

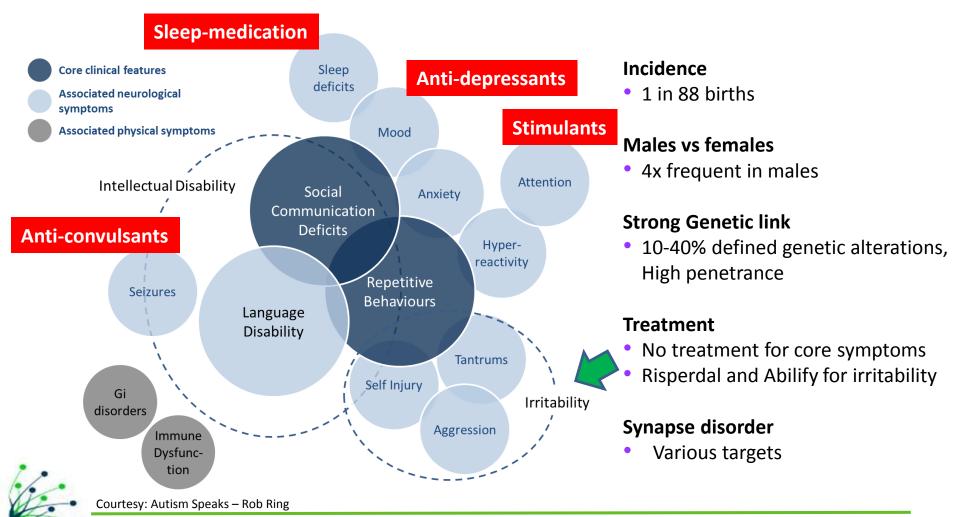


European Autism Interventions - A Multicentre Study for Developing New Medications (EU-AIMS)

Will Spooren and Declan Murphy
F. Hoffmann-La Roche/IoP, King's College

Autism spectrum disorders (ASD): Current situation







Autism spectrum disorders (ASD): European Situation 2012



- No major strategy defined within Europe
- No major or concerted efforts in drug discovery
- No pre-clinical network
- No clinical trial network
- No translational network
- No regulatory strategy
- Late diagnosis and poor awareness (adults)
- Poor knowledge of patients needs across life-course (teens into adulthood)
- Wide range in treatment strategy with no evidence of efficacy

a concerted effort of key stakeholders is needed Private Public Partnership





Neurodevelopmental disorders: from sin to synapse

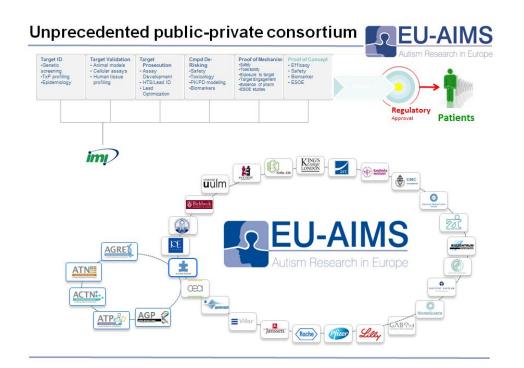


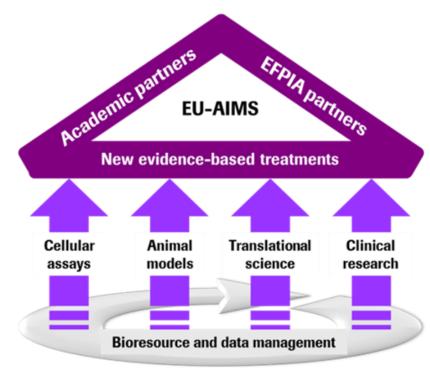
Synaptic dysfunction: development, maturation and stabilization of synapses. Clinical outcomes and diagnostic BM Genetics of mendelia forms Mechanistic therapeutics Fragile X syndrome Rett syndrome Molecular pathophysiology Translational models and target id.



EU-AIMS







Launched April 2012







12 publications in 1st year of the project

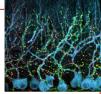
DOWJONES

WALL STREET JOURNAL

Animal Study Offers Prospect Of Autism Treatment

Roche, in collaboration with Seaside Therapeutics, is testing treatments for autism spectrum disorders, targeting the mGlu receptors.





The Emerging Biology of Autism Spectrum Disorders

"Synaptic connections in the brain of an autistic mouse"

Roche: New Findings From A Preclinical Study Of Autism

"The pharmaceutical company Roche along with the Biozentrum has discovered new insights to the study of autism."

New Scientific Research Attacks Behaviors In Autism

"According to Swiss drug maker, Roche Holding, Changes in the brain caused by autism can be reversed in mice, a new preclinical study showed, opening a potential path to develop a treatment for the incurab



Including toplines journals:

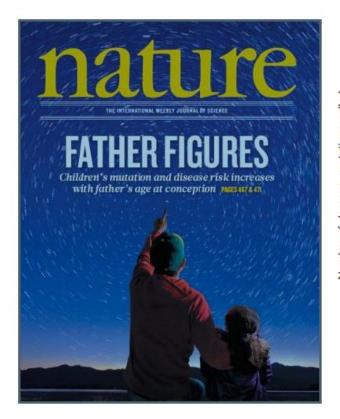
Nature, Nature – Drug Discovery, Science, PNAS, Molecular Psychiatry, JAMA, TiPS

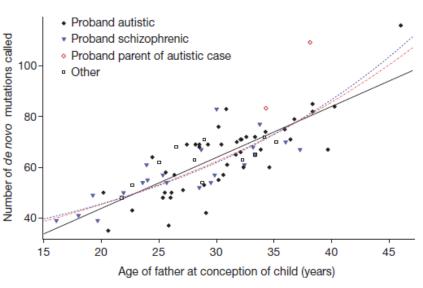
Mean Impact factor: 13.8











Age father is a risk factor for child to have deleterious mutation leading to autism



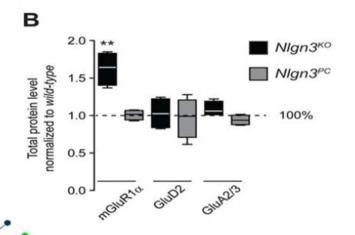


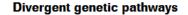


Shared Synaptic Pathophysiology in Syndromic and Nonsyndromic Rodent Models of Autism

Stéphane J. Baudouin, ¹ Julien Gaudias, ¹ Stefan Gerharz, ^{1*} Laetitia Hatstatt, ¹ Kuikui Zhou, ² Pradeep Punnakkal, ¹ Kenji F. Tanaka, ^{3,4} Will Spooren, ⁵ Rene Hen, ³ Chris I. De Zeeuw, ^{2,6} Kaspar Vogt, ¹ Peter Scheiffele¹t







ingle variants/gene(s) Comorbid Phenotype ~10% Neuroligin 3

 Fragile-X · Rett Syndrome

Tuberous Sclerosis

Chromosome anomalies ~ 10%

> Trisomy Chr. 21 · Chromosome 15

Chromosome 22

CNV/gene(s) ~ 65-75%

· Chr 2q, 5p, 7q Chr 9p, 11p, 15q

Chr. 17q, 19q, 22q

Group I mglu receptors 1

Convergent pathway







· SHANK 3





- 1 21 March 2013
- 2 EMA/CHMP/40896/2013
- 3 Committee for Medicinal Products for Human Use (CHMP)
- 4 Concept paper on the development of Medicinal
- 5 products for the treatment of Autism Spectrum
- 6 Disorder

7

Agreed by CNS Working Party	February 2013
Adopted by CHMP for release for consultation	21 March 2013
Start of public consultation	4 April 2013
End of consultation (deadline for comments)	4 July 2013

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Comments should be provided using this <u>template</u>. The completed comments form should be sent to CNSWPSecretariat@ema.europa.eu

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Keywords	Autism, Paediatric population, Asperger's Disorder, Rett's disorder,	
	Childhood disintegrative disorder and Pervasive Developmental	
	Disorder - not otherwise specified	

Regulatory guidance document

initiated in

March 2013

Concept paper expected in 2014







WP1 Cellular Assays

- Robust methods to reprogram keratynocytes into iPSc from autism patients and controls
- Shank3 iPSc and control lines delivered to EU-AIMS partners

WP2 Animal models

- Central Animal (TG) repository established (Harlan, Italy)
- A multi-site study for standardization and cross-site comparison completed (Oxytocin)
 - First results indicate excellent correlation between sites across Europe
- Genetic rescue of NRL3 KO phenotype
- $-\hspace{0.1cm}$ New target identified for the treatment of autism (mGlu1 lackbreaklar)
 - Pharmacological intervention studies initiated
 - Findings are being translated to man including potential PET study (new being nogatiated)
- Agreement with Sage labs for (50%!) discount for TG autism rats
- First TG rats at various behaviour groups of EU-AIMS testing initiated







WP3 Translational Sciences

- New data: evidence that brain functional abnormalities can be reversed and in adults with ASD by modulating brain 5HT. Currently tested as an outcome predictor for clinical trials.
- New data: ASD patients have significant differences in GABAalpha5 binding.

WP4 Clinical Sciences

- New data: individuals with ASD have significant differences in cortico-cortical anatomical connectivity.
 This is now being tested as an outcome predictor for clinical trials.
- New data: infants at risk for autism have significant differences in brain functional response to emotional sounds. This is now being investigated as a risk predictor
- New data: Published first study of EU wide prescribing in ASD demonstrated; 1) very low prescription rates for associated symptoms, and 2) very wide variation across countries (NL highest).
- Regulatory succes: Worked with EMA to launch concept paper for regulatory guidance (2014).

WP5 Biorepository and data base

- Biorepository ready to receive samples
- Data base nearly functional







What next?





Identify biomarkers of ASD which precede onset of clinical symptoms – High-risk sibling study



- OBJECTIVE: To investigate patterns of brain development that may be associated with early detection of ASD
- white matter development
- infant brain processing of human voice sounds (+ emotions).





- Participants: n=300 HR infants, n = 100 LR infants
- infants between 3 and 7 months





Validate biomarkers of ASD in children and adults

Accelerated Longitudinal Study



Large scale multi site clinical study

Total 480/450 **ASD**, 320/330 **HC**)

Four schemes:

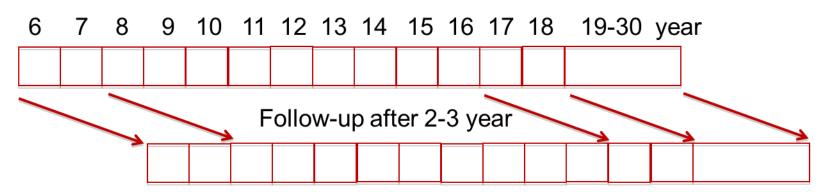
A: HFA adolescents/adults (100 + 100)

B: HFA children (100)

C: LFA adolescents / adults (100)

D: MZ and DZ twins (50 + 30)

Follow-up 18 months



knowledge of patients needs across life-course

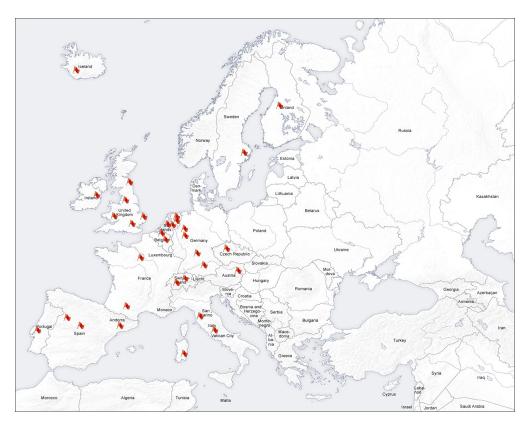




Build clinical trial network



- Contacted existing partners
 European networks; ECNP, ESSEA
 COST Action, & other major clinical
 research centres across Europe
- **48** invitation letters sent to **41** sites
- Response
 - 34 sites from 15 countries showed willingness (83% of sites contacted)
- Second wave of letters focus on Eastern Europe and scandinavia





May indicate multiple sites in the same city

EU-AIMS is now being contacted by centers that want to participate!





Develop inventory and data mining of clinical databases



Develop European inventory of ASD patients & measures

- Potential sites will complete a brief online survey in early summer 2013
- Focus on patient numbers, characteristics, & routine assessments
- Help data compatibility

Coordinate large clinical databases

- Collect historical anonymised data from clinical research network
- Analyse datasets to answer informative questions
- Help identify potential outcome measures





Remaining challenges



- Heterogeneity in patients
 - Stratification and relevant biomarker
- Evidence based therapy
- Clinical endpoints
- General awareness of ASD across Europe





EU-AIMS – Full details





EU-AIMS: a boost to autism research

Will Spooren, Declan Murphy

Nature Reviews Drug Discovery 11, 815-816 (November 2012)







Thank you



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Major goals for Europe





Development and validation of translational approaches for the advancement of novel therapies to treat ASD



Setting new standards in research and clinical development to aid the drug discovery process



Identification and development of expert clinical sites across Europe to run clinical studies and trials, and the creation of an interactive platform for ASD professionals and patients.



