

Improving patient empowerment and self-management of care in mental health¹

A Call to Action

This Call to Action aims to highlight the urgent need to stimulate the empowerment of those affected by mental ill health and to recognise the vital role patients can play in the management of their treatment and care, in partnership with health professionals and other care providers.

"In the current, Europe- wide efforts aiming to develop sustainable, patient-centred and cost-effective health systems, empowered patients are an essential part of the solution.

Therefore, investing in patient empowerment should be seen as a priority".

Marco Greco, President European Patients' Forum

Why a Call to Action?

The burden of mental ill health is huge across the EU

It is increasingly recognised that the burden and prevalence of mental health conditions is huge: At least 27 % of the European (EU, Switzerland, Iceland and Norway) adult population experience mental disorders every year². The proportion of the national health budget devoted to mental health ranges between 4% to 13% across the EU. Mental health conditions account for 22% -25% of the EU burden of disability, with neurologic and psychiatric disorders being among the third leading causes of disability-adjusted life-years in the WHO-Europe^{3,4}. Social and economic inequality and exclusion are both a cause and a consequence of mental ill health.

Mental ill health can affect persons at any age in a variety of forms (e.g. depression, bipolar disorder, schizophrenia, ADHD...). These conditions go hand in hand with substantial stigma, costs and consequences that impact individuals, families and carers, health and social systems, society and the economy.

Evidence shows that life expectancy can be reduced by mental disorders, sometimes with a greater impact than smoking or obesity. In particular, mental disorders can be the precursors to chronic diseases, consequences of them, or the result of interactive effects. In addition to a higher risk of physical illness such as coronary heart disease, Type 2 diabetes or respiratory disease, mental disorders (such as

¹ http://psychopathology.imedpub.com/empowering-people-with-mental-illness-within-health-services.php?aid=17223, http://psychopathology.imedpub.com/empowering-people-with-mental-illness-within-health-services.php?aid=17223, https://www.health.gov.au/internet/main/publishing.nsf/content/CFA833CB8C1AA178CA257BF0001E7520/\$File/servpri.pdf,http://www.ccom/ssantementalelillefrance.org/sites/ccoms.org/files/pdf/Empowerment%20and%20Mental%20Health%20in%20Community.pdf

http://www.euro.who.int/en/health-topics/noncommunicable-diseases/mental-health/data-and-resources

https://www.mentalhealthandwellbeing.eu/assets/docs/publications/Framework%20for%20action_19jan%20(1)-20160119192639.pdf

depression) often exacerbate unhealthy life styles including smoking, substance abuse, physical inactivity and insufficient sleep^{1,2}.

On the other hand, poor physical health increases the risk of mental ill health: the risk of depression is doubled for people with diabetes, hypertension, coronary artery disease and heart failure, and tripled in those with stroke, end-stage renal failure and chronic obstructive pulmonary disease. Children experiencing a serious or chronic illness are also twice as likely to develop emotional disorders.

There is an obvious need to address mental ill health in the most effective way

Fortunately, good mental health is increasingly recognised as important for economic growth and social development; poor mental health is consistently associated with unemployment, less education, low income or standard of living, poor physical health, adverse life events and poor quality of life. However, across the EU, there is a lack of adequate mental health services and support available.

Empowered patients should be seen as a resource in this respect

Patient empowerment is defined 'as a multi-dimensional process that helps people gain control over their own lives and increases their capacity to act on issues that they themselves define as important'³. This concept comprises many elements, such as health literacy, confidence, coping skills and self-management of treatment and care. It is opposed to 'disempowerment' which refers to taking away power from an individual thus leaving them feeling helpless, without control over their lives and less likely to succeed⁴.

While the notion and need for patient empowerment has been increasingly accepted when it comes to patients affected by physical health conditions, this is not the case for those affected by mental ill health; many individuals and organisations have highlighted the disempowering attitudes and behaviours towards people affected by mental ill health which exist in society.

Disempowerment occurs within health care, education, the workplace and society at large. While antidiscrimination legislation, promoting empowerment of people affected by mental ill health has been put in place in some countries, and while several organisations/campaigns have contributed to positive change in this respect, discrimination and disempowerment towards people with mental ill health remain widespread⁵. The stigma attached to having a mental health issue – as opposed to having a physical health issue - is pervasive and widespread; this stigma occurs at individual, organisational as well as at societal levels and is regarded as one of the main barriers to patient empowerment.

Self-management of care and treatment is an important element within the context of patient empowerment⁶. Self-management refers to an 'active engagement of the health care consumer in dealing with his or her disorder, meaning that the person with the disorder is an active participant in care, rather

www.euro.who.int/en/health-topics/noncommunicable-diseases/mental-health/data-and-resources

² http://www.ox.ac.uk/news/2014-05-23-many-mental-illnesses-reduce-life-expectancy-more-heavy-smoking

³ http://www.eu-patient.eu/globalassets/campaign-patient-empowerment/briefing_paperpatient-empowerment_final_external.pdf

 $^{{\}color{blue}^4\underline{\,https://psychopathology.imed pub.com/empowering-people-with-mental-illness-within-health-services.pdf}}$

 $[\]frac{5}{\text{http://psychopathology.imedpub.com/empowering-people-with-mental-illness-within-health-services.php?aid=17223}{\text{http://psychopathology.imedpub.com/empowering-people-with-mental-illness-within-health-services.php?aid=17223}}$

⁶ https://www.mentalhealth.org.uk/a-to-z/s/self-management-mental-ill-health,
http://www.heretohelp.bc.ca/visions/self-management-vol1/self-management-in-the-mental-health-field,
http://www.euro.who.int/ data/assets/pdf file/0018/124560/E94377.pdf.pdf, https://www.rethink.org/living-with-mentalillness/recovery/tools-for-recovery/self-management, http://mentalhealthrecovery.com/info-center/mental-illness-self-management-throughwellness-recovery-action-planning/,

than someone who simply follows recommendations and complies with the treatment plan developed by a health professional'.¹ Interestingly, self-management can have as positive an impact on mental health as medical treatment, enabling people to lead fuller, more active lives. Self-management of chronic conditions is increasingly seen as a tool that may help struggling healthcare systems reduce costs and at the same time empower patients². Many people with physical health problems already self-manage their care and treatment and are seen as competent and capable to do so.

However, when it comes to mental ill health, questions are raised as to whether individuals affected by mental ill health are able to meaningfully and effectively self-manage their care and treatment. In fact, for many of those affected by mental ill health, lack of legal capacity is still the reality.

However, most people with mental ill health, most of the time, <u>do</u> have the competence to self-manage their treatment and care; they are able to comprehend the required information and to apply this to making rational decisions. They are able to develop the skills needed to manage their conditions: the majority of individuals affected by mental ill health have the full capacity to manage their care^{3,4}.

This Call to Action highlights what can be done to empower patients and to engage them as partners in care

Years of under-investment in mental health and in those that are affected by mental ill health means that patients often experience poorer access to services and lower quality of care than those with physical health conditions. Clearly, mental ill health needs to be better addressed as a matter of urgency and at a variety of different levels. Involving those affected by these conditions with the treatment and care decisions that affect them will ensure patient-centredness as well as treatment relevance. This in turn will translate into more effective treatment and care, which will contribute to sustainable health systems and budgets. This in turn will help increase the self-confidence and facilitate the integration on people affected by mental ill health in the community.

Patients should be seen as resource in this respect; and empowering patients will ensure the best possible 'use' of that resource, while putting patients at the heart of care provision.

 $^{{\}color{blue}1} \underline{\text{http://www.heretohelp.bc.ca/visions/self-management-vol1/self-management-in-the-mental-health-field}}$

 $^{{}^2\}underline{\text{ http://www.eu-patient.eu/globalassets/campaign-patient-empowerment/pe-conference-report.pdf}}\\$

https://www.rethink.org/living-with-mental-illness/recovery/tools-for-recovery/self-management

⁴ http://www.heretohelp.bc.ca/visions/self-management-vol1/self-management-in-the-mental-health-field

A Call to Action

The overarching principle:

The need for parity of esteem of mental and physical health:

Parity of esteem is the principle by which mental health must be given equal priority to physical health¹. In some countries, the government requires health services to work for parity of esteem to mental and physical health. However, there are many areas where parity of esteem has not yet been realised. For instance, in the UK, mental ill health account for 28% of the burden of disease but only 13% of NHS spending.

This lack of parity is a key element in disempowering those affected by mental ill health.

The undersigned call on the following stakeholders to take action as a matter of urgency in order to turn patient empowerment into a reality:

Policymakers/governments should put into place health systems...

- ...which are patient-centred, personalised, recovery-oriented and geared towards patient
 empowerment. They should be easy to access (e.g. affordable, limited waiting times, limited
 distances, provide reliable information on what is available and what patients and carers are
 entitled to) and take account of the expertise of patients in relation to their condition.
- ...which are less fragmented; there should be better coordination and communication between health and social services as well as between the various medical specialists.
- ...which aim to keep patients in the community to the largest possible extent; community-based services need to be strengthened and developed.
- ... which effectively support the civil rights as well as the capacity and competence of people affected by mental ill health to manage their condition
- ...which cater for a more holistic, proactive and preventative approach. Early detection and diagnosis is key; secondary and tertiary prevention are also crucial.
- ...which officially recognise peer support as beneficial to patients.
- ...which are geared towards supporting patients in the decision-making processes and aim for the least restrictive treatment and care alternative, safeguarding the freedom of the patient as much as possible.
- ...which grant patients the right of appeal of against decisions that are taken for him/her.
- ...which help organisations to build capacity and support their efforts to amplify the voice of
 patients as well as the practical services these organisations offer to patients and carers (e.g.
 psycho-education, peer support, dissemination of good practice.
- ...which offer people using mental health services (patients and carers) around the clock access to a crisis team.
- ...which meaningfully engage in dialogue with patients with respect to their needs and how the system is providing the care and treatment they need, giving patients and their families a voice in the development of services.

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¹ https://www.centreformentalhealth.org.uk/parity-of-esteem

- ...which recognise and take account of the pivotal role played by family carers in terms of care and support offered to relatives affected by mental ill health. Research has indicated that the risk of developing a mental health condition is 20% higher for carers who do not receive any support. Systems need to ensure that there are support services in place, such as information, training and respite care. Assessment of carers' needs and situation, and support in cases where the burden is too heavy, is also vital.
- ...which allocate appropriate resources to assure that people with mental ill health are empowered to manage their condition is so far as they are capable to do, having access to the most appropriate and innovative treatment and care.
- ...which provide reassurance to individuals in the form of distance monitoring and the use of 'E' and 'M' Health.

Policy makers should also take action with respect to the identification and dissemination of good practice – between professionals, patients, family carers and other stakeholders at local, regional, national and European levels – as this would help to share and develop evidence-based models and enable positive change.

They should also provide incentives to employers to enable people affected by mental ill health (and their carers) to either enter, participate and stay in the labour market.

Governments should launch awareness campaigns to educate society about mental ill health and facilitate more informed views and opinions.

Health professionals should...

- ...be aware of their crucial role in facilitating and stimulating the empowerment of those affected by mental ill health. They should work in partnership with their patients and treat them with dignity, compassion and respect, working towards the highest level of shared decision making.
- ...legally and ethically maximise the autonomy of those affected by mental ill health and make use of non-invasive, non-traumatic treatments.
- ...inform and involve family carers in treatment and care plans of their relatives as they play a pivotal role in the implementation and success of these plans.
- ...take the potential of peer support into account when developing and implementing treatment and care plans.
- ...have a positive outlook and address the strengths of patients rather than focus on their limitations. Patients should be viewed in a holistic way; the focus should be broader than their condition only. Treatment should have a recovery focus and should be multi-modal (medication, psychotherapy, coaching, life-skills...)
- ...view the provision of information to patients and carers as a key professional task; they can contribute to health literacy a crucial element in self-management of treatment and care. Information should be accessible (easy to obtain and easy to understand). Where and when needed, patient and carers need to be informed of the specific condition and its warning signs, medication prescribed and possible side effects of pharmacological treatment.
- ...regularly check the physical health of people affected by mental ill health and the mental health
 of those affected by physical health problems. Being diagnosed with a specific (mental or physical)
 condition should not lead to a sole focus on that condition: in many cases, patients are affected
 by multiple conditions and these should all be addressed in parallel.
- ...put in place better ways of communication and cooperation with the different medical disciplines involved (GPs, psychiatrists, other medical specialists or health care professionals) as well as with social services.

- ...pay special attention to children and their mental well-being. Early detection is key; the earlier the detection, the better chance of recovery and effective treatment.
- ...ensure a seamless transition when patients move from childhood to adult mental health services.
- ...should strive for informed consent and avoid coercion to the largest extent possible.

Patients should...

- ...make efforts to get better organised in order to strengthen their voice as well as allowing for the provision of peer-support (counselling, information, practical support, psycho-education...).
 Peer support networks are self-empowering as well as helpful to patients.
- ...take an active stance and learn about their condition and its management, as experts of their condition and their needs and requirements.
- ...become and be more aware of the positive role they can play in managing their care and treatment, in so far as they are capable of doing so.
- ...engage in meaningful dialogue with policymakers (regarding services development and implementation) health professionals and family carers (working in partnership towards shared decision making).
- ...participate in training and educational programmes, sharing the views and experiences of patients. This will help bridge the gap between patients and health professionals.

Informal carers should....

- ...advocate for recognition of their role, contribution in the care provision and knowledge of their
 relatives affected by mental ill health. This contribution needs to be recognised by
 policymakers/governments and catered for by health systems and health professionals.
- ...call for better access to information regarding the condition of the patient they care for, of the basic rights and entitlements, and the treatment plan in place.
- ...advocate their needs for support, in practical as well as psychological terms.
- ...request access to training to help them with the daily care needs of their relatives.
- ...advocate the option NOT to provide care. Caring should not be seen as an obligation but rather as an informed choice.

Social services should...

- ...ensure the involvement of patients and their carers when deciding on what services are required (e.g. supported housing, contacts with social workers...) and how these should be provided.
- ...take a holistic approach and take the entire situation of the patient into account.
- ...work with the other health professionals involved in order to ensure coordination and integration of care.

Health insurances should....

- ...ensure equal priority to physical and mental health in relation to reimbursement of the best and most appropriate treatment and care. Mental health remains under-resourced across the EU.
- ...recognise that prejudice plays a significant part in decisions that relate to reimbursement of mental health treatment and care. The coverage in this area is much less that the coverage for
- physical health conditions. This holds true for medical interventions and treatment as well as for psychotherapy, counselling, peer support and other forms of treatment.

• ...address the lack of parity of esteem between physical and mental health as a matter of urgency, also in view of the fact that there is increasing evidence that the return on investment is highest in the field of psychiatry.

Medical schools should...

- ...aim to ensure a holistic approach in medical training and education curricula, both those aimed
 at medical students as well as those targeting retraining and life-long learning curricula.
 aim to stimulate an open mindset, in order for health professionals to take the views of patients
 into account and be open to the concept of shared decision making and patientcentredness.
- Patients can be involved in education with respect to shared decision making and patient-centred
 care, which is useful for medical students and health professionals on the one hand side as well
 helpful to empowering patients on the other.

Primary and secondary schools should

- ...address mental health in their educational programmes and encourage children to speak about their emotional problems. This would help 'normalise' mental ill health.
- ...change their policy and, rather than trying to get all children to do the same things, treat children as individuals and help them to find their individual strengths.

Research institutes should...

- ...engage in more dedicated research in the field of mental health. The EU-funded ROAMER¹ project resulted in a roadmap for priority research, focusing on social, societal and medical aspects of mental ill health and provides a useful guide towards useful research. Examples: patients and their specific needs and requirements, the impact of interventions and treatment (medical, psychological, peer support...), decision making models.
- ...not forget about carers when undertaking research in the field of mental health.
- ...redress the balance and ensure that mental health receives the same amount of attention as physical health in health research.
- ...include patients, and their family carer(s), where possible, in an advisory capacity to ensure the relevance of research to those directly affected.
- ...pay attention to the translation of research findings into (good) practice.

Employers/social partners should...

- ...offer adapted working conditions to enable people affected by mental ill health to enter, participate and remain in the labour market, as good work is empowering and beneficial for mental health and well-being. Work can support recovery.
- ...address the lack of understanding as well as the lack of knowledge about how to support people
 in the workplace. Some examples of support measures: flexible working hours or patterns,
 changes in the physical environment, support with work load and support with/from colleagues.
- ...share examples of good practice of what works.

¹ https://cordis.europa.eu/result/rcn/171328 en.html

Annex 1: LOGOs of endorsing organisations to be added

Annex 2: How this Call to Action was developed

On 7 March 2018, GAMIAN-Europe convened an expert workshop on the issue of empowerment and self-management of mental health care and treatment, as a first and crucial step towards the development of this Call.

This meeting brought together key stakeholders with an interest in empowerment of patients and better management of their care (this includes patient organisations, health professionals organisations (psychiatrists, psychologists, GP's, mental health nurses), carers and policy makers and had the following objectives:

- 1. explore the current situation in relation to the empowerment of people affected by mental ill health and their ability to self-manage care and treatment;
- 2. identify the barriers to empowerment and self-management of care, both to patients as well as their carers;
- 3. come forward with recommendations to improve the situation.

The outcome and content of the discussions was used as the basis of a Call to Action on patient empowerment and self-management of care, which was then further elaborated; participants had the possibility to take a critical look at a draft and provide input and comments. This second was discussed in a meeting of the European Parliament Interest Group on Mental Health, Well-being and Brain Disorders and finalised on the basis of input received on that occasion.

Annex 3: Workshop registered participants

1. AD	OHD-Europe	Joanne Norris	Board Member
2. AD	OHD, ASC&LD Belgium	Beverley Sinton	Vice -President
3. An	noiksis	Leo Finn	Board Member
4. EU	JFAMI	Margaret Walker	Secretary General
5. Eu	ırocarers	Stecy Yghemonos	Director
6. Eu	ropean Brain Council	Fred Destrébecq	Executive Director
	uropean Federation of sychologists Associations	Ruta Sargautyte	Associate Professor Psychology, Vilnius University
8. Eu	ropean Patients' Forum	Lyudmil Ninov	
	uropean Psychiatric ssociation	Mari Fresu	Policy and Scientific Officer
	uropean Psychiatric ssociation	Ulrike Kool	Trainee
11. Fe	errer	Alfredo Gracia	Corporate Director, Patient Advocacy & Public Affairs
12. GA	AMIAN-Europe	John Bowis	Board Member
13. GA	AMIAN-Europe	Dolores Gauci	Board Member
14. GA	AMIAN-Europe	Lina Papartyte	Secretariat
15. GA	AMIAN-Europe	Christine Marking	EU Advisor and workshop organiser
16. Me	ental Health Europe	Marcin Rodzinka	Project Manager
	ospital Gregorio aranon	Alfredo Calcedo	Professor of Psychiatry
	anding Committee of uropean Doctors	Markus Kujawa	EU Policy Advisor
	nion Européenne édicaux Spécialistes	Marc Hermans	President Psychiatry Section
20. Un	niversity of London	Angela Coulter	Freelance researcher
21. Up	os and Downs	Erik Van der Eycken	Board Member
22. Up	os and Downs	Anouk Drieskens	Board Member