammasome activation.

Despite recent extensive efforts, the molecular mechanisms underlying the crosstalk of these processes remain poorly understood. Mounting evidence suggests that the imbalance in the mutual regulation of autophagy and NLRP3 inflammasome activation under hypoxia plays a role in the development of IBD, and that signalling molecules involved in this balance could represent novel therapeutic targets for the treatment of IBD.

In this project, we will study the effects of hypoxia in DSS-treated wild-type, NIrp3-/-, IL10-/- and double-knockout IL10-/-NIrp3-/mice. Additionally, we will culture lipopolysaccharide-treated human intestinal epithelial cells (HT29 and Caco-2 cells) and monocytes (THP1 cells) under normoxia or hypoxia in order to characterise the molecular mechanisms of hypoxia-associated autophagy and inflammasome activation under conditions of intestinal inflammation. Clarification of the molecular mechanisms governing the mutual regulation of autophagy and inflammasome in the hypoxic environment of the inflamed mucosa will improve knowledge of the pathogenesis of IBD and may provide novel therapeutic targets for its treatment.

# The ClinCom School for Clinical Trialists

Following the success of the 1<sup>st</sup> School for Clinical Trialists at the 11<sup>th</sup> Congress of ECCO, held in Amsterdam in March 2016, ECCO will run a further workshop in 2017

his session at the annual ECCO Congress aims to help delegates understand the different types of clinical trial, to share ideas on how to optimise recruitment and examples of best practice and to consider what the future holds for IBD clinical trials. Delegates include clinicians, research fellows, trial coordinators, IBD nurses and research nurses.

The initial talk on different trial designs and terminology will review the difference between efficacy and effectiveness and address how the statistical question and the design of the trial dictate the size of the clinical trial required. Looking to the future, new types of trial design may be useful, with cluster randomisation and more adapted designs.

All trials eventually sink or swim by the success of their recruitment. The workshop will include helpful tips on how to optimise recruitment, such as developing a research culture amongst the whole team, including patients, and only taking on trials that match the patient population available. Recent surveys have confirmed that IBD patients are enthusiastic supporters of research. Delegates thinking of setting up national or international trials would be well advised to look at the clinical trials toolkit website *www.ct-toolkit.ac.uk/ routemap*. This is a comprehensive webtool to assist with trial set up.



1<sup>st</sup> School for Clinical Trialists, ECCO Congress, Amsterdam 2016© ECCO

Members of the hugely successful research team from Leuven will share some of the practical tips and tricks that help trials start up smoothly and minimise screen failures whilst boosting participation. In summary, being well prepared at each step of the process and having the time and energy to see the trials through to a successful conclusion are the key messages.

In future, IBD clinical trials will move towards having more co-primary end points, central reading of endoscopy, histology and MRI scans, and tight control of disease and damage assessment. Drug trials are increasingly being designed as head-to-head studies and incorporating biomarker and therapeutic drug monitoring.

ClinCom is proud to present the ECCO 2017 School for Clinical Trialists to equip delegates with valuable information, ideas and skills to help deliver clinical trials in IBD.

The programme of the 2<sup>nd</sup> School for Clinical Trialists is available in the Congress section of this issue of ECCO News, on page 8.

### JOHN MANSFIELD ClinCom Member

# The 45<sup>th</sup> ECCO Educational Workshop

### A report from the 45<sup>th</sup> ECCO Educational Workshop in Siófok, Hungary – June 3, 2016

he workshop was held in the popular holiday destination of Siófok on the southern coast of Lake Balaton in Hungary, where there are several excellent hotels. The 45<sup>th</sup> ECCO Educational Workshop was organised by the ECCO National Representative for Hungary, Tamas Molnar, from the University of Szeged, in collaboration with the chief of the local organising committee, Ferenc Szalay.

More than 80 delegates, mostly from Hungary, attended the workshop, including both young, talented future gastroenterologists and more experienced colleagues. **Abraham Eliakim** and **Janindra Warusavitarne** were the expert faculty from ECCO. Thanks to the representation of different professions – gastroenterology and surgery dedicated to IBD – all points of view concerning the selected hot topics were discussed.

After **Tamas Molnar**, head of the local organising committee and host for the workshop, had welcomed the delegates, Janindra Warusavitarne from London, on behalf

of ECCO, gave a presentation on ECCO's history and the organisation of the ECCO Committees, ECCO Guidelines and ECCO Consensuses. He also described the success of ECCO EduCom over the past decade in arranging workshops throughout Europe and also intercontinentally in order to foster the implementation of ECCO Guidelines by gastroenterologists worldwide.

# During the workshop day a total of seven cases were presented and discussed, as follows:

**Case 1:** Optimising therapy – Abraham Eliakim, Tel Aviv, Israel

**Case 2:** Acute Severe Colitis – Zoltan Szepes, Szeged, Hungary

**Case 3:** Pouchitis – Janindra Warusavitarne, London, UK

**Case 4:** How to deal with side effects – Pal Miheller, Budapest, Hungary

**Case 5:** IBD and cancer/comorbidity – Janindra Warusavitarne, London, UK

**Case 6:** Stopping drugs, exit strategy – Klaudia Farkas, Szeged, Hungary

**Case 7:** Pregnancy and IBD – Tamas Molnar, Szeged, Hungary

The atmosphere was very interactive between the faculty and the delegates during the workshop, with many discussions, critical comments and constructive answers.

Finally, Abraham Eliakim gave an excellent State of the Art Lecture about the importance of mucosal healing, before Tamas Molnar closed the meeting. After a short break all the ECCO Workshop delegates moved on to the National Gastroenterological Congress, held at the same venue at the Hotel Azúr.

We hope to be able to welcome you all back to Siófok in the future – at any time, and whether for business or pleasure.

Our thanks are due to the secretaries, **Phillip** and **Flavia**, and ECCO EduCom for their effective organisation before and during the workshop!

TAMÁS MOLNÁR ECCO National Representative, Hungary

# 15<sup>th</sup> IBD Intensive Advanced Course

The next IBD Intensive Advanced Course will take place over 1.5 days on Wednesday, February 15, 2017 and Thursday, February 16, 2017 before the start of the main ECCO Congress

his highly popular course is now in its 15<sup>th</sup> year and, based on the success of previous courses, will follow a similar format covering the core curriculum by means of a variety of teaching methods including lectures, interactive case discussions and seminars.

Active participation of attendees in the discussions is integral to the success of the course and this aspect is facilitated by the relaxed and friendly atmosphere in which attendees from Europe and the rest of the world are encouraged to interact.

The faculty is carefully chosen not just for their expertise in the areas on which they are invited to speak, but also for their ability as educators. The course covers a wide curriculum including cutting-edge science as well as advanced clinical practice and also allows participants to choose areas of particular interest on which to focus.

Whilst the course has always received positive feedback, the members of the Education Committee of ECCO pay keen attention to suggestions for improvement and have therefore included the following amendments to the course:

- A new concept of deep review of IBD drug management focussing on mechanisms of action, optimisation (personalised medicine) and practical aspects of the use of the medication.
- Seminar sessions in smaller groups covering topics including: management of extra-intestinal manifestations and pregnancy

in IBD, complications associated with anti-TNF use and endoscopy in IBD.

• A greater emphasis on **case-based discussions** and **interactive sessions** rather than didactic lectures, with several interesting new cases (e.g. refractory pouchitis).

We are looking forward to seeing young keen gastroenterologists at the 15<sup>th</sup> IBD Intensive Advanced Course in Barcelona in 2017!

> PASCAL JUILLERAT AND PETER IRVING EduCom Members and ECCO IBD Intensive Advanced Course Directors

# ECCO Ultrasound and Endoscopy Workshops at ECCO 2017:

Advancements and Use in Clinical Practice

iagnostic imaging and endoscopy represent the main procedures for the detection and appropriate management of patients with IBD. In expert hands, their usage ranges from early diagnosis and detection of complications to the prediction and assessment of therapeutic response. Intestinal ultrasound has taken hold in a number of countries, where it is performed by gastroenterologists as a practical and patient-friendly diagnostic tool that allows rapid clinical decision making, improved management of patients with suspected IBD, effective monitoring of disease activity and assessment of complications. Endoscopy has resulted in great progress in diagnosis and therapy and, in the hands of gastroenterologists, continues to occupy a central position in the clinical management of IBD patients. In February 2017, ECCO will offer two Educational Workshops devoted to the features and advancements of bowel ultrasonography and of diagnostic and therapeutic endoscopy, placing their use within the clinical context as reflected by real cases.

In spite of the great success and high acceptance among gastroenterologists that have been achieved by the ECCO Workshop on bowel ultrasonography since it was first held in Copenhagen in 2014, we felt that it was time for a change. We shall be moving on from the practical hands-on experience of the previous editions, where we introduced bowel ultrasound

to IBD specialists with little or no experience in the technique. Now the focus will be on the role of bowel ultrasound in the management of IBD patients in real life, highlighting its performance and recent advancements. On this occasion, after introductory lectures there will be interactive video case presentations on the role of ultrasound in patients with active and guiescent Crohn's Disease and Ulcerative Colitis, mural and extramural complications of IBD and postoperative Crohn's Disease as well as on the advancements in ultrasound as applied to IBD, such as contrast enhancement and sonoelastography. This workshop is targeted at physicians with routine experience in intestinal ultrasound but also at IBD specialists interested in the potential of this fascinating technique. With respect to the development of education on ultrasound, ECCO would particularly like to acknowledge the valued cooperation with the European Society of Gastrointestinal and Abdominal Radiology (ESGAR) and the European Federation of Societies of Ultrasound in Medicine and Biology (EFSUMB).

Next year ECCO will once again offer the ECCO Workshop on endoscopy for IBD specialists, aimed primarily at those who already have some experience in the technique. Given the great success of the previous edition, the same formula will be adopted: four sessions covering endoscopic activity, surveillance, small bowel endoscopy and operative endoscopy. After an introductory talk by experts in each field, real cases on workstations will be discussed, translating the updated theoretical knowledge into real practice. At the end of the course, participants should: be able to write a report; be able to discriminate and describe the endoscopic activity in IBD and to use endoscopic indexes; know the salient technical aspects of and the indications for chromoendoscopy; understand the indications for videocapsule endoscopy and double balloon enteroscopy when assessing the small bowel and be able to interpret the acquired images; and know the indications, relevant technical aspects and potential complications of operative endoscopy in IBD when used for such procedures as balloon dilatation, polypectomy, mucosectomy, and management of dysplastic lesions.

I hope that you will find these activities interesting and constructive and that they will assist you in your work and your career. Please register in time to experience these interactive workshops and share your knowledge and expertise with international experts. We are looking forward to seeing you in Barcelona 2017.

GIOVANNI MACONI EduCom Member



Volume 10 Issue 9 September, 2016

# **JOURNAL of CROHN'S and COLITIS**



Variability in endoscopic scoring of Crohn's disease. See editorial page 999 and articles pages 1001 and 1006

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www.ecco-ibd.eu

# ECCO Survey – Registries across Europe

### Epidemiological research potential and beyond

### Introduction

At ECCO we pride ourselves working on collaboratively. ECCO has 36 Member Countries; not only does this equip us with an array of IBD-ologists, but also a vast combined patient population. The EpiCom mission is to conduct and support epidemiological research across Europe. In particular we aim to evaluate differences in care across Europe, and subsequent impact on patient outcome.

In order to achieve these aims, we first need to know what information is available and how to access it. With this in mind, the EpiCom survey was undertaken. This article outlines the key findings; a complete analysis will be submitted to the Journal of Crohn's and Colitis.

### The EpiCom Survey

National Representatives from each of the 36 ECCO Member Countries were asked to complete a survey requesting information about registries within their home country. In addition, DG SANTE National Contact Points for these 36 countries and also for Iceland and Luxembourg were contacted with the same survey.

The survey enquired about population registers, birth and death registers, disability benefits registers, hospital registers, adverse events and prescription registers, surgery registers, cancer registers, twin and multiplex registers and IBD patient registers and biobanks. When possible, National Representatives were asked to provide a contact for each register and to comment on whether the register is a public or private facility.

### Who responded?

National Representatives and/or Contact Points from 33 countries responded at least in part. When responses were incomplete or absent, EpiCom members searched for missing information online. Figure 1 shows the countries from which information was acquired.



# Hospitalisation databases and patient registers

Twenty-nine countries surveyed have a hospitalisation database, of which 22 are thought to be public. Fewer countries have patient registries containing detailed information about treatment and diagnosis. Fourteen reported national patient registries and two, local registries. It is likely that the remainder have at least some form of local hospital record or discharge information. Twenty-one countries reported some form of adverse event register.

The majority only required mandatory reporting of events from trials as opposed to clinical practice. Fifteen countries have a register of specific medications, such as biologics; however, access to and completeness of these registries are variable.

Figure 1: Participating Countries © ECCO Participating Countries

### **National statistics**

Population registers, including birth and death registers, are kept by the majority of countries. Twenty-one countries have a disability pensions register, although these do not store details of disease or disability.

### **IBD** registries

Seventeen surveyed countries reported an active IBD registry (Figure 2). In many cases joining is voluntary, and the majority of IBD sufferers are not included. The information included by each country often contrasts. That said, local or "opt in" registers can be valuable epidemiological research tools; the EpiCom inception cohort included data from 31 countries.

### Twin and multiplex registries

Twins and multiplex families offer invaluable insights into the relative importance of genetic



and environmental factors. Sixteen of the surveyed countries have some form of twin research cohort. The Scandinavian countries have comprehensive registries of all twin births, linkable with medical history. Applications can be made to search the database to invite twins to participate in research.

However, the majority of European twin cohorts are made up only of twins who have responded to recruitment advertisements for a specific purpose. They do not contain population data or links to health records.

### Biobanking

Fourteen countries have biobanks either actively running or in development. The International IBD Genetics Consortium have undertaken meta-analysis of genotyped IBD patients from across the world. Within the UK, the IBD Bioresource has been launched as a subset of the established UK NIHR Bioresource. The UK IBD Twin Registry is currently creating an IBD twin biobank, and the Danish Twin Registry has already stored DNA, RNA, plasma, serum, blood and stool from all monozygotic IBD twins willing to participate.

# Disease progression – surgery and cancer registries

Only 13 countries reported a register of surgical procedures. Several of these were local datasets, and it is as yet unknown whether such registers contain details of underlying medical conditions or are linkable with medical records.

Perhaps the most complete registers across Europe are the cancer registers. All 38 countries surveyed have a cancer register. The majority mandate entry of all cancer diagnoses.

# How useful are these registries?

The diversity and at times incompleteness of the registries across Europe can seem overwhelming when considering research potential. National data collection varies considerably, as do legal and cultural attitudes towards confidentiality, consent and central records.

The immediate challenge for EpiCom is to form links with as many of these registries as possible, and to ascertain exactly what data is stored and how it can be accessed.

### The future

IBD databases, registries and biobanks are growing across Europe. As these resources develop, ECCO wishes to facilitate early collaboration. The success of the International IBD Genetics Consortium is testament to the importance of pooling expertise, resources and samples.

The cancer registries provide further inspiration; they give hope that, with the correct infrastructure, the aims of EpiCom are achievable throughout Europe. In addition to epidemiological research, a close network of IBD registries and biobanks will also facilitate both translational science and trial recruitment. All are required for us to best serve our current and future IBD patients.

### Acknowledgements

Many thanks to ECCO National Representatives and DG SANTE Contact Points from the following countries that completed the survey: Austria, Belgium, Bosnia and Herzegovina, Bulgaria, Croatia, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Israel, Ireland, Italy, Latvia, Luxemburg, Malta, Moldova, The Netherlands, Norway, Poland, Portugal, Romania, Serbia, Slovenia, Spain, Sweden, Switzerland, Turkey, United Kingdom.

> EBBE LANGHOLZ EpiCom Chair HANNAH GORDON External Support

# ECCO e-Guide – New Enhancements

he ECCO e-Guide has been conceived and developed to help clinicians of any level of expertise in IBD when approaching IBD patients in their daily practice. It is intended to be both easy to use and complete, summarising all the useful information contained within the ECCO Guidelines. In the last few years, a very motivated ECCO Taskforce, led by Marcus Harbord, has done a huge amount of hard work to set up algorithms, to include useful information about IBD and all aspects of its management and to incorporate educationally supportive material into the e-Guide on the ECCO Website. During recent months, the main challenge has been to harmonise and update all the contents in line with the newly published ECCO Guidelines, and to select useful supporting educational tools for each section. This task has now almost been completed. In early July, during the e-Guide Taskforce meeting, all those members who are deeply involved in this ambitious project discussed and agreed on the final run to complete it. In particular, three aspects have been finalised.

First, all the contents of the e-Guide will be checked, updated and amended, if needed, prior to the next UEGW. Then, a new structure will be finalised and ready for release just prior to UEGW, and then again before ECCO '17. Probably, in subsequent years, UEGW and the ECCO Congress will continue to serve as two time points for regular updating of the e-Guide content. The new structure will include new topics recently covered by the ECCO Guidelines, such as pregnancy and lactation, extraintestinal manifestations, malignancies and the management of paediatric IBD. Second, Marcus Harbord is working very hard especially on the paediatric section, developing the diagnostic and therapeutic algorithms that need to be



ECCO e-Guide Taskforce © ECCO

included, drafting supporting material and background text, and also splitting the entire section into two more detailed sections about Paediatric Crohn's Disease and Paediatric Ulcerative Colitis. In this context, new Guidelines content, such as the Nurse Consensus or new drugs that have recently been introduced into clinical practice, have been included in the project. Third, the Taskforce decided to choose two "e-Guide champions", one from GuiCom and one from EduCom, who are required to know the e-Guide extremely well and will accordingly be able to make content updates easily when new Guideline updates are released. In fact, the publication of updates of the ECCO Guidelines on Crohn's Disease and Ulcerative Colitis will require appropriate harmonisation and updating of the e-Guide in time for each of its biannual releases

The e-Guide project will prove a great success thanks to the efforts of all members of the Taskforce, and specifically Marcus and Neil,

who are working very hard with the support of all of us. Our hope is to provide clinicians with a useful tool that allows standardisation of IBD management at any level, in accordance with the evidence-based ECCO Guidelines. We invite all ECCO Members to widely use and test the e-Guide content in the coming months and to provide us with feedback and suggestions. Even if the structure is close to being finalised, the e-Guide will always be a work in progress.



GIOIIdta FIOIIIIO © ECCC

GIONATA FIORINO GuiCom Member and on behalf of the ECCO e-Guide Taskforce

# N-ECCO Activities in Barcelona 2017

We are swapping the wonders of canals and bicycles of Amsterdam for the sunny climate of Barcelona for N-ECCO 2017

nce again, N-ECCO is offering an outstanding range of clinical and educational opportunities for nurses attending the ECCO'17 Congress. The programmes for each activity continue to be developed from the evaluation forms of the previous year to meet your needs.

This year the **8<sup>th</sup> N-ECCO School** will again be joined by up to 20 dietitians. Some changes have been made to the agenda to include the role of nutrition in the care of patients with IBD and we are keeping the interactive workshops in the afternoon. The **11<sup>th</sup> N-ECCO Network Meeting** will once more deliver a wide variety of subjects presented by expert speakers from several centres across Europe. The theme of the opening morning session is quality of care, featuring contributions from eminent speakers covering quality indicators in IBD, the use of PROs (Patient Reported Outcome Measures) and the hot topic of remote monitoring in IBD. The mid-morning session continues by addressing special considerations in IBD, with topical presentations exploring fertility and pregnancy, travel and IBD and the low FODMAP diet. Nurses presenting their research abstracts are being allotted a longer time, 15 minutes, in order to encourage discussion. In the final session, a presentation on therapeutic algorithms for the management of IBD patients will be followed by a lively debate of mono vs. combination therapy, a debate I am sure will be extremely interesting!

# The **4<sup>th</sup> N-ECCO Research Forum** will offer a range of sessions. Experienced researchers from across Europe will present their own research concentrating on the methodology of their work, covering systematic reviews, the development of a questionnaire and measurement of conditions. Using the topics identified from the

research priority Delphi survey, the outlines of three to five research projects will be presented and then, in small groups, these projects will be developed into final research protocols. The aim is for experienced researchers to work together with less experienced nurses to develop a full research protocol which may be used when submitting an application for project funding.

The Research Forum will be of special interest to nurses already undertaking research and to nurses wishing to learn more about research or to perform research themselves. The programme allows a generous amount of time for discussion and for networking with experienced research nurses from the United Kingdom, Scandinavia, and outside of Europe. The complete programmes for the 8<sup>th</sup> N-ECCO School, 11th N-ECCO Network Meeting and 4<sup>th</sup> N-ECCO Research Forum can be reviewed on pages 7, 8, 12 and are available on the ECCO'17 Congress Website (www.ecco-ibd.eu/ecco17). There are several other Workshops which can be attended by nurses, such as the 2<sup>nd</sup> D-ECCO Workshop, the 4<sup>th</sup> P-ECCO Educational Course and the 10<sup>th</sup> Y-ECCO Career Workshop, as well as, of course, the main Scientific Programme of the ECCO Congress throughout the rest of the week.



N-ECCO Network Meeting at ECCO'16 © ECCO

At the ECCO'17 Barcelona Congress, one N-ECCO Committee Member will be stepping down: Karen Kemp, current Chair of N-ECCO. Karen was a N-ECCO National Representative prior to joining the N-ECCO Committee in 2014. She has since helped to further develop its profile and portfolio of clinical and educational nursing activities, in particular the introduction of the N-ECCO Research Forum. We would like to thank Karen for her hard work and contribution to N-ECCO. With many forums where up-to-date IBD therapy and research can be explored, Barcelona is the place to be in February 2017! We very much look forward to seeing you there for another excellent educational event and an ideal opportunity for networking with colleagues from around the world.

> KAREN KEMP N-ECCO Chair On behalf of the N-ECCO Committee: Palle Bager, Kay Greveson, Usha Chauhan, Kathleen Sugrue

# Quality indicators for running a paediatric Inflammatory Bowel Disease Centre: Time to set the bar!

### Managing IBD is a complex process, especially in children, who represent a sensitive population

he move towards personalised medicine and the advent of new therapies and technologies require access to an increasing number of laboratory assays and to a diverse multidisciplinary team with expertise in different aspects of the disease.

With the assistance of a Delphi group of 110 paediatric gastroenterologists, P-ECCO tabulated and graded for the first time in children 60 items that any paediatric IBD centre must include, ranked on a scale of importance. Some of the highest ranking items were a minimum of 50 patients or at least 10 new patients per year, a dedicated stand-alone weekly IBD clinic, a 24/7 help line, the accommodation of urgent clinics, the availability of a wide range of multidisciplinary team and access to IBD-specific lab tests such as drug levels.

Several previous surveys reported a significant gap between the recommended IBD management guidance and clinical practice. Patients often receive suboptimal dosing of immunomodulators and 5-ASA and prolonged use of corticosteroids. Many do not undergo appropriate infectious screening before initiation of anti-TNF treatment and there is wide variation

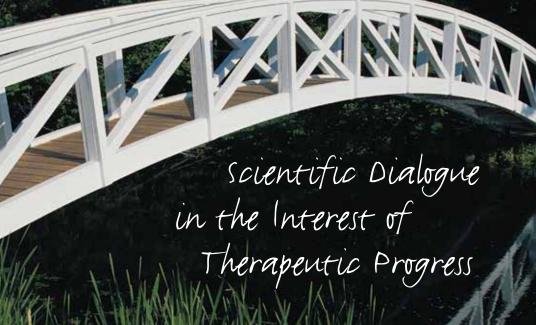
in treatment algorithms between centres. Many IBD centres lack important specialists such as a dedicated IBD nurse, a stoma specialist, a psychologist/social worker and a radiologist. The problem is worse in paediatrics since the number of children with IBD is one-tenth of the adult number. This naturally leads to smaller centres and many do not attain a critical mass of patients to justify the required resources. On the other hand, paediatric-onset disease is more often extensive and its course may be somewhat more challenging to manage. The presence of pubertal delay, growth impairment or osteopenia should dictate aggressive treatment with the aim of promoting mucosal healing. In these cases, the use of nutritional therapy is particularly attractive and requires a specialised team. The use of support groups and other psychosocial interventions is advocated as a means to develop coping skills and improve quality of life. Approximately half of adolescents do not comply with the recommended treatment and thus a particular emphasis should be placed on engaging the adolescent in the treatment plan while enhancing gradual transition to adulthood.

Caring for IBD patients in a dedicated multidisciplinary centre has been shown to improve clinically important outcomes. In addition to a gastroenterologist and a colorectal surgeon, the team should include an IBD-specialist nurse, a nutritionist, a stoma specialist, a radiologist, an endoscopist, a pathologist, a psychologist and a social worker, all with IBD experience. Several other aspects of an IBD centre may improve clinical care, including a 24/7 hot line, a transition clinic, structured weekly clinical rounds, establishment of local treatment algorithms, regular self-assessment of clinically important outcomes, an interactive patient registry and performance of IBD-related research.

Formal IBD multidisciplinary centres and proactive management may reduce disease burden, improve outcomes and thus reduce the disease-related costs. We are hopeful that the P-ECCO quality indicators project will promote standardisation of the care that children with IBD receive globally, while setting the bar at a minimal accepted standard.

> DAN TURNER P-ECCO Chair

# International Symposia and Workshops





Workshop **Future Perspectives in Hepatology:** From Basics to Clinics Essen, Germany January 19 – 20, 2017



Symposium 206 From the New and Complex Concepts to the Real Patient: Science and Clinic Madrid, Spain March 31 – April 1, 2017



Symposium 207 **Mucosal Microbiome and Mucosal** or Systemic Dysfunction: Mechanisms, Clinical Manifestations and Interventions Brisbane, Australia May 19 – 20, 2017

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### IX Gastro-Conference Berlin, Germany October 4 - 7, 2017



Symposium 208 **Eosinophilic Esophagitis** – **Medical and Dietary Treatment** Berlin, Germany October 4 – 5, 2017



Symposium 209 **IBD 2017 – Therapeutic** and Biological Barriers Berlin, Germany October 6 - 7, 2017



Workshop Workshop on Oral, Gastrointestinal and Pulmonary GvHD Regensburg, Germany November 17 – 18, 2017

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# A quick summary of Research Gaps in Diet and IBD

hile the current doctrine of IBD pathogenesis suggests an interaction between environmental factors and gut microbiota in people who are genetically susceptible to develop the illness, dietary exposures have attracted recent interest and are likely to explain the rapid rise in the incidence of IBD. The D-ECCO Working Group, along with other ECCO Experts with expertise in nutrition, microbiology, physiology and medicine, recently reviewed all state-of-the-art literature looking at the role of diet and nutritional therapy in the onset, perpetuation and management of IBD. In this Topical Review, which has reached the late stage of completion, we summarised research gaps in the area of IBD and diet within three major thematic domains:

# 1) Diet as an environmental factor in IBD aetiology

While the literature contains numerous scientific studies exploring the role of host genetics, gut microbiota and diet as isolated factors in the onset of IBD, there is scarce evidence for the interplay between these factors. The advent of systems biology and bioinformatics tools now allows exploration of research questions which were left unanswered previously. Similarly, very few studies which have portrayed gut microbiota in IBD patients have accounted for dietary influences, an established determinant of gut microbiota composition and metabolic activity. More evidence is also required on how early-life dietary exposures might shape gut microbiota colonisation and modify risk of IBD later in life, using a longitudinal cohort design.

There is a need for more basic science to delineate mechanisms by which dietary components increase or reduce the risk of IBD. Extrapolation of findings from in vitro and animal-based models of colitis to human IBD is essential before a causal association can be established. Moreover, studies of the role of diet in the pathogenesis of IBD now need to be expanded from major nutrients to nonnutrient components of habitual diet and others introduced into the human food chain via industrialisation, processing and storing of food.

### 2) Diet as therapy in IBD

The role of diet in the management IBD encompasses the following main of therapeutic modalities: The use of exclusive (EEN) and supplementary enteral nutrition for induction and maintenance of remission or as treatments in CD and the use of recently developed whole food-based elimination diets for induction and maintenance of remission. EEN is now the established induction treatment of active CD, with an excellent safety profile and high response rate. Understanding the epigenetic, immunological and microbiological mechanisms that potentially mediate the mode of EEN is a research priority which will offer the paradigm for development of alternative therapies. Studies monitoring food reintroduction following EEN need to be performed to identify potential dietary triggers of disease relapse. More evidence is also required to evaluate the use of EEN in adult patients with CD and its potential use in UC, complicated CD, and pre- and postoperative settings. Exploring the effectiveness of supplementary enteral nutrition as a monotherapy or in combination with medical therapy for prevention of relapse in IBD is of particular interest. The optimal regimen of supplementary enteral nutrition for maintenance of CD, including the dose, composition, duration, method of delivery of feeding, and nature of the accompanying oral diet, needs to be studied. While there is emerging interest from patients and health professionals in the use of elimination diets, well-designed clinical trials are still required to evaluate efficacy and understand the mechanisms of such novel dietary therapies.

# 3) Nutritional assessment and supportive nutritional therapy in IBD

Although most of the focus around nutrition in IBD has primarily concentrated on the previous two thematic areas, nutritional assessment and supportive nutritional therapy are cornerstones in the management of IBD patients. There are limited data from good quality studies on the progression of undernutrition following diagnosis and whether this is predictive of disease outcomes, particularly in the elderly. Development of novel biomarkers of micronutrient status is needed to overcome limitations of plasma measurements in the presence of systemic inflammatory response. Supportive nutritional therapy is frequently required in IBD patients with short bowel syndrome or fistulising disease. There is a need for development of better biomarkers to predict or diagnose short bowel syndrome in IBD and for better evidence on optimal preand postoperative nutritional management of such patients and those with fistulising CD. Identification of functional bowel symptoms in inactive IBD is of utmost importance to ensure that unnecessary and potentially harmful treatment strategies are avoided. Conversely, presence of active IBD should be excluded before determining that symptoms are functional in nature. There is an interest in the use of a low FODMAP diet, extending beyond irritable bowel syndrome (IBS) to functional IBD. High-quality clinical trials are needed to demonstrate whether a low FODMAP diet is effective (in terms of symptom resolution, quality of life and acceptability) and safe (with respect to gastrointestinal health and nutritional adequacy) for the management of IBS in patients with inactive IBD.

### KONSTANTINOS GERASIMIDIS D-ECCO WG Member

# Activity Scores in Inflammatory Bowel Disease (IBD): A Challenge for Pathologists

In patients with Ulcerative Colitis (UC), mucosal healing has emerged as a major therapeutic goal. Disease activity in UC has traditionally been evaluated on the basis of clinical, haematological and endoscopic parameters. Endoscopic bowel healing is associated with favourable long-term outcomes in UC. However, the correlation between histological and endoscopic mucosal healing is not very good, with persistence of microscopic inflammation in 16%–100% of patients with endoscopically quiescent disease [1]. There is growing interest in the assessment of histological disease activity based on the concept that histological healing is associated with better clinical outcomes in UC and may be the ultimate therapeutic goal in UC. For this reason, the ECCO Consensus on the Histopathology of IBD recommended that "the pathology report should include some information on the level of activity in the biopsies in order to assess both the effect of therapy and the risk of relapse" [2].

However, before histological disease activity is accepted as a useful parameter in research and clinical care in IBD, a validated instrument for its measurement must be developed. Although more than 20 indices for the assessment of histological disease activity have been developed, their operating properties have not been systematically validated. The two most commonly used histological scores are the Geboes Score (GS) and the Riley Score (RS). More recently, Gramlich et al. and Gupta et al. developed other systems for scoring. Most of these scores of activity have numerous items, may combine chronic and acute changes, and assess epithelial and inflammatory features. In a recent study, Bressenot et al showed that intra-observer reproducibility and inter-observer agreement for these four main available histological UC activity indexes are very good and that correlation between these histological indexes is strong [3]. Histological items that showed better correlation were "erosion/ulceration" and "acute inflammatory cells infiltrate/neutrophils in lamina propria". In a second step, the same group developed a simple descriptor scoring system, the "Nancy Index", which shows good intra- and inter-observer reliability [4]. Briefly, the Nancy Index is composed of three histological items ("ulceration", "acute inflammatory cells infiltrate" and "chronic inflammatory infiltrate") defining five grades of activity ranging from grade 0 (absence of significant histological disease activity) to grade 4 (severely active disease). Another interesting histological index, the Robarts Histopathology Index (RHI), has been developed concomitantly by another group [5]. Component items of this latter index are "chronic inflammatory infiltrate", "lamina propria neutrophils", "neutrophils in the epithelium" and "erosion and ulceration".

It is worth noting that these two histological indices have been tested by highly experienced and intensively trained pathologists on the item scoring conventions. Thus, prospective cohort studies should define the feasibility of their application among non-expert pathologists who report gastrointestinal pathology. Moreover, further studies are needed to define the predictive value of these systems in assessing outcomes in UC and correlating histological with endoscopic scoring.

Besides the need for a validated instrument to measure histological disease activity in IBD, the necessity of a correct methodological approach in the evaluation of colonic biopsies should be emphasised, as mentioned by V. Villanacci (H-ECCO WG member) in a previous letter. In this

context, the adequate and correctly oriented number of biopsies is of paramount importance (as highlighted in the ECCO statements 4A and 4B) [2].

In summary, histological assessment of disease activity and severity is likely to become an integral part of the management of UC and constitutes a substantial challenge for pathologists.

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MAGALI SVRCEK H-FCCO WG Member

# <sup>th</sup> S-ECCO IBD Masterclass

### February 16, 2017

The 6<sup>th</sup> S-ECCO IBD Masterclass, the meeting of the surgeons' branch of ECCO, will take place on February 16, 2017 in Barcelona, within the European Crohn's and Colitis Organisation's annual Congress. The S-ECCO IBD Masterclass is the annual scientific forum for professional discussions on the surgical aspects of the treatment of patients with IBD, emphasising the multidisciplinary approach in these patients. This unique meeting is devoted to IBD surgery and is attracting an increasing number of surgeons from around the globe each year owing to the high-quality scientific content and the interchange of knowledge and ideas between surgeons and other professionals who take care of IBD patients.

The upcoming S-ECCO IBD Masterclass will focus on "Difficult decision making in **IBD**" and will include multiple debate sessions in which two or more talks address the same topic, often one presented by a surgeon and the other by a gastroenterologist. The topics to be debated will include the timing of surgery in Acute Severe Colitis, the best surgical technique for patients with Ulcerative Colitis, the approach to patients with Crohn's Disease following successful percutaneous drainage of intra-abdominal abscess, surgical technique for ileocolonic anastomosis and the approach to fissures and haemorrhoids in Crohn's patients. In addition, the scientific programme will include sessions focussing on cutting-edge technical



5<sup>th</sup> S-ECCO IBD Masterclass at ECCO'16 © ECCO

aspects of surgery for Inflammatory Bowel Diseases, including top-down versus bottom-up pouch surgery and TAMIS (trans-anal minimally invasive surgery). For the first time, the scientific programme will include a free paper session of abstracts related to surgery of IBD, and the trials update session, which was very successful last year, will briefly present ongoing trials in IBD surgery.

The 6<sup>th</sup> S-ECCO IBD Masterclass faculty includes a large number of leading IBD surgeons from Europe and globally and leading

gastroenterologists, which assures a fascinating programme for both IBD surgeons and all other professionals caring for patients with IBD. This meeting is also a good opportunity to meet old and new colleagues and friends. We look forward to seeing you in the 6<sup>th</sup> S-ECCO IBD Masterclass hall in Barcelona.

# Dear Y-ECCO Friends,

t's the middle of the summer, but already our thoughts are turning to the next ECCO Congress. Abstract submission opens on August 17 and there are a number of opportunities that Y-ECCO Members need to be aware of. As in previous years, there will be five awards for the best abstracts submitted by a Y-ECCO Member, with two of these reserved specifically for Basic Science Abstracts. The five winners will each receive free registration at the next ECCO Congress. Entry for these awards is simple - just make sure to tick the box indicating that you're a Y-ECCO Member at the time you submit your abstract. Y-ECCO Basic Science Abstracts will also all be considered for the Y-ECCO Basic Science Workshop taking place on Wednesday, February 15 (the day before the main Congress). This workshop will focus on increasing the clinical relevance of research findings with two key themes: "Genetics in IBD - Moving beyond GWAS" and "Identifying novel targets in IBD using patient samples". State of the art talks and expert moderation will be provided by our two outstanding invited speakers, Miles Parkes and Gijs van den Brink.

And if you're still undecided about whether to come to Barcelona a day early for the workshop (frankly, who needs a better reason – this is Barcelonal), the Y-ECCO Basic Science Workshop will be followed by our Y-ECCO Career Workshop, which this year will focus on the practical skills and tips required to submit the perfect grant application. Again, we are delighted that we will be joined by an expert panel, which will include Séverine Vermeire, Gerhard Rogler and Helen Terry. As usual, we'll all be heading to a local meeting spot for some drinks and food after the workshop – a great chance to relax with fellow Y-ECCO Members.

Lastly, two important and exciting things to tell you. First, we are inviting Y-ECCO Members to submit their proposals for a new joint initiative with ClinCom that will support a proposal for a new study from a Y-ECCO Member. This doesn't have to be something huge (save your big ideas for a grant submission) but rather something simple, interesting and, above all, deliverable. What's that quirky idea you've always wanted to test in IBD? See details of the call below – but don't for one second put off applying! And finally – Y-ECCO is now on Twitter. Or about to be – we've set up the account but need to get going! We plan to use the feed to communicate with you (and others who aren't lucky enough to be Y-ECCO Members), particularly around the time of the Congress. Follow us @Y\_ECCO\_IBD. As ever, do get in contact (*ecco@ecco-ibd.eu*) – or tweet us!



Tim Raine © ECCO

Isabelle Cleynen © ECCO

ISABELLE CLEYNEN Y--FCCO Literature Review Admin

TIM RAINE Y-ECCO Chair

# Y-ECCO Literature review

### Dear (Y-)ECCO Members,

We are happy to welcome you in the Y-ECCO Literature Review section of ECCO News. In this section, Y-ECCO Members highlight and summarise recent landmark articles within the field of IBD. The articles cover different topics, including basic science, epidemiology, clinical phase 3 trials, endoscopy, surgery... As we are always looking for people that want to participate in this initiative, Y-ECCO Members that are interested can contact Isabelle (isabelle. cleynen@kuleuven.be). The idea is you choose a recent article, and summarize the key findings and importance of the paper in max 1000 words. Together with the review, a short selfdescription and picture will be published.

### Gene-microbiota interactions contribute to the pathogenesis of Inflammatory Bowel Disease

Chu H, Khosravi A, Kusumawardhani IP, Kwon AH, Vasconcelos AC, Cunha LD, Mayer AE, Shen Y, Wu WL, Kambal A, Targan SR, Xavier RJ, Ernst PB, Green DR, McGovern DP, Virgin HW, Mazmanian SK

Science. 2016 May 27;352:1116–1120.

### Introduction

Large genetic studies investigating susceptibility to Inflammatory Bowel Disease (IBD) have uncovered the important role of impaired microbial handling by the gut immune system in the pathogenesis of IBD [1]. Gene-microbiota interactions have since been the subject of investigation. Important IBD susceptibility genes have been linked to alterations in the gut microbiota composition: the intracellular bacterial peptidoglycan sensor gene NOD2 [2], regulation of autophagy genes ATG16L1 [3] and IRGM4 [4], the immune response gene CARD9 [5, 6] and the intestinal mucus layer gene FUT2 [7].

This functional study aimed to explain the aberrant immune response to commensal microbiota in patients with functional IBD susceptibility variants in NOD2 and ATG16L1 by investigating the interactions between these two genes and the commensal bacterial species Bacteroides fragilis of the family Bacteroidaceae. Bacteroides fragilis produces polysaccharide A (PSA), which is packaged in vesicles called outer membrane vehicles (OMVs). These OMVs with PSA can induce the production of the anti-inflammatory cytokine IL-10 in CD4+Foxp3+ regulatory T cells (Tregs). This mechanism was studied in combination with alterations in NOD2 and ATG16L1 in mouse dendritic cells, mice and human dendritic cells.

### **Experiments and key findings**

In the first experiment, bone marrow-derived dendritic cells (BMDCs) from wild-type and ATGI6L1deficient mice were pulsed with either (1) wild-type Bacteroides fragilis OMVs with PSA, (2) OMVs from mutant Bacteroides fragilis lacking PSA or (3) purified PSA not encapsulated in OMVs. These BMDCs were all co-cultured with CD4+ T cells. The OMVs from wild-type Bacteroides fragilis were able to induce IL-10 production in wild-type BMDCs, but not in ATGI6L1-deficient BDMCs. PSA-deficient OMVs were not able to induce significant IL-10 production in either wild-type BMDCs or ATGI6L1-deficient BDMCs. However, purified PSA not encapsulated in OMVs was able to induce LI-10 production in wild-type BMDCs as well as in ATGI6L1-deficient BMDCs.

In the second experiment, the autophagy pathway used by the OMVs to induce CD4+Foxp3+IL-10+ Tregs was investigated. There are different autophagy

pathways: canonical and non-canonical. ATG16L1, ATG5 and ATG7 participate in both the canonical and non-canonical autophagy pathways. The experiment using BMDCs showed that the canonical autophagy-specific genes Ukl1, Fip200 and Atg14 were not required for OMVs with PSA to induce CD4+Foxp3+IL-10+ Tregs. The non-canonical autophagy pathway, LC3-associated phagocytosis, is specifically activated by microbial ligands that are delivered as particles instead of soluble molecules. LC3-associated phagocytosis requires RUBICON, which represses the canonical autophagy. In BMDCs lacking RUBICON, Treg responses could not be induced.

In the third experiment, colitis was induced in wildtype mice, ATGI6L1-deficient mice and NOD2-deficient mice using 2,4-dinitrobenzenesulfonic acid (DNBS). All these mice were subsequently treated with OMVs from wild-type Bacteroides fragilis via oral gavage. The wild-type OMVs were able to ameliorate the colitis in wild-type mice treated with DNBS. However, in neither ATGI6L1-deficient mice nor NOD2-deficient mice were OMVs from wild-type Bacteroides fragilis able to cause the same anti-inflammatory effect.

In the last experiment, human monocyte-derived DCs from both Crohn's Disease patients and healthy controls with either the ATG16L1 T300A risk variant or the ATG16L1 T300 protective variant were pulsed with OMVs from wild-type Bacteroides fragilis, OMVs lacking PSA and purified PSA. Consistent with the mouse experiments, the induction of IL-10 from Foxp3+ Tregs was dependent on having the ATG16L1 T300 protective variant and was not influenced by the disease status of the monocyte-derived dendritic cells.

### **Conclusions and commentary**

This study shows that both mouse and human DCs require functional ATG16L1 and NOD2 for the induction of anti-inflammatory CD4+Foxp3+IL-10+ Tregs in response to Bacteroides fragilis PSA in OMVs. In addition, the experiments show that the Bacteroides fragilis OMV-based induction of CD4+Foxp3+IL-10+ Tregs uses the non-canonical LC3associated phagocytosis pathway.

It is very likely that there are more gene-microbiota interactions underlying the pathogenesis of IBD, many of which are still unknown. Moreover, a large number of known genetic variants and gut microbiota alterations are associated with IBD [1, 8, 9]. Therefore, there is a need for more functional gene-microbiota interaction studies to clarify the role of the gut microbiota in the aetiology and pathogenesis of IBD. References

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### Floris Imhann

Floris Imhann obtained his MD from the University of Groningen, the Netherlands, in 2011. Currently, he is a PhD candidate at the Department

of Gastroenterology of the University Medical Center Floris' Groningen. research focusses on the gut microbiota in Inflammatory Bowel Disease patients



Floris Imhann © Floris Imhann

### Proteomic analysis of ascending colon biopsies from a Paediatric Inflammatory Bowel Disease inception cohort identifies protein biomarkers that differentiate Crohn's Disease from UC

. Starr AE, Deeke SA, Ning Z, Chiang CK, Zhang X, Mottawea W, Singleton R, Benchimol EI, Wen M, Mack DR, Stintzi A, Figeys D. Gut. 2016 May 23. doi:10.1136/gutjnl-2015-310705

### Introduction

Accurate differentiation between Crohn's Disease (CD) and Ulcerative Colitis (UC) is important to ensure early and appropriate therapy. Especially in children, this differentiation can be challenging due to an atypical or less distinctive presentation [1]. Biomarkers have been sought to complement current IBD diagnostic tools, such as endoscopy, imaging and histology, in order to reduce ambiguous diagnosis of IBD. Several studies have successfully identified promising biomarkers in stool, serum and tissue. However, none of the identified biomarkers have been implemented in clinical daily use [2]. This study reports the first proteomic analysis in paediatric biopsies, aiming to identify candidate biomarkers that may contribute to the accuracy of IBD diagnosis.

### Methods

This cross-sectional study included 99 children (39 control, 30 CD, 30 UC), from whom biopsies (n=124) were taken at the ascending colon before initiation of any treatment. If possible, biopsies were also obtained from non-inflamed regions in the case of IBD patients (in 23 CD and in 2 UC patients). Frozen biopsies were lysed by mechanical homogenisation and proteins isolated. Relative quantification of peptides (obtained through tryptic digestion) was performed using an Orbitrap Elite mass spectrometer (MS). Raw MS data were analysed against the human UniProt database. Proteomic data of an inception cohort (n=50) were used to develop a mathematical model for disease classification via three different models [Random Forest, Support Vector Machine and Partial Least Squares Discriminant Analyses (PLSDA)]. Proteins identified in all three models were considered to be candidate biomarkers (252 in total). Biomarker panels [to distinguish IBD from control (panel 1) and CD from UC (panel 2)] were designed with PLSDA models using a step-forward method. An independent cohort (n=49) was used to validate the discovery-trained PLSDA models.

With the intent of translating their findings into the clinical setting, the authors finally measured the absolute amount of two of the identified candidate biomarkers using ELISA in the validation cohort biopsy samples. Importantly, a requirement of the study design was that all candidate biomarkers had to be quantified in ≥95% of the biopsies. This is in contrast to previously studied biomarkers like pANCAs or ASCAs, which are identified in a limited number of patients (range 2%-85%) [3]. Although this prerequisite may have reduced the number of candidate biomarkers, the approach reinforced the potential applicability of these prediction panels in diagnosis at initial endoscopy.

### **Key findings**

Proteomic analysis of the intestinal biopsies led to the identification of a panel of 12 proteins which can be used to differentiate paediatric patients with CD from those with UC [area under the curve (AUC) of 86%, sensitivity 73% and specificity of 87% in the validation cohort]. One of the panel proteins, metallothionein-2 (MT2), a metallothionein previously linked to IBD with contradictory findings [4], was quantified in all validation cohort samples by ELISA and was significantly higher in patients with CD compared to those with UC. Pathway analysis of the initial 252 diagnostic candidate biomarkers suggested elevated fatty acid metabolism in paediatric CD and elevated energy and amino acid metabolism in paediatric UC A panel of five proteins was sufficient to differentiate patients with and without IBD (AUC 99%, sensitivity 95% and specificity 97% in the validation cohort). Visfatin, a novel adipokine that is also known as pre-B cell colony-enhancing factor and has recently been associated with the development of osteoporosis in IBD [5], was measured by ELISA in 24 samples of the validation cohort and found to be significantly higher in IBD compared to control biopsies. No significant difference was found in the amount of visfatin between inflamed and non-inflamed biopsies in CD patients. The identified canonical pathways differing IBD from healthy controls seemed mainly related to metabolic processes and catalysis. Also, the energy metabolism was clearly represented in the IBD-related proteome, including inorganic pyrophosphatase, visfatin and UDP-glucose 6-dehydrogenase. Both the first and the second panel contained components of the fatty acid metabolism, which is not necessarily a surprise as its contribution to the pathogenesis of IBD has already been established [6].

Looking beyond the issue of differential diagnosis, the authors observed 118 proteins showing a correlation with disease severity (defined by either PCDAI or PUCAI for CD and UC respectively). Thirty-nine of them were identified as biomarker candidates, four of which were part of one of the two previously discussed biomarker panels. The relative expression of panel 2 protein MT2 correlated with CD severity, whereas HNRNP H3 (heterogeneous nuclear ribonucleoprotein H3) expression was inversely related to UC severity. Likewise, the expression of visfatin and inorganic pyrophosphatase, both members of panel 1, showed a significant correlation with disease severity in CD.

### Conclusions

The results of this study clearly strengthen the potential development of an accurate diagnostic

tool in paediatric IBD, though further research is required in order to translate these findings into the clinical setting. Among the upfront challenges are the development and validation of a tool allowing absolute quantification of the identified proteins and the development of an appropriate multivariable prediction model for disease classification. However, confirmation of the current panels in a validation cohort and the ELISA results for two of the identified proteins suggest the likelihood of translating these findings to the clinic.

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### Bram Verstockt

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Department of Medicine at Cambridge University, UK.



Bram Verstockt © Bram Verstockt

### Ozanimod induction and maintenance treatment for Ulcerative Colitis

Sandborn WJ, Feagan BG, Wolf DC, D'Haens G, Vermeire S, Hanauer SB, Ghosh S, Smith H, Cravets M, Frohna PA, Aranda R, Gujrathi S, Olson A N Engl J Med. 2016;374:1754-62

### Introduction

Ozanimod is a new oral agonist of the sphingosine-1-phosphate receptor sub-types 1 (s1p1 receptor) and 5 (s1p5 receptor) that induces peripheral lymphocyte sequestration, potentially decreasing the number of activated lymphocytes circulating to the gastrointestinal tract [1]. S1p1 receptor agonists induce internalisation and degradation of the s1p1 receptor, rendering the B and T lymphocytes incapable of migrating from secondary lymphoid tissues, which leads to a reversible reduction in circulating lymphocytes in the blood [2-4]. Ozanimod has been used in the treatment of patients with relapsing multiple sclerosis and showed a dose-dependent reduction in circulating lymphocytes that was associated with significant reduction in inflammatory and neurodegenerative brain lesions [5]. This study was a phase 2 trial of induction and

maintenance therapy at 57 centres across 13 countries using ozanimod in 197 adults with moderate to severe Ulcerative Colitis. The trial was over a 32-week period in which patients were randomly assigned in a 1:1:1 ratio to receive ozanimod at 0.5 mg or 1 mg or placebo daily. Inclusion criteria: Patients of 18–75 years of age who had UC with a Mayo Clinic score of 6–12 and an endoscopic sub-score of 2 or 3. Treatment with oral aminosalicylates or prednisolone was required to be at stable doses. Patients receiving biologics, azathioprine, mercaptopurine or methotrexate were required to discontinue these agents five half-lives before starting the trial.

The primary outcome measure was clinical remission, defined as a Mayo Clinic score of ≤2 with no subscore >1 at 8 weeks. Secondary outcome measures at week 8 were clinical response in the Mayo Clinic score  $\geq$ 3 points and  $\geq$ 30% from baseline, with a decrease in the rectal bleeding sub-score  $\geq$ 1 point or a subscore ≤1. Explorative outcome measures at week 32 included clinical response, histological remission, clinical remission, mucosal healing and change in the Mayo Clinic score. Absolute lymphocyte count, CRP, calprotectin and lactoferrin were also measured.

### **Key findings**

Clinical remission occurred in 16% of patients who received 1 mg of ozanimod and in 14% of patients who received 0.5 mg of ozanimod, as compared with 6% of those receiving placebo (p=0.048 and p=0.14 respectively for comparison of the two doses with placebo). There were therefore no statistically significant treatment differences in primary outcomes between ozanimod and placebo.

Concerning secondary endpoints, clinical response at week 8 occurred in 24/65 (37%) in the placebo group compared with 35/65 (54%) who received 0.5 mg of ozanimod (p=0.06) and 38/67 (57%) who received 1 mg of ozanimod (p=0.02). Mucosal healing at week 8 occurred in 8/65 patients (12%) in the placebo group as compared with 18/65 (28%) in the group that received 0.5 mg of ozanimod (p=0.03) and 23/67 (34%) in the group that received 1 mg of ozanimod (p=0.002). Histological remission, defined by a Geboes score of less that 2, at week 8 occurred in 7/65 patients (11%) in the placebo group as compared with 9/65 (14%) in the group that received 0.5 mg of ozanimod (p=0.63) and 15/67 (22%) in the group that received i mg of ozanimod (p=0.07). Absolute lymphocyte counts in blood decreased by a mean of 32% from baseline to week 8 in patients who received 0.5 mg of ozanimod, and by 49% in patients who received 1 mg of ozanimod.

No differences in safety were observed among the groups. One patient in the 0.5 mg ozanimod group who has evidence of pre-existing bradycardia had first-degree AV block and sinus bradycardia after day 8. This event was asymptomatic and resolved without the need of intervention

### Conclusion

Ozanimod at a dose of 1 mg was associated with a slightly higher rate of clinical remission in patients with moderate to severe Ulcerative Colitis as compared with patients receiving placebo, without the difference reaching statistical significance. At week 32, patients receiving 1 mg of ozanimod continued to have higher rates of clinical remission, clinical response, mucosal healing and histological remission when compared with the placebo group. Limitations of the study included the time point analysed in the primary outcome. Eight weeks was used, and this time point may not have been long enough for the drug to target the lymphocyte

tracking. Another limitation is that it is not possible to assess the long-term safety profile of the drug due to the relatively small number of patients analysed and the short follow-up period. Interestingly, ozanimod was associated with an absolute drop in lymphocyte count, which is a finding consistent with the mechanism of the drug. This, however, raises issues about long-term risk of opportunistic infections. Ozanimod shows some potential for treatment in moderate to severe Ulcerative Colitis but larger studies with longer follow-up data are needed.

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### Jonathan Segal

Jonathan Segal is а gastroenterology trainee and research fellow at St. Marks Hospital in London. He has a particular interest in the microbiota in IBD, and his current research is understanding the role of the microbiota in the development of Pouchitis.



Jonathan Segal © Jonathan Sega

# Y-ECCO/ClinCom Call for Y-ECCC researc h propo

### An exciting new opportunity for Y-ECCO Members to propose and perform a brief, focussed research project with ClinCom/ Y-ECCO guidance and full ECCO endorsement and support.

At the ECCO Congress in Amsterdam, Y-ECCO and ClinCom jointly conducted a survey of attendees, assessing access to investigations in IBD across Europe and beyond (the ACCID survey). Targeted responses from a large sample of gastroenterologists, surgeons, nurses and paediatricians generated interesting results that are currently being prepared for abstract submission and publication.

Based on this success, we are now asking Y-ECCO Members to propose a small research study (which may or may not be a survey). The best proposal (as assessed by members of the Y-ECCO Committee) will then be supported by the ClinCom experts and developed with ClinCom and ECCO Governing Board support. If the winning proposal is a survey, this will be performed at the ECCO Congress 2017

and the winners will receive the full support of the ECCO Team in distributing, collecting and collating data, with additional support as needed during data analysis.

This is a fantastic opportunity for Y-ECCO investigators to access the resource of the ECCO Office and the entire ECCO Membership. It is anticipated that the successful study will be submitted for publication, e.g. in the Journal of Crohn's and Colitis.

Applications need to be well thought through and structured, and should address a current and pertinent question in IBD. The research question does NOT need to be huge or overambitious - we would welcome simple but interesting proposals that are deliverable.

Please submit proposals (maximum 1 side of A4) structured as:

- Background
- Aim(s)
- Methodology
- Other matters (including any relevant financial details)

Please send all proposals to ecco@ecco-ibd.eu. Please keep an eye out for details and specific deadlines in our regular eNewsletters. No covering letter is required. Just do it!

Please note that although no financial support is provided, reasonable printing costs and activities related to any distribution or collection of materials and data entry will be borne by ECCO.

> NIK SHENG DING Y-ECCO Member

# ECCO Country Member Profiles



### **Identity card**

### Country: Denmark

- Name of group: no specific interest group for IBD, Gastroenterologists with an interest in IBD are organised in "Dansk Selskab for Gastroenterologi og Hepatologi" (DSGH), which is our national 💈 society of gastroenterology and hepatology
- Number of active members: approx. 750 members of DSGH
- Number of meetings per year: one annual meeting lasting 2 days, three minor meetings throughout the year, some of which have an IBD focus
- · Name of president and secretary: Inge Nordgaard-Lassen (President)
- National Representatives: Jørn Brynskov, Torben Knudsen
- · Joined ECCO in: Denmark has been a longstanding member of ECCO, having joined in 2004
- Incidence of IBD in the country (if available): Crohn's Disease approx. 10/100,000, Ulcerative Colitis approx. 15/100,000

### **Identity card**

### Country: United Kingdom

- · Name of groups: British Society of Gastroenterology (BSG) IBD
- Section, IBD Clinical Research Group (CRG) • Number of active members: The BSG IBD section has 695 full members and 228 trainee members. There are 398 UK Members of ECCO.
- Number of meetings per year: BSG IBD section committee meetings, monthly teleconference, and annual full BSG meeting.
- Name of president and secretary: Barney Hawthorne (IBD section chairman), John Mansfield (Secretary, IBD CRG chairman)
- National Representatives: Barney Hawthorne; Tariq Ahmad
- Joined ECCO in: 2003
- Incidence of IBD in the country (if available): There are no national level incidence studies but regional UK studies show a continuing rise in the incidence of both CD and UC, with rates of 6.6-8.3×105/yr (CD) and 12–14×105/yr (UC).

### **Identity card**

### Country: Norway

- Name of group: The Norwegian IBD interest group is organised within the Norwegian Gastroenterological Association (Norsk Gastroenterologisk forening NGF)
- Number of active members: approximately 370 active members, The IBD interest group consists of 8 members elected at the NGF general assembly (four members elected every second year)
- Number of meetings per year: the IBD-interest group has four meetings per year: one national meeting, one to two regional meetings in the five different health regions in Norway
- Name of president and secretary: Birgitte Seip (President), Christine Slinning (Secretary), Petr Ricanek (Leader of the IBD interest group)
- National Representatives: Kristin Kaasen Jørgensen, Marte Lie Høivik
- Joined ECCO in: 2005
- Incidence of IBD in the country (if available): UC 12,8/100.000 persons/ year, CD 6/100.000 persons per year. These are data from 1994 (the IBSEN study). No updated incidence numbers are available.

### **Identity card**

### Country: Poland

- Name of groups: Polish Society of Gastroenterology, IBD Section, Polish Paediatric Society of Gastroenterology, Hepatology and Nutrition, IBD Section
- Number of active members: 30 adult/30 paediatric
- Number of meetings per year: One meeting every 2 years
- Name of president and secretary: A. Dabrowski (President), J. Kwiecien (Secretary)
- National Representatives: Edyta Zagórowicz, Jarosław Kierkuś
- Joined ECCO in: 2004
- Incidence of IBD in the country (if available): About 700 cases of UC per year and about 180-250 CD cases per year. The total number of patients with IBD exceeds 50,000

### Questionnaire – DENMARK



# What has changed since your society became an ECCO Country Member?

There are many Danish participants in ECCO Events. Danish clinicians and researchers are active on ECCO Committees and take part in ECCO-initiated research activities and projects.

### What are the benefits to you of being an ECCO Country Member?

Participation in educational events organised by ECCO and networking regarding science.

Is your society making use of the ECCO Guidelines?

Yes. The Guidelines are used in the daily work at the hospitals and they are cited in many of our national guidelines.

Have you developed research projects with other countries through your ECCO Country Membership?

Yes.

Have you developed educational activities with other countries through your ECCO Country Membership?

Copenhagen has hosted an ECCO Congress

What are your main areas of research interest?

- Epidemiology
- Patient-reported outcomes in IBD
- Cell biology and immune pathogenesis
- Therapeutic drug monitoring

# Does your centre or country have a common IBD database or bio bank?

A national database covering patients treated with biologics has just been established.

Torben Knudsen © Torben Knudsen

What are your aims for the future?

We will try to establish a national Danish interest group covering IBD. *How do you see ECCO helping you to fulfil these aims?* 

Inspiring discussions and inputs from members of national groups in other ECCO Member Countries

What do you use ECCO for?

Networking, congresses, scientific collaboration

How do you use the things/services that ECCO has to offer?

Participation in congresses, meetings and educational events. Use of website solutions.

JØRN BRYNSKOV TORBEN KNUDSEN ECCO National Representative, Denmark

### **Questionnaire – NORWAY**



# What has changed since your society became an ECCO Country Member?

The general awareness of and knowledge on IBD is increasing among Norwegian gastroenterologists. IBD research is also increasing both in quality and quantity. This is reflected in an increasing number of Norwegian delegates at the ECCO-conferences.

### What are the benefits to you of being an ECCO Country Member?

- Access to a large society of researchers
- Access to the ECCO platforms for education and research, including the clinical guidelines
- The possibility to host ECCO work-shops
- The possibility to build an international network for research and to contribute in multinational research projects
- ECCO is a source of inspiration for fellows aspiring a career in IBD research

### Is your society making use of the ECCO Guidelines?

Yes. The Guidelines are used in the daily work at the hospitals and they are cited in many of our national guidelines.

Have you developed research projects with other countries through your ECCO Country Membership?:

### Yes.

Have you developed educational activities with other countries through your ECCO Country Membership? Copenhagen has hosted an ECCO Congress

What are your main areas of research interest?

- Epidemiology
- Patient-reported outcomes in IBD
- Cell biology and immune pathogenesis
- Therapeutic drug monitoring



Norwegian Gastroenterological Association © Norwegian Gastroenterological Association

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A national database covering patients treated with biologics has just been established.

### What are your aims for the future?

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Inspiring discussions and inputs from members of national groups in other ECCO Member Countries

What do you use ECCO for?

Networking, congresses, scientific collaboration

How do you use the things/services that ECCO has to offer?

Participation in congresses, meetings and educational events. Use of website solutions.

KRISTIN KAASEN JØRGENSEN MARTE LIE HØIVIK ECCO National Representative, Norway

### Questionnaire – POLAND

### What has changed since your society became an CCO Country Member?

Thanks to ECCO Membership, Poland has become a more significant country in the IBD field, both in the decision-making process and in terms of international collaboration.

# What are the benefits to you of being an ECCO Country Member?

ECCO Membership has opened new opportunities to the Polish IBD Team. Trainees annually participate in the IBD

Intensive Advanced Course, Polish representatives have participated in the creation of ECCO Guidelines (on Paediatric UC and CD, Malignancies, Surgical UC and CD, and Ulcerative Colitis), and one Y-ECCO Member took advantage of the ECCO Travel Award 2015. New collaborations and partnerships have been established.

### *Is your society making use of the ECCO Guidelines?* Yes.

Have you developed links with other countries through your ECCO Country Membership?

Yes.

Have you developed research projects with other countries through your ECCO Country Membership? Yes (France – EO-IBD, for example)

Have you developed educational activities with other countries through your ECCO Country Membership? Yes (e-CCO Learning Platform – malignancy panel)

*Has your country been involved in a fellow exchange through ECCO?* Yes (ECCO Travel Award 2015)

### What are your main areas of research interest?

- Biologic therapy (both originators and biosimilars)
- Nutritional therapy in IBD
- IBD and PSC
- Histopathology in IBD
- Surgical treatment in paediatrics
- Epidemiology (including EO-IBD)

# Does your centre or country have a common IBD database or bio bank?

### Yes – centre database

# What are your most prestigious/interesting past and on-going projects?

- Safety and efficacy of biologics and biosimilars in children with IBD
- Impact of biologic therapy on mucosal healing
- Participation in clinical trials to evaluate new biologics in IBD (golimumab)

### Which ECCO Projects/Activities is the group currently involved in?

- Transition Topical Review
- I-CARE

### What are your aims for the future?

- Exit Strategies for IBD Treatments Topical Review
- IBD and PSC





Edyta Zagórowicz Source: courtesy of Edyta Zagórowicz

Jarosław Kierkuś © ECCO

### How do you see ECCO helping you to fulfil these aims?

Through ECCO Grants and/or Travel Awards and collaboration with other Member Countries

# What do you use ECCO for? Network? Congress? How do you use the things/services that ECCO has to offer?

Each year more and more physicians, especially Y-ECCO Members, participate in annual ECCO Congresses and Workshops (IBD Intensive Advanced Course) to deepen their knowledge on IBD. Also, the number of ECCO Members from Poland is increasing every year. Our Members try to be actively involved in various ECCO Projects (e.g. Guidelines Working Groups, e-CCO Learning Platform) and various IBD fields (surgery, histopathology). The number of publications on IBD from Poland in peer-reviewed journals has been steadily growing.

EDYTA ZAGÓROWICZ JAROSŁAW KIERKUŚ ECCO National Representative, Poland

### **Questionnaire – UNITED KINGDOM**



# What has changed since your society became an ECCO Country Member?

The UK National IBD Audit Programme has undergone 4 rounds between 2006 and 2015. It has documented (and helped to drive) a significant improvement in services, with a significant increase in IBD specialist nurses, more multidisciplinary team working and enhanced service quality throughout the UK.

### What are the benefits to you of being an ECCO Country Member?

The UK has found enormous value in ECCO (reflected in the high UK membership), with opportunities for benchmarking IBD clinical practice across Europe, high-quality education and collaborative research.

### Is your society making use of the ECCO Guidelines?

ECCO Guidelines are widely used and referred to in the UK, although the BSG continues to develop UK IBD Guidelines (due in early 2017).

# Have you developed research projects with other countries through your ECCO Country Membership?

Yes: the SPARE/BIOCYCLE study, the I-CARE study and the ASTIC study of autologous stem cell transplantation.

# Have you developed educational activities with other countries through your ECCO Country Membership?

We have hosted a recent ECCO Regional Workshop in Edinburgh and hope to do so again in 2017.

*Has your country been involved in a fellow exchange through ECCO?* Several visiting fellows have been funded through ECCO since 2012.

### What are your main areas of research interest?

There are many and varied research interests in the UK, from basic science to microbiome studies, clinical trials and particularly genetics studies through the UK IBD genetics consortium.

# Does your centre or country have a common IBD database or bio bank?

As well as the UK IBD genetics consortium, the MRC IBD Bioresource has been established. The National IBD Registry has been launched as a UK database to allow real-time clinical data entry – succeeding the National IBD Audit as a means of data collection for audit and quality improvement across the UK.

# What are your most prestigious/interesting past and ongoing projects?

Collaborative genetics studies (past and current) are of great importance. Current projects include studies of pharmacogenetics of serious drug side-effects (PRED4), studies of monogenic causes of early IBD and twin/ multiplex registries. Amongst many other studies, it is worth highlighting the previous studies from Cambridge of gene expression profiling in CD8+ cells to predict prognosis in UC and CD, with a new CD8 biomarker stratified trial of early aggressive therapy in CD starting now. The TOPPIC trial, completed recently, was the largest study of postoperative therapy with mercaptopurine. The PANTS study is ongoing, collecting reallife data and biomarkers of anti-TNF response in CD. Many UK sites are also participating in I-CARE, a European prospective longitudinal observational multicentre cohort study of anti-TNF therapy.





Tariq Ahmad © Tariq Ahmad

Barney Hawthorne © Barney Hawthorne

*Which ECCO Projects/Activities is the group currently involved in?* See above.

### What are your aims for the future?

The BSG is committed to improving the quality of IBD care in the UK by promoting ongoing collaborative research through the IBD CRG and expanding the use of the IBD registry with a view to continuing national audit and quality improvement.

### How do you see ECCO helping you to fulfil these aims?

Despite the disappointing outcome of the UK referendum on EU membership, the UK IBD community remains committed to developing ever closer clinical, research and organisational links with ECCO. Future success for our IBD patients depends on bigger collaborations to generate and test ideas, and to develop better ways of treating our patients.

# What do you use ECCO for? Network? Congress? How do you use the things/services that ECCO has to offer?

ECCO gives us the opportunity to network with old and new European friends, as well as to engage in cutting-edge education, collaborative research and audit.

BARNEY HAWTHORNE TARIQ AHMAD ECCO National Representative, United Kingdom

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# Inflammatory Bowel Diseases



# 12<sup>th</sup> Congress of ECCO February 15-18, 2017

- CCIB Barcelona, Spain
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