



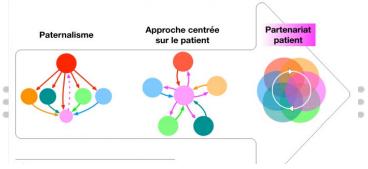
How the Partners in **Care Approach Promotes Mental** Health in the community and viceversa:

the example of the Health Partnership project at UHG Paris

Context



Du paternalisme au partenariat patient





ENTRETIENS JACQUES CARTIER

3 novembre 2020 par visioconférence

Contact : Pre. Marie-Pascale Pomey - email : marie-pascale.pomey@umontreal.ca

Inscriptions : <u>plateforme.entretiensjacquescartier.com</u> Pour toute question, vous pouvez contacter les Entretiens Jacques Cartier à l'adresse suivante : eje®centrejacquescartier.com



VERS UN PARTENARIAT USAGERS-ÉQUIPES DE SOINS RÉUSSI POUR AMÉLIORER L'EXPÉRIENCE DES USAGERS ET DES PROFESSIONNELS

Comment développer le partenariat usagers-professionnels dans les établissements de santé ?

Guide de recrutement de patientsintervenants





Véronique Gauthier Marc Tremblay

Une expérience de patient-partenaire ayant une maladie rare en recherche

HAUTE AUTORITÉ DE SANTÉ

CHIATRIE &

SYNTHÈSE

Soutenir et encourager l'engagement des usagers dans les secteurs social, médico-social et sanitaire

Validée par le Collège le 23 juillet 2020

RECOMMANDER LES BONNES PRATIQUES



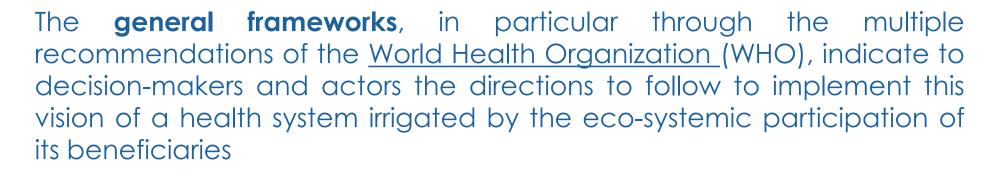
For about 40 years, the **involvement of users and relatives in care** has helped initiate health democracy (France and elsewhere) (Gross and Gagnayre, 2017; Merle et al., 2021; Simon et al., 2020; Tourette-Turgis, 2019).

This phenomenon is visible in:

- research (Jouet, 2014),
- healthcare (Fournier and Troisoeufs, 2018; Gardien, 2018; Pétré et al., 2020),
- training (Berkesse et al., 2020; Pétré et al., 2020; Towle, 2019)
- user control (related to user skills) (Coirié and Delanoë-Vieux, 2021).







"WHO's vision is to put the complete needs of individuals and populations, not just diseases, at the center of health systems and to empower people to play a more active role in their own health" (Coldefy and Maugiron, 2022)



Context



Increasingly robust studies show the positive effects of user participation in care, health, and support systems.

(Gross, 2017; Heijboer, 2020; Merle et al., 2021; Tourette-Turgis, 2019).



The patient partnership - general definition

"In the field of health care, it refers to a person with a specific mission.

The term 'patient partner' is a fairly general term for people who collaborate with care teams on a regular basis to either improve the quality and safety of care and organizations, or to conduct teaching or take part in research."

Guide HAS – Soutenir et encourager l'engagement des usagers dans les secteurs social, médicosocial et sanitaire – 23 juillet 2020



Montreal Model - The patient partnership – Basic Principles

- The construction and value of the **experiential knowledge** of the people concerned by a situation of vulnerability are **recognized and legitimate**;
- The patient becomes a partner in care, i.e. he/she is considered and experiences him/herself as a person progressively empowered, in the course of his/her clinical journey, to make free and informed health choices;
- The user has become a caregiver for himself;
- The partnership relationship between professionals and users has as its main objective **the realization of the patient's life project by recognizing and mobilizing the knowledge of all parties**, including that of the patients/relatives considered as members of the clinical team in the context of their care;
- **Co-construction** is favored to work in partnership: a collaborative method that favors the emergence of a common basis of understanding between patients, family members, stakeholders (health professionals) and managers (administrators);



8

MontreaL Model - The patient partnership – Basic Principles

- Health care partners come **together** and **work together**: all the actors involved in health and social services choose to co-construct the care environments of tomorrow. They may be patients and users, their families, health and medico-social professionals (internal or external to the institutions), and administrative staff in the health and medico-social sectors;

- The individual and collective aim is to encourage the process of empowerment;

- It is a perspective that refers to **health** rather than illness (salutogenic versus pathogenic);
- **Recovery** (versus healing) is the basis of care and support practices;
- Peer support practices are supported and promoted.



MontreaL Model - The patient partnership – Basic Principles

...

		Patient resource	
	Level 1 - The patient as a partner in his care	Level 2 - The patient trainer, the patient intervener, the patient researcher	Level 3 - The patient co- leader, the patient co-worker
Objectives	The patient is a co-builder of his or her care, his or her care program, his or her life project; the patient caregiver	The patient who has become competent for himself is trained to intervene with others: peer helper, patient trainer, "peer practitioner", patient researcher	The patient is involved in hospital governance and health democracy, the patient as co- worker
Roles	 >> Receive information about their illness and its consequences, be able to question it, >> To make their opinion known on therapeutic choices and their preferences in order to reach a mutual agreement on care; to share the decision, >> To benefit from personalized care, >> To be trained in self-care and to practice it >> Complete advance directives and joint crisis plans, >> Participate in education programs, >> Participate in caregiver/caregiver meetings. 	 >> Become a peer helper, peer health mediator >> Co-produce and animate ETP programs >> Co-produce and lead training programs for patients, professionals and future professionals >> Participate in research and evaluation 	 >> Participate in the governance of the hospital and in health democracy > Participate in the hospital's decision-making processes (steering groups and thematic work) >> To intervene as an expert/resource on specific projects

MontreaL Model - The partnership with carers – Basic Principles

The carers involved in the accompaniment of a patient also have a role to play at all levels.

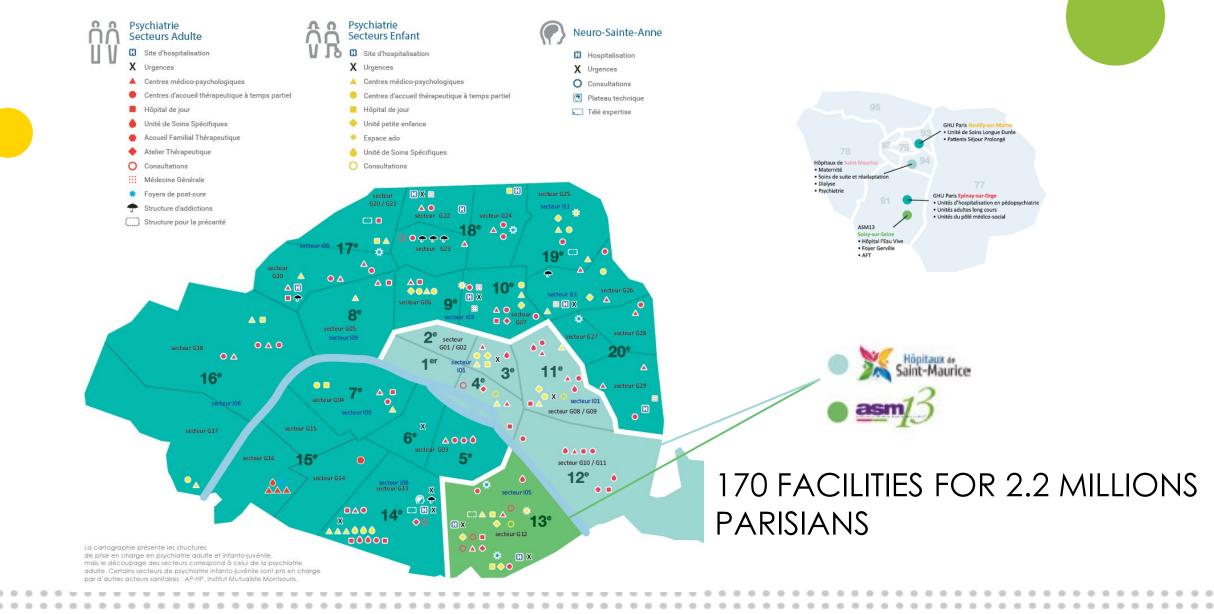
		Carer Resources		
	Level 1 - The carers partner in healthcare	Level 2 - The carer trainer intervener, researcher	Level 3 – The carer co-leader, co-counselling/builder	
Objectives	Depending on the situation and with the patient's agreement, the family member/Carer participates in defining the care program and the life project	The family member/carer is trained to intervene with others, especially family members/carers: peer helper, family/carer trainer, family/carer researcher	The family member/carer is involved in hospital governance and health democracy, the family member is a co-worker	





Reminder: what is UHG Paris?







Why work on the public patient partnership at UHG Paris?

- An **international dynamic** (Quebec, Belgium, Great Britain, United States, Switzerland, Catalonia...)
- Increasingly present in French public policies (Law on the modernization of the Health System 2016, HAS recommendations July 2020, Ma santé 2022, ...)
- Who is starting to find practical variations in **France** (University of Patients, Patient-Partner Program of the Institut Curie ...),
- Which is based, in psychiatry, on a new approach to care: recovery,
- Which has objective effects on the quality and organization of care.







The objectives of the implementation of the public patient partnership at the UHG

- Improving the quality of care: "There is no possible optimization of the quality of care and support without the active participation of the people concerned"* and in particular: Improve people's experience and experience when they are accompanied by the GHU: satisfaction to see their preferences taken into consideration, less suffering related to poorly lived practices, etc.
- **Promote the empowerment of users**, i.e. strengthen their ability to acquire greater control over decisions and actions affecting their health
- Improve the GHU care offer and pathways through better alignment with the needs and expectations of users
- Improve the working conditions of professionals, in particular by improving caregiver/patient relationships

* Source : HAS – Projet stratégique 2019-2024







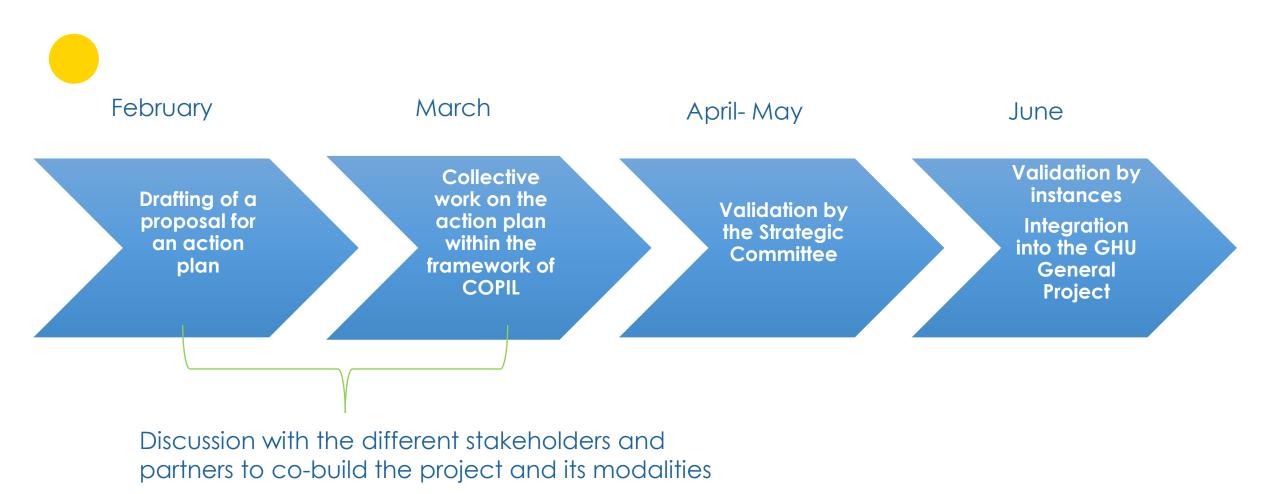
Methodology and timetable







Public Patient Partnership Action Plan 2021







Adoption of the 2021-2025 establishment project A course: better patient care and improved quality of life at work



Public Patient Partnership Action Plan 2022



- ✓ Training of Health professionals in patient partnership,
- Strengthening Public Patient Partnership programs involving patients at all stages: design, facilitation, evaluation,
- ✓ The deployment of health peer supports in the services,
- ✓ The involvement of patients in decision-making spaces and projects





- Favorable conjunction, with the support of the medical community, strong institutional support and the presence of competent professionals already invested in these issues;

- At the beginning of the process of this cultural transformation;



- Inside the hospital:

- the presence of motivated professionals committed to this partnership dynamic is not yet the norm → the dissemination in the training of future health professionals is only a few years old and not totally disseminate in the French universities;
- The appropriation of these new models implies changes in practices and social representations that it is essential to accompany with a reflexive look



- Inside the hospital:

- Despite the presence of associations, their roles are still very limited to the representation and the fight for rights (complementary but different from the health partnership)
- Representations still very focused on the disease (even if a lot of change with the effects of covid on awareness
- Still many doubts about the legitimacy, value and place of experiential knowledge in institutions
- At the moment: health professionals career crisis, lack of nurses and psychiatrists, decrease of beds without a dynamic and structured health community policy + Fragile governance of public health between primary, secondary and medico-social care



- Outside the hospital:

- Links with municipalities, care in the city and city medicine: always difficult
- Patient partnership:
 - a model too focused in illness and not health, to close to the medical knowledge, (the name: patients, try to have it called patients and the publics...)
 - lack of people and culture diversity (cultural communities, LGBTI communities, ageing communities not so much taken into account in the model at this stage)
 - Not addressing the main issues such as coercion, seclusion, human rights...
 - Leaving groups of people behind: programs not adapted to specific needs
 - Risk of tokenism and staying at the surface





Question for the future:

Links with the communities:

Obstacles? Facilitators? For the partnership

