CRITICAL REVIEW AND INVITED COMMENTARY

Advancing research toward faster diagnosis, better treatment, and end of stigma in epilepsy

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Abstract
Seven large European Union (EU)-funded epilepsy-related research projects joined forces in May 2018 in Brussels, Belgium, in a unique community building event—the epiXchange conference. During this conference, 170 investigators from the projects DESIRE, EpimiRNA, EPISTOP, EpiTarget, EpiXchange, and EpiPGX as well as the European Reference Network EpiCARE, met up with key stakeholders including representatives of the European Commission, patient organizations, commercial partners, and other European and International groups. The epiXchange conference focused on sharing and reviewing the advances made by each project in the previous 5 years; describing the infrastructures generated; and discussing the innovations and commercial applications across five thematic areas: biomarkers, genetics, therapeutics, comorbidities, and biobanks and resources. These projects have, in fact, generated major breakthroughs including the discovery of biofluid-based molecules for diagnosis, elucidating new genetic causes of epilepsy, creating advanced new models of epilepsy, and the pre-clinical development of novel compounds. Workshop-style discussions focused on how to overcome scientific and clinical challenges for accelerating translation of research outcomes and how to increase synergies between the projects and stakeholders at a European level. The resulting advances would lead toward a measurable impact of epilepsy research through better diagnostics, treatments, and quality-of-life for persons with epilepsy. In addition, epiXchange provided a unique forum for examining how the different projects could build momentum for future novel groundbreaking epilepsy research in Europe and beyond. This report includes the main recommendations that resulted from these discussions.

KEYWORDS
biobanks, biomarker, comorbidities, databases, delivery systems, disease-modification, e-health, epileptogenesis, genetics, personalized medicine, research strategy, stigma, therapy
1 | INTRODUCTION

Epilepsy affects over 6 million people in Europe, with an estimated annual health care and societal cost of 16 billion euros. This condition presents as an exceptionally multifaceted cluster of diseases, which vary in semiology, etiology, pathophysiology, progression, outcome, and comorbidities.\(^2\) The epilepsy research community, in partnership with industry and other stakeholders, has had many successes, including the development and approval of over 20 different antiseizure drugs, which provide seizure control for two-thirds of patients.\(^3\) However, none of these drugs prevent the development, or alter the course, of epilepsy, and 30% of patients remain refractory to currently available treatment—a figure that has not changed over the last decades.\(^2,4\) In addition, more than 30% of treated patients experience adverse events that compromise their quality of life. In addition, they have concurrent cognitive, psychiatric, and other comorbidities, as well as stigma, which is still widely related to this condition.\(^5,6\) It often takes many years before a patient receives a correct diagnosis or the appropriate medication.\(^7\) Research to elucidate molecular mechanisms of epilepsy is now reaching the clinic, with gene panels and exome/genome sequencing guiding some treatment and care pathways.\(^8,9\) However, the majority of patients still do not receive a genetic diagnosis. Another key unmet need is that there has been a two- to threefold increase in mortality of persons with epilepsy, with sudden unexpected death in epilepsy (SUDEP) being the major cause of premature loss of life in affected persons.\(^10\)

Together, these challenges mandate transdisciplinary expertise and coordinated actions in brain research that integrate other scientific disciplines and emerging technologies, in order to advance our understanding of the cause(s) and treatment of epilepsy. Concurrently, unique features of epilepsy as a medical condition provide invaluable opportunities for fundamental studies of brain network function, improved imaging and diagnostics, and the search for disease mechanisms and novel drug targets. This could also lead to new therapies for other conditions such as neuropsychiatric disorders,\(^11\) neurodegenerative diseases\(^12\) and autism.\(^13\) Consequently, research into the treatment of epilepsy also provides a unique insight over the broader neuroscience field.

2 | INVESTMENT IN EPILEPSY RESEARCH UNDER FRAMEWORK PROGRAMME 7

There has been historical underinvestment in epilepsy research relative to its disease burden.\(^14\) Compounding this, various manufacturers of antiseizure drugs have recently disengaged from research and development in the area of epilepsy therapeutics.\(^15\) Tackling these challenges received a critical boost in 2013, when, after determined and successful efforts to put epilepsy on the EU research agenda, a specific call was included in the last year of Framework Programme (FP) 7, inviting large collaborative research projects to address the challenges posed by epilepsy and to advance our understanding of mechanisms, diagnosis, and treatment. Projects had to be multidisciplinary in nature and feature collaborative work by scientists and clinicians as well as other technical experts from outside the field. Finally, each project required a significant involvement of small- and medium-sized enterprises (SMEs). Four projects received funding—a total investment of approximately 50 million €—each with unique, as well as complementary, research approaches blending basic and translational science, animal models, and clinical studies and trials. The projects were called DESIRE, EpimiRNA, EpiSTOP, and EpiTarget. In addition, a number of other European epilepsy research projects overlapped within this same timeframe, including EpiPGX and Epixchange, which were funded by earlier funding rounds in FP7, and EpiCARE, a European Reference Network approved and funded by CHAFEA, the EU Consumers, Health, Agriculture and Food Executive Agency, to enhance and better network clinical care and research in epilepsy.

3 | EPIXCHANGE 2018—A UNIQUE COMMUNITY-BUILDING EVENT FOR EPILEPSY RESEARCH IN EUROPE

Soon after the projects were launched, discussions began between the project coordinators to bring together these
projects when results began to emerge. Although the epilepsy research communities in Europe and across the world meet regularly through events such as the European, International, and American epilepsy congresses, the project coordinators recognized that an event bringing together these specific projects would be unique, focusing on the achievements, the discoveries made, and the breakthroughs still to come. It would also provide a forum to identify remaining roadblocks and challenges to progress and discuss ways to secure research funding through future European and other programs.

The epiXchange conference was held on May 23rd 2018 in Brussels, Belgium. Altogether 170 investigators from 18 European countries and 5 non-EU countries met with several stakeholders, representatives of the European Commission, patient organizations, nonprofit research institutes, and other key European and international groups, to present the results from their ongoing research. The conference was organized into a series of thematic sessions, chaired by clinicians and scientists from the different projects. The thematic areas were based on shared research priorities that were specifically addressed by the EU call: biomarkers; genetics; therapeutics; comorbidities; and databases and biobanks.

The epiXchange meeting clearly captured the scale of progress made by the different consortia. This included traditional scientific outputs such as new knowledge about the causes and treatment of epilepsy, novel genetic causes of epilepsy, diagnostic biomarkers and precision medicine approaches, transformative new in vitro and in vivo models (eg, induced pluripotent stem cells and use of model organisms such as zebrafish), structural and functional brain imaging, and new therapeutic targets moving into preclinical development. All these results and outputs were published (or in submission) in leading peer-reviewed journals (Table 1). The meeting also highlighted other important achievements including the infrastructure and expertise developed to date, refinements of molecular and other biomedical research technologies, establishing best research practices, expanding bioresources such as DNA and tissue banks, creating new databases that can transform the power of pathway discovery, and significantly for future progress, the development of new intersectorial partnerships. There was specific attention paid to how these projects have helped support, and have been supported by, small- and medium-sized enterprises (SMEs) in Europe, a key impact metric for the FP7 projects. Each project clearly benefited from this innovative research approach and recognized the key role of industry in accelerating translation of basic science toward actual life-changing patient care and the resulting socioeconomic contribution to Europe. A number of the industry-based researchers also presented at the meeting.

Finally, the meeting provided a forum to discuss the way forward for accelerating epilepsy research in Europe and beyond. EpiXchange was both a reflection on what was achieved and what was foreseen emerging in the near future. It was (it is) a means for reinforcing the existing collaborations between projects and creating new ones that will lay the foundations for success in delivering transformative research with direct impact for the citizens of Europe with epilepsy. More information about these various research projects, presentations, and videos from the conference can be found at www.epixchange2018.eu.

4 | OVERVIEW OF PROJECTS AND SUMMARY OF PROJECTS’ ACHIEVEMENTS

Through oral presentations and posters, project coordinators and presenters at epiXchange summarized key achievements. The main highlights of these achievements are included in Table 1. Details of each project can be found at their various hosting websites.

The DESIRE project (http://www.epilepsydesireproject.eu/) focused on uncovering developmental brain processes to understand the genetic mechanisms underlying epileptogenic developmental disorders (ie, early onset epilepsies in children). The research in DESIRE comprised work packages focused on discovering mutations in such patients and understanding the mechanisms of disease. This research has added value through the development of novel diagnostics and treatment strategies where use of gene therapy and reprogramming of somatic cells into neurons was also being pursued, along with new recording and analysis tools for diagnosing epileptiform activity in patients. Another function of the project has been to continue to expand the collection of pathologic brain specimens in the European Brain Bank. The project has also resulted in the launch of a multicenter longitudinal study of patients with Dravet syndrome (DS).

The EpimiRNA project (https://www.epimirna.eu/) focused on molecules called microRNAs (miRNAs), which serve important functions in controlling gene expression. EpimiRNA’s research has focused on cataloguing which miRNAs are active in animal models of epilepsy and in brain tissue from adult patients with epilepsy. In addition, the project focused on identifying miRNAs in biofluids, such as diagnostic biomarkers, on solving the mechanisms by which miRNAs affect brain excitability, the use of systems biology to predict miRNA-target interactions, looking for possible genetic variation in miRNAs or their targets in epilepsy patients, and exploring the potential of miRNA-based treatments. The project also included clinical studies on brain stimulation and recording technologies.

EPISTOP (http://www.epistop.eu/) was the first human study aimed at prospectively identifying biomarkers for epileptogenesis, beginning from its latent phase, through active epilepsy and chronic disease (tuberous sclerosis complex...
### TABLE 1 Summary of major achievements reported across the thematic areas by the FP7-funded large-scale collaborative research projects on epilepsy at epiXchange

<table>
<thead>
<tr>
<th>Thematic area and project output reported at meeting</th>
<th>Related publications</th>
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<tbody>
<tr>
<td><strong>Translational research in biomarkers</strong></td>
<td></td>
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<tr>
<td><strong>DESIRE</strong> reported that:</td>
<td>PMID: 27694961</td>
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<tr>
<td>Epigenetic (DNA methylation) marks are biomarkers used to classify different types of focal cortical dysplasia. A longitudinal study of 330 patients with Dravet syndrome (DS) suggests early diagnosis and appropriate intervention may benefit prognosis and has also clarified to what extent early clinical and genetic findings predict outcome.</td>
<td>PMID: 27864847</td>
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<tr>
<td>A system for high-performance recording and analysis of brain signals that may predict surgical outcomes has been awarded a European conformity mark.</td>
<td>PMID: 2652080</td>
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<tr>
<td>It also highlighted the diagnostic yield from exome sequencing and novel imaging modalities, including ultra-high MRI field, for diagnosis.</td>
<td>PMID: 29668857</td>
</tr>
<tr>
<td>Implications for diagnostic strategies and genetic counseling in relation to germline and mosaic mutations have been elucidated for several mTOR-related disorders.</td>
<td>PMID: 28202706</td>
</tr>
<tr>
<td><strong>EpimiRNA</strong> identified:</td>
<td>PMID: 29069555</td>
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<tr>
<td>Sets of microRNAs in blood and cerebrospinal fluid that support a diagnosis of epilepsy or status epilepticus and companion technology for their rapid detection in a point-of-care setting</td>
<td>PMID: 26778405</td>
</tr>
<tr>
<td><strong>EpiSTOP</strong> identified:</td>
<td>PMID: 27834078</td>
</tr>
<tr>
<td>EEG changes that are a biomarker of emerging epilepsy in infants and children with TSC.</td>
<td>PMID: 25258368</td>
</tr>
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<td><strong>EpiTarget</strong> demonstrated that:</td>
<td>PMID: 24763967</td>
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<tr>
<td>In animal models and human epilepsy, serum HMGB1 is a prognostic and mechanistic biomarker, predicting with high fidelity the onset of epilepsy in animal models and pharmacoresistance in patients.</td>
<td>PMID: 27042956</td>
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<tr>
<td>Preclinical MRI and MR spectroscopy have also been used to identify novel epilepsy biomarkers.</td>
<td>PMID: 31074842</td>
</tr>
<tr>
<td>It also established common data elements (CDEs) and case report forms (CRFs) for harmonization of preclinical data collection and analysis in epilepsy research.</td>
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<tr>
<td><strong>EpiPGX</strong> identified:</td>
<td>PMID: 28607431</td>
</tr>
<tr>
<td>Genetic risk factor for rash associated with antiepileptic drug treatment, and treatment options for a common treatment-resistant epilepsy.</td>
<td>PMID: 28496112</td>
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<td>It also provided the first evidence that multidrug resistance has a genetic contribution (heritability) - unpublished data.</td>
<td>PMID: 26699132</td>
</tr>
<tr>
<td>Biomarkers for hyponatremia have been evaluated; and a genetic biomarker study is in press (no PMID yet).</td>
<td>PMID: 30396857</td>
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<tr>
<td><strong>Genetics of epilepsy—therapeutic implications</strong></td>
<td></td>
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<tr>
<td><strong>DESIRE</strong> identified:</td>
<td>PMID: 28053010</td>
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<tr>
<td>Several new genetic causes of developmental and epileptic encephalopathies and elucidated the mechanisms by which these may drive brain malformations. This includes AKT3, GABRB3, SPTAN1, and ATP6V1A.</td>
<td>PMID: 26520804</td>
</tr>
<tr>
<td>It identified a high frequency of mosaic mutations including in the regulatory subunit of the PI3K-AKT-MTOR pathway, PIK3R2, as a cause of the epileptogenic brain malformation, perisylvian polymicrogyria.</td>
<td>PMID: 29668857</td>
</tr>
<tr>
<td>It also found how mutations in the NEDD4L gene lead to abnormal neuronal positioning (gray matter heterotopia) and the underlying pathogenic mechanism through mTORC1, AKT, and Smad2/3 activities.</td>
<td>PMID: 29050398</td>
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<tr>
<td><strong>EpimiRNA</strong> reported:</td>
<td>PMID: 29286390</td>
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<tr>
<td>The first analysis of genetic variants in microRNA biogenesis genes, microRNAs, and ~15,000 targets in over 3500 patients and controls.</td>
<td>PMID: 25552301</td>
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<tr>
<td>The discovery of multiple new microRNAs regulating brain excitability and their targets/mechanisms in animal models and human epilepsy.</td>
<td>PMID: 26631939</td>
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<tr>
<td><strong>EpiSTOP</strong> reported:</td>
<td>PMID: 29509898</td>
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<tr>
<td>Several discoveries on gene mutations and the molecular mechanisms of TSC-related epilepsy</td>
<td>PMID: 28487411</td>
</tr>
<tr>
<td>It identified strong causal associations between TSC and aberrant expression of microRNAs.</td>
<td>PMID: 27840225</td>
</tr>
<tr>
<td><strong>EpiTarget</strong> showed:</td>
<td>PMID: 27681419</td>
</tr>
<tr>
<td>Functional epigenomic influences in the human and corresponding mouse models</td>
<td>PMID: 29196317</td>
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(Continues)
### Thematic area and project output reported at meeting

#### EpiPGX contributed:
- To discoveries in genetic generalized epilepsies, and to the largest ever GWAS in epilepsy (in press, no PMID yet).

#### Innovative therapeutics and translation

**DESIRE developed:**
- A number of therapeutic strategies for epileptic encephalopathies. This includes use of an optogenetic technique for control of transcription factor REST and genetic techniques for direct conversion of fibroblasts into inhibitory neurons.
- The effectiveness of established and new third-generation antiepileptic medication approaches has been critically reviewed.

**EpimiRNA discovered:**
- A potent disease-modifying (antiepileptogenic) microRNA-targeting oligonucleotide and advanced it into preclinical development (patent awarded, US 9 803 200 B2).
- Novel delivery routes have been validated for microRNA-based drug delivery.
- New small molecule identified with antiseizure effects in preclinical development with a project SME.
- A new combined stereo-EEG electrode for simultaneous recording of deep brain activity, microdialysis, and delivery of therapeutics is in preclinical testing.

**EpiSTOP reported:**
- Updates on a prospective clinical study of TSC infants treated with antiepileptic drugs at the onset of subclinical seizures in comparison to children treated only after clinical seizures appear. Results suggest the early intervention is disease-modifying.
- Preclinical studies have demonstrated gene therapy using hamartin can ameliorate behavioral phenotypes in a mouse model of TSC.

**EpiTarget demonstrated:**
- Combinatorial approaches using antiinflammatory, antioxidant, and microRNA combinations produce significant and potent disease-modifying effects in animal models of epilepsy.
- Gene therapy tools and Molecular Envelope Technology (MET) were developed to allow short- and long-term delivery of genes targeting distinct mechanisms of epileptogenesis.
- rAAV-mediated expression of a dominant-negative MTF1 was found to abolish SE-induced CaV3.2 mRNA upregulation and attenuates epileptogenesis.

**EpiPGX identified:**
- Mechanisms associated with cortical thinning and disease severity in the epilepsies, with proof of concept of preventative therapy (in collaboration with EpiTarget): unpublished data

**EpiXchange developed:**
- New gene therapy and cell therapy tools for the transfer of large therapeutic molecules, like neuropeptides or neurotrophic factors, directly in the diseased brain tissue.
- Human brain slices obtained from surgery of epilepsy patients were used to select the most effective treatments and thereby address the translational potential of therapies developed in animal models.

### Understanding comorbidities in the epilepsies

**DESIRE has:**
- Paid particular attention in measuring to what extent the genetic alterations and structural abnormalities cause intellectual disability and behavioral problems per se, and what is the specific weight of the superimposed intractable epilepsy in relation to its severity and age at onset. Dravet syndrome and epilepsy with continuous spikes and waves during slow sleep have been chosen as models.
- A European registry for Dravet syndrome has been launched as the basis for a long-term perspective study assessing associated comorbidities

**EpiSTOP reported:**
- Early deviation in developmental trajectories in relation to seizure onset in TSC. It was shown that some clinical biomarkers may predict autism onset already at 6-12 months, enabling early treatment strategies.
In the project, risk factors and biomarkers of epilepsy development were identified in newborns and infants with TSC. A key part of the study was determining whether early preventive treatment can block or delay the onset of epilepsy and comorbidities such as cognitive and language developmental problems. Research within the project also included investigations into the cell and molecular mechanisms by which genetic mutations cause TSC.

**EpiTarget** (https://www.epitarget.eu/) focused on identifying biomarkers and multiple basic mechanisms for epileptogenesis in adults, and translating these findings to the clinic by validating the biomarkers in human samples. The project focused on testing combinatorial therapeutic approaches such as combining antioxidant or antiinflammatory treatments in preclinical models. Common data elements (CDEs) were devised to enable harmonization of procedures between laboratories in preclinical trials. The biomarker research included identification of proteins and RNAs in biofluids as well as using imaging techniques. Finally, the project explored gene therapy approaches and advanced drug-delivery tools.

In addition to the four main projects, epiXchange featured presentations from three other projects. The FP7 Thematic area and project output reported at meeting

<table>
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</table>
| **EpiTarget** demonstrated: Systems genetic evidence for a convergence of epilepsy and its comorbidities on shared molecular pathways. The bioinformatic characterization of potential novel therapeutic targets predicted the tyrosine kinase receptor Csf1R and was validated in complementary mouse model and human epileptic hippocampi. | PMID: 27955713  
PMID: 30177815 |
| **EpiPGX** used: GWAS and polygenic risk score methods, and demonstrated significant genetic correlations between multidrug resistance and neurodegenerative processes (unpublished data) | PMID: 29766029 |
| **EpiXchange** demonstrated that: Certain neurotrophic factors, when released continuously in the hippocampus of chronically epileptic rats through an encapsulated cell biodelivery system, strongly attenuate epileptic seizures and comorbidities. | PMID: 29069555 |
| **Biobanks and databases—basis for translational research**

**DESIRE** reported on:
The European Epilepsy Brain Bank, which recently collected findings on nearly 10,000 epilepsy patient resections from 36 epilepsy centers across 12 European countries. This resource supports diagnosis and future research on genetic markers for disease classification and personalized treatment. | PMID: 26748106  
PMID: 27839653  
PMID: 28331471  
PMID: 30525113 |
| **EpimRNA** established: A clinical biobank containing DNA, tissue and biofluids from over 850 TLE patients and another 1000 epilepsies and over 1700 controls. Preclinical biobank comprising tissue and biofluids from three different animal models of epilepsy. It contributed to the European epilepsy brain bank consortium. It set up a database for tracking microRNA - epilepsy associations for the scientific community (www.epimirbase.eu). This includes a molecular brain tissue atlas that features data on levels of all active miRNAs in three models of epilepsy and a database containing all human microRNAs and their targets in human TLE brain. Systematic reviews on microRNAs as biomarkers and therapeutic targets. | PMID:29338461  
PMID: 27042238  
PMID: 27750396  
PMID: 27425893  
PMID:28808237  
PMID: 27014996  
PMID: 29384235 |
| **Epistop** reported on: Biobanking in tuberous sclerosis complex (TSC) and challenges and opportunities for understanding epilepsy and cognitive and behavioral comorbidities. TSC represents a model disease for investigating mTOR-related epileptogenesis. TSC molecular networks highlighting evidence for shared cellular and molecular mechanisms underlying severe pediatric epilepsies. | PMID:29291240  
PMID: 27912503  
PMID: 28912503  
PMID: 27750396  
PMID: 27425893  
PMID:28808237  
PMID: 27014996  
PMID: 29384235 |
| **EpiTarget** established: An Epilepsy Preclinical Biomarker Bank (EPBB), including imaging and EEG data, and blood samples from various animal models of epilepsy, using REDCap database for biobanking information. It also performed meta-analyses of microRNA dysregulation across animal models and humans to identify conserved microRNAs as biomarkers and therapeutic targets. | PMID: 29338461  
PMID: 28912503  
PMID: https://doi.org/10.1002/epi4.12275 |
EpiPGX project (https://www.epipgx.eu/) aimed to identify genome-based biomarkers for use in clinical practice to personalize treatment of epilepsy, by stratifying patients for clinical trials, preventing relapse, and reducing adverse drug reactions. EpiXchange (www.epixchange.eu) was an FP7-IIAPP that explored innovative cell and gene therapies for the treatment of partial epilepsies. EpiCARE (https://epi-care.eu/) is a DG Sanco European Reference Network focusing on rare and complex epilepsies and aims to develop and deliver highly specialized diagnostics and clinical care in order to improve interventions and outcome in individuals with rare and complex epilepsies initiated in 2017. EpiCARE is an extension of a previously DG Sanco-funded project, E-PILEPSY (2014-2016), a pilot network of cooperation in epilepsy surgery and refractory epilepsy that continues as the surgical therapeutic arm of EpiCARE.

Altogether, the presentations and additional achievements of the various projects clearly indicated the enormous advances in our understanding of the pathogenesis, diagnosis, and treatment of epilepsy. Collectively, the projects have led to dozens of new discoveries about the cellular and molecular mechanisms of epilepsy and have driven the preclinical development of dozens of potential new therapies. A wide range of diagnostic approaches are, in fact, now at various stages of validation and entering clinical testing. The impacts from these projects can thus extend beyond the scientific advances and the new technologies to patient care, including improved diagnosis, better understanding of side effects, and new community resources fostering increased competitiveness of SMEs in Europe. However, to ensure that these achievements are of concrete benefit and reach persons with epilepsy, further research and innovation efforts are still required by the community of researchers, clinicians, and industry. This collaborative effort should also involve patients, as described in the next section.

5 | EPIXCHANGE: RECOMMENDATIONS FOR THE FUTURE

In addition to the presentations of the achievements of the various projects to date, the epiXchange meeting provided a new forum for a discussion about the future of epilepsy research. A series of recommendations for future research emerged, aimed at ensuring a broadening of the knowledge base and achieving the necessary impact to improve the lives of persons with epilepsy or those at risk of developing epilepsy (Figure 1). Particular emphasis was also made on the engagement of persons with epilepsy and their caregivers in research and innovation actions.

5.1 | Higher throughput translational models for diagnostics and therapy target development

It is recognized that seizures are a symptom of epilepsy. Preclinical seizure models have a strong track record of delivering benefits for persons with epilepsy, having been instrumental in the discovery of many of the currently used, effective antiseizure drugs. Recognizing that these traditional models have led to somewhat blinkered approaches, however, it has become clear that we need different types of animal models. New, higher throughput models and refined “old” models will help to discover disease-modifying therapies and to de-risk failure in translation. Thus ongoing research is resulting in the identification of several genetic and acquired in vitro and in vivo animal models of human epilepsies, leading to a better understanding of the mechanisms of development of the various epilepsies and types of seizure generation. To make faster progress and increase the translational value of preclinical research, epiXchange investigators agreed that there is an urgent need for:

- The development of new, high throughput, translational in vitro, in vivo, and in silico models through the application of innovative technologies including advanced optogenetics and chemogenetics, genome modifications (CRISPR-based), human induced pluripotent stem cells, and human brain organoids.
- The refinement of the existing epilepsy models and the development of innovative preclinical and clinical study designs. Closer matching of models to human clinical phenotypes may help ensure more direct and immediate clinical translatability.
- Further and expanded efforts to discover new targets through hypothesis-driven and unbiased “omic” approaches using animal and human models, in order to continue to elucidate the mechanisms of epileptogenesis.

These improvements in modeling will address the urgent need for an evidence-based disease classification, integrating clinical, pathological, and genetic findings in order to reduce time to diagnosis and personalized treatment.

5.2 | Target-led diagnostics discovery

The rate of misdiagnosis of epilepsy remains high. This does not relate only to diagnosis of epilepsy itself, but also to the recognition of the epilepsy type and etiology, processes that may take several years. Improved diagnostic techniques may be expensive, but a more accurate diagnosis would result in prompt application of correct treatments and, therefore, would be cost effective in the long term. Long delays in accurate diagnosis could be related to a poor knowledge
of epilepsy among health care professionals, lack of accurate etiology-based diagnosis, and incomplete understanding of disease mechanisms in a patient with a given epilepsy diagnosis. Thus, target-led diagnostics discoveries will lead to:

- An increase in the mechanism-led multimodal diagnostic and prognostic tools, including genetic, epigenetic, and proteomic tests for antiepileptogenesis and epilepsy, possibly at bedside.
- Identifying readily accessible and easy to evaluate and interpret biomarkers for diagnostics, prognostics, advancement, and discovery of new and personally adapted treatment regimens.
- A better understanding of the mechanisms of SUDEP (sudden unexplained death in epilepsy) and the development of innovative methods for its prevention.

These activities would result in a better overall stratification of people enrolling in preclinical and clinical trials. It would thus subsequently lead to greater predictive power of possible adverse events, comorbidities (such as attention-deficit/hyperactivity disorder or depression), and treatment efficacy toward (faster) individualized patient-centered care.

5.3 | Digitalization and new wearable materials in epilepsy diagnostics and monitoring

We still have limited understanding of the impact of everyday events and circumstances on seizure control in the epilepsies. Environmental and other influences undoubtedly exist, but are difficult to determine and measure, since digitization of this information on a widespread scale is still missing. The development of digital datasets offers the possibility of progress in this key area through the:

- Development of digital information and communications technologies in health care such as e-health and m-health technologies for personalized diagnosis of epileptogenesis, as well as monitoring of progression and treatment of epilepsy.
- Development of innovative virtual storage solutions such as Mycloud, possibly with other neurologic diseases.

These would lead to a better understanding of the effect of environmental, lifestyle, and other factors on epilepsy, and establish a Big Data discovery platform to improve the treatment and quality-of-life of persons with epilepsy and of their caregivers.

5.4 | Personalized medicines and delivery systems

Despite the introduction of about 10 new antiseizure treatments over the past 15 years, the proportion of people with poorly controlled epilepsy has not decreased. This could be related to an incomplete understanding of disease mechanisms in a patient with a given epilepsy diagnosis. Therefore, there is an urgent need for:

- The development of evidence-based precision medicines, integrating clinical, imaging, electrophysiologic, pathologic, and genetic findings for antiepileptogenesis and epilepsy.
- The development and application of innovative drugs, technologies, and preclinical and clinical trial designs.
- The development of new technologies for targeted treatment delivery such as gene therapy, oligonucleotides for targeting miRNA, siRNA, and small vesicle nanocarriers for proteins or peptides.
- Developing personalized therapeutic monitoring across genders and all ages, through the collection of information from brain activity.
- The development of noninvasive applications for multimodal monitoring of the efficacy of treatments and epilepsy surgery.

This would lead to the development of novel personalized technologies for therapy and for monitoring the efficacy of therapy, drug delivery, and compliance.

5.5 | Epilepsy data ecosystem

The seven EU-funded projects contributing to epiXchange have each individually generated a number of unique data-banks, tissue biobanks, and databases. The lack of harmonization and standardization in data collection and storage across the projects makes integration of these “banks” a complex process, hindering Big Data analysis and access from these key datasets. The new data protection GDPR regulations ensure that each person is now the steward of his/her own data. Innovative IT solutions provided by various SMEs or existing EU-funded data platforms can be used to address these requirements and will implement GDPR requirements about when and how consent is obtained and what data are stored. Therefore, epiXchange is proposing:

- The integration of these preclinical and clinical genetic, tissue, electroencephalography (EEG), and neuroimaging banks into a virtual, controlled access European Epilepsy Data Ecosystem. This would lead to the development of ancillary informatics tools, Big Data analyses and mining, and mathematical modeling of large multimodal datasets. Standardization in this field will greatly facilitate the guiding of breakthrough prospective diagnostic and therapy development studies.
- To ensure that this European Epilepsy Data Ecosystem can subsequently contribute to larger global existing databases such as the European Open Science Cloud, BBMRI (the European research infrastructure for biobanking), Human Brain Project, and ELIXIR (an intergovernmental organization that brings together life science resources from
across Europe). This would lead to a complete clinical-to-network-to-cell-to-molecule phenotyping of the epilepsies, which takes into account life-style factors, demographics, genetics, and other complex data, and provides unprecedented opportunities for preclinical, clinical, technological, and societal research and innovation activities.

5.6  Comprehensive European and worldwide epilepsy care

In addition to addressing traditional scientific questions, the epiXchange meeting also recognized the opportunity to develop new infrastructural models that would lead to a comprehensive European and worldwide epilepsy care for all ages. There must be better access across all Europe and worldwide to diagnostic tools such as genetic testing, brain magnetic resonance imaging (MRI), positron emission tomography (PET), and magnetoencephalography (MEG). Although some of these issues are being addressed through the European Reference Network, EpiCARE, with an emphasis on utilization of virtual technology, this is an area that is still very much in development, due to a lack of structured resources. We envisage:

- The creation of Virtual Epilepsy Centers would greatly help to alleviate some of these demands, and also address cognitive and behavioral comorbidities often seen in persons with epilepsy.
- The fostering of data-driven regulatory requirements and the harmonization of health care regulations at a European level, providing secure patient management platforms and financial support for holistic care.

5.7  Integration to society and regulatory space

The enormous medical, economic, and social burden of epilepsy can only be addressed through the coordination and integration of research among private and public partnerships. There should be a multi-stakeholder dialogue for priority setting in epilepsy research in order to define the goals/aims/opportunities for a leverage-synergy-maximal impact. This would allow patient/family participation from the outset of the mission, and would ensure that the results really address the needs of the epilepsy community. This can be achieved through multiple actions:

- Persons with epilepsy should have a key role to play in the drug-discovery process, the development of new devices, the assessment of new neurosurgical techniques, and the planning of clinical trial designs. They also have a key role to play in the application of the research findings.
- The high incidence of epilepsy as a comorbidity in various brain diseases, such as traumatic brain injury, stroke, and Alzheimer disease, can address some of the crisis the pharmaceutical industry is presently facing, since this mission would be a unique opportunity to join forces across various research and patient communities in order to implement joint research initiatives.
- Healthcare providers and policymakers should also take part in the selection process of future research directions. They are key enablers, facilitating the provision of initiatives such as starter grants for SMEs and the funding of clinical trials. Health Insurance providers can be involved in the development of new models on the best and sustainable use of limited resources. This is an area that is still very much underexploited.
- Public educational programs on epilepsy for patients, different sectors of society, and for all stakeholders should be developed.
- Supporting actions to facilitate and maintain interactions between various research initiatives and between different stakeholders, eg, through the organization of joint events and workshops, ongoing educational activities for early stage researchers in epilepsy, and the development of collaborative short- and long-term research strategies, should be established, also with the contribution of patients.
- A global initiative for independent living should be undertaken in a worldwide effort to reduce stigma, using validated tools and indicators, which can thus provide clear outcome measures and address discrimination.

5.8  Engagement of actors in the cross-disciplinary epilepsy mission across multiple sectors of society

The development of innovative diagnostic tools and treatments, as described above, requires not only trained and experienced clinicians, but also experts in genetics, cell biology, chemistry, pharmacology, electrophysiology, imaging, mathematics, computer sciences, artificial intelligence, engineering, nanotechnologies, robotics, personalized drug design, and surgery. In addition to the regulatory implications of such research, the implementation of innovations furthermore mandates an entrepreneurial spirit and the involvement of (patent) lawyers, financial experts, and product designers. These various actors must align and synergize cross-disciplinary research at all levels of the innovation chain for the discovery of novel diagnostic tools and therapies. This is essential if new treatments to address unmet needs are to reach the market. There are also unique opportunities for industry to leverage targets and assets that were developed for other disease areas.

In addition, it is key to recall that epilepsy can start at any age and is often a life-long condition. It affects various aspects of life, from education to employment, creating problems not
only just for the person with epilepsy, but also for caregivers. These, and the wider lay community, need to be informed and educated about the condition in order to prevent misperceptions, discrimination, and stigma, and bring persons with epilepsy out of the shadows. These educational programs should be conjointly designed by various actors, including persons with epilepsy themselves as well as professionals from the medical, educational, and public-relation spheres.

The building and financing of comprehensive and safe epilepsy care requires the interaction of different actors: from the patient to the researcher and from the regulatory to policymakers in different sectors of the society (educational, health care, legislative). Moreover, it requires understanding of societal changes, which are leading to the greater expectation by patients for the highest possible level of epilepsy care, across borders with Europe-wide access.

6 | CONCLUSIONS

EpiXchange clearly showed that there remains an urgent need to unravel the basic molecular and cellular mechanisms of the epilepsies and integrate this information with data from population and patient cohorts. This research must continue in parallel with translational approaches in order to drive innovative solutions for therapy. The meeting acknowledged the important role of other EU-funded epilepsy research projects such as E-epilepsy and ESBACE, and the two early stage researcher training networks ECMED and EUGliaPhD in the debate. The epiXchange conference has shown how such an integrated approach can maximize the outputs and speed up the innovation process. This can be achieved through the:

- Sharing of technologies, concepts, novel mechanisms, biomarkers, and therapies;
- Sharing of common infrastructures (eg, Epilepsy Data Ecosystem) in a FAIR (fairness and accuracy in reporting) data management manner;
- Identification of synergies with other brain disease-oriented programs, where epilepsy can occur as a comorbidity, such as traumatic brain injury;
- Identification of synergies in different projects to engage patients and caregivers into the digital development and the design of applications for research and innovation activities in diagnostics and monitoring, and optimizing clinical study designs;
- Providing digital applications and innovative holistic epilepsy care models for the implementation of affordable comprehensive care.
Coordination of an open-access Epilepsy Data Ecosystem on a global level would also greatly increase the impact of efforts both within and outside Europe and provide unprecedented opportunities for international cooperation. Only such a cohesive framework of knowledge acquisition and sharing with regulatory, policy, infrastructural, and other supportive activities can lead to tangible innovative results within the mandate of the timeframe indicated. Facilitating synergies between existing European and international research infrastructures and consortia will leverage European research in this area. This would thus ensure a global role for EU epilepsy research in the worldwide research arena by 2030. In addition, these solutions will have a much wider global impact, since they can also be used to address the unmet needs of other neurologic conditions.

Therefore, the epiXchange meeting recommended the following immediate lines of action:

- Providing continuity in resources needed for the most promising innovations so as to ensure their efficient development from discovery to market and patient care.
- Supporting and aligning national strategies (eg, exit funding) to further develop the research and innovation initiated by the Horizon Europe funding program.
- Creating funding instruments that will ensure the continuation of the most successful projects and/or their confluence into other neuroscience projects/SMEs with potential to expand the impact of innovation.
- Synergize research funding activities in Europe with global initiatives to facilitate international collaboration and thus maximal impact of its research outputs.

This message will be all the more powerful if the whole epilepsy community works beyond our traditional remits in clinics, labs, and support organizations, so as to engage those key players who are in a position to provide financial support and investment in research. All this can only be achieved if we remain bold and inspirational with a wide societal relevance, yet, at the same time, focusing on the well-being of persons with epilepsy.

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CONFLICT OF INTERESTS

J.H.C. has been an investigator in clinical trials sponsored by GW Pharma, Zogenix, Vitaflo, Takeda, and Marinus. She has participated in advisory boards for GW Pharma, Zogenix, and Nutricia. She has been a speaker in events sponsored by GW Pharma, Zogenix, and Nutricia. All remuneration has been paid to her department. D.C.H. holds US patent no. US 9,803,200 B2 “Inhibition of microRNA-134 for the treatment of seizure-related disorders and neurologic injuries.” D.C.H. received funding for research described in the review article (European Union’s FP7 (602130)). He was co-organizer of the epiXchange conference. A.P. received funding for research described in the review article (European Union’s FP7 (602102)). S.S. is the Chair of the Epilepsy Advisory Group of the Association of British Neurologists, UK and Lead of the Clinical Genetics Testing Task Force of the International League Against Epilepsy. J.M. is Member of the Epilepsy Advocacy Europe and Member of the International Bureau for Epilepsy, International League against Epilepsy. We confirm that we have read the Journal’s position on issues involved in ethical publication and affirm that this report is consistent with those guidelines.

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