IMPACT OF THE FREE INFORMATION SERVICE MY CANCER NAVIGATOR ACCORDING TO PATIENTS AND THEIR RELATIVES

AN ONLINE SURVEY

CN20

<u>Gabry Kuijten</u>¹, Jorn Hermans², Liese Vandeborne¹, Klara Rombauts¹, Guy Buyens¹, Manuel Bollue³, Patrick Deroost³, Rosanne Janssens², Isabelle Huys²

¹Anticancer Fund, Meise, Belgium; ²Department of Pharmaceutical and Pharmacological Sciences, Catholic University Leuven, Leuven, Belgium; ³Optigage, Keerbergen, Belgium

BACKGROUND

My Cancer Navigator (MCN) is a personalised service of the Anticancer Fund*, aiming to address information needs and contribute to informed and shared decision-making (SDM). To this end, a team of scientists and physicians, answers therapy-related questions of cancer patients and their relatives in an evidence-based manner.

An explorative and descriptive cross-sectional study was conducted to assess the perceived impact of MCN on users' emotions, knowledge level and communication with healthcare professionals (HCP), all factors contributing to SDM^{**} .

METHODOLOGY

- Online surveys consisted of quantitative (Likert scale, multiple choice) and qualitative (open-ended) questions, based on a structured literature review.
- Surveys were disseminated in February-March 2023 via email to adult patients and their relatives who used the MCN service in the previous year.
- Patient demographics were extracted from the MCN case files obtained as part of the standard procedure to which all service users consented.
- Catholic University Leuven ethics approval (ref S67258).

RESULTS

- Of 253 invited MCN users, 109 (43.1%) filled out the survey. Respondents were 64.2% patients and 35.8% relatives.
- Overall knowledge level improved for 86.3% of respondents, with no change for 13.7%.
- Regarding impact on emotions, 72.5% of respondents reported a positive change overall; 16.5% reported no and 11.0% a negative change.
- Of all patients among respondents, 68.5% indicated an improved ability to cope with their disease after MCN while 76.4% of relatives said they were able to better support the patient.
- Regarding impact on communication with HCPs, 37.8% reported a positive change.
 48.5% of patients among respondents said discussing treatments with HCPs got easier and 44% found joint decision-making easier.
- According to 52% of respondents, disease management changed after having used the MCN service.

CONCLUSION

MCN supports shared decision-making in cancer treatment for both patients and caregivers, by positively impacting users' emotions, coping skills, knowledge level and communication with HCPs.















