



Review article

The Obligation of Informed Consent and the Best way Contact with a Patient Who is Diagnosed with a Gynecological Cancer

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Summary

Human relations are a meeting point for many fields of science and of the natural and social sciences, especially when studying the relation doctor and patient. Closely linked to communication are the concepts of satisfaction or dissatisfaction of the patient, but also of his compliance or non-compliance. The phenomenon of dissatisfaction expressed by the patient towards the literature for several decades. The main factor of dissatisfaction was the ambiguity in communication between doctor and patient. Although the importance of detailed patient information is now recognized or has been enshrined in law, however, in everyday clinical practice the provision of it is not systematic or does the tendency to conceal the truth prevail? Also, there is also the informed consent which constitutes a necessary condition for conducting a medical procedure, but also the provision of health service.

Keywords: Consent, Patients, Communication, Cancer, Consent, Informal.

Import

Gynaecological cancer is the most common form of cancer and the need arises for ongoing assessment and appropriate supportive care, as well. Patient information has been the subject of many research studies. There are different practices in many countries when it comes to inform patient support. The focus is on the use of theoretical models with the aim of improving communication between the doctor and the informed consent of patients are protected by clear legislative settings.

Also, communication and information about the disease and the upcoming are increasingly considered to be of paramount importance to the prognosis and treatment of the disease. However, many health professionals tend to underestimate the amount of information patients need and remain sceptical about informing patients.

Informed Consent

Doctor-patient communication is not just a mean medical work, nor just social convention and necessity. It also constitutes ethical

obligation of the doctor and should respond to specific requirements of the Code of Medical Ethics. The obligation of the doctor medical procedures without the patient's consent, as well as his obligation to the patient information about the general state of his health, on the one hand the purpose and risks of impending medical procedures.

Information Methods

The most well-known methods of patient information, which have been studied in recent decades, are the "ABCDE" method, the "SPIKES" method, Hitoshi Okamura's method and the method of Peter McGuire and Ann Fa. The first two methods are considered the most important of these, while mainly suggestions / instructions for proper and effective information. OBSTACLES TO PATIENT INFORMING Unfortunately, all studies that refer to Greece or abroad, they agree that this is a process with particularly and various obstacles. According to Beckman, these obstacles fall into two categories. a. Obstacles for social reasons. b. Obstacles concerning the patient and which are referred to health professionals. The information of the patient and the time, place and manner in which the diagnosis will be announced seems to be of significant concern to health professionals worldwide. They believe that to achieve quality information, education is required responds to current

data and patient demands. The lack of this due to education is the main reason for doctors' decision to conceal the truth from cancer patients. The same is true of nurses. Although that nurses seem to be the most suitable health professionals for them patients, to share their thoughts and feelings, they themselves feel inadequate to communicate with them. Specifically, the issues on which What health professionals are lagging behind is the announcement of b of mourning, support for relatives and management of emotional cries cause, therefore, the importance of education in information and announcement of the unpleasant news, it is necessary to conduct more regarding patients' opinions and preferences on this specific topic.

Refusal of Consent

OR consent of patient, such as already has was emphasizes necessary for thed, <legalization> of the medical procedure to which he is about to undergo respect for the patient's will as a manifestation of his self-determination the patient's will is negative regarding the performance of the medical If we have a refusal to provide consent, then the doctor must respect it In the past it was very doubtful whether such a refusal should be respect especially when it concerned the sole salvation of life, in which case it i self-determination, a form of disposition of the highest hierarchically le is a source and condition for the existence and exercise of other fund am But such a refusal must be strictly personal, that is, given only by the patient himself, must have been fully informed beforehand The patient's decision must be free, uninfluenced, mature, well-balance It is granted after proper and thorough information. Of course, it should be noted that when the patient refuses medical intel must make efforts to persuade him. The patient's will is proposed to it is found from a large-scale multi-disciplinary commission where it works effectively, so that an in-depth psychological diagnosis of the real the patient's will, with objective criteria. The patient's will in the case o she must be fully informed. The act to which the patient "consents", the discontinuation of treatment, in some respects, can be considered "the benefit" of the patient, since the patient seeks this and considers it to b himself.

Legislation

The principle of informed consent, which aims at the legitimacy of health care, tends to reflect the concept of autonomy and self-determination decisions of the person who needs and requests medical interventions. type, in recent years, has gained not only significant space but also important processing and approaches to doctrinal texts, as well as legal interpret influencing the daily activities of the medical profession. The informed consent is still a subject of ongoing exploration, not only with regard to the already confirmed theoretical profile, but also on the practical and consequential aspect (book 1,2,5,6,7,8). The principle of co is a

relatively new condition. In fact, already since the time not only of the civilization, but also of the Greek and Roman, documents have been fou the doctor's intervention had to, in some way, first be approved by the Plato (Law IV) had already foreseen the problems, procedures and way information that are, in composition, at the root of the principles of the informed consent and related the practice of information and of consent with the quality and social status of the patient. A form can b recognized, in Hippocrates, who cared about the patient's pain, but net neglected to take care of his own result, trying to avoid involved in the failure and death of a patient. Hippocrates respected a professional responsibility that was more religious and moral in nature from a legal point of view, very weak, to the extent that it depended on compiled by humans. Christianity was incorporated into this unified vi the sacred character of medicine and medical practice, which has not c essentially the Hippocratic type of ethical behaviour. Therefore, invested the power that came from his professional role and from his own work, he considered it his duty to guide the patient, deciding also for him (book 1,2,3,4,5,6). The socio-medical environment is changing. Toda must fully explain the planned medical treatment and any alternative solution to the patient, who has the right to choose the treat take into account the benefits and side effects. The physician has a "duty patient. Accordingly, the patient has the right to information regarding which are necessary for him to make his decisions regarding the condo medical procedure to which he is about to undergo. This emerges explicitly and clearly from the Code of Medical Ethics why was passed in 2005 and also by the Oviedo Convention. The Medical Co Ethics includes a special chapter on this subject. Based on the ethical principle of patient autonomy provides that <The physician mu informs the patient about their health status, as well as about any med which concerns him and obtain his consent. The code also provides that Consent is at any time revocable at the patient's sole discretion. In add From the very beginning of the Code, it is stated that the practice of the an interpersonal relationship between doctor and patient that must be mutual trust. Important articles related to the issue of informing the patient in the Code of Medical Ethics is article 1 paragraph 4b. Also the Article 9(1) states that the doctor gives priority to the protection of heal of the patient, while part of article 11 legislates that the doctor general truth to the patient. Furthermore, article 12 states that the doctor is not proceed with the performance of any medical procedure without the p patient.

Research Data

OR updated consent constitute basic element of modern medicine, reflecting the patient's fundamental right to participate in decision-making concerning their health. Despite its critical importance, there is a lack o about real-world practices regarding informed consent of patients in the context of the modern high-pressure medical environment. JAGO/NOGGO survey on practices in

daily routine from 24 February 2022 to September 14, 2022, exploring practices and challenges concerning informed consent in hospitals throughout Germany, Austria, and Switzerland, using a specially designed questionnaire. The purpose of this study was to evaluate the practices and challenges of the real world regarding informed consent in gynaecology and gynaecological oncology in various types of hospitals. Based on over 200 responses from gynaecologists, the research demonstrates the critical need for structured discussions about informed consent, with over 80% of participants expressing interest in courses related to this aspect. It is worth noting that a significant percentage of doctors (59.9%) reported discussions about procedures they had never personally witnessed. Observed important differences between types of hospitals and of professional groups in the frequency of discussions on the updated consent, with limitations arising from factors such as time constraints, language barriers and insufficient resources for patient education. In a psychological burden experienced by physicians after discussions about consensus highlights the need for systematic changes to mitigate concerns about patient safety, legal implications and patient satisfaction.

A descriptive study was published in Greece to document the needs supportive care of patients with gynaecological cancer before surgery in *Hellenic Journal of Nursing* in 2022. This is a descriptive study with 150 women diagnosed with gynaecological cancer. The patients were approached preoperatively during their hospitalization for planned surgery at an oncology hospital in Attica. Admission criteria were age > 18 years, the ability to speak and understand the Greek language, to have scheduled for surgery as well as to consent to the completion of the questionnaires. Women with breast cancer were not included because specific group of patients there are many relevant studies in contrast to other types of gynaecological cancer (article 3). The average age of the patients was 61.1 years old, 56.9% were married, retired (33.3%) with children. The most common diagnosis was endometrial cancer (47%). The stage of highest percentage of Grade I (39.5%). Median time since diagnosis was 2 months. The vast majority of women in the sample (94.8%) reported a need for care, while the average number of needs was 10.9. Patients with gynaecological cancer in the present study expressed more needs compared with oncology patients with advanced disease. They found that 83% of at least one moderate to high level need. In general, as stated in a systematic review, 25-60% of patients with gynaecological cancer report physical, social or psychological needs due to the consequences of cancer treatment regardless of the phase of treatment. The three groups. The most frequent occurrence was the information needs related to diagnosis and prognosis (86.3%), communication needs (81%) and need support (79.2). Correspondingly, the three groups of needs with the low appearance were the needs for home infrastructure (23.5), the needs of treatment (60.4) and needs for

financial matters (67.3). The three needs a higher percentage was the need for more information about the future situation (85.3%), the need for more explanations about the treatment need for others to be less upset about their situation (69.6%).

In contrast, the three needs with the lowest scores were the need for help to eat, dress, go to the bathroom (12.7%), the need to talk to priest (19.6%) and the need for better hospital services (23.5%). The present study highlights supportive care needs of women with gynaecological cancer immediately after diagnosis and before surgery. At the clinical nursing practice, the significance of the findings lies in highlighting the necessity nurses' focus on taking measures to identify and cover the specific needs, providing adequate and understandable information to areas, encouraging patient communication with the team of healthcare as well as providing appropriate personalized supportive care.

Conclusions Patient information is a matter of global interest, it concerns societies, regardless of their cultural particularities, and is linked to the informed consent and the announcement of unpleasant news. Further crucial and integral part of the therapeutic relationship with the patient care, while requiring particular interpersonal skills and communication health professionals. In clinical practice, the relationship between the doctor and patient changes from a traditionally submissive relationship in which the doctor's authority in a mutual contractual relationship based on the doctor and the patient's consent. The doctor must fully explain the planned and alternative treatments and the patient has the right to choose after considering the benefits and side effects. Well-documented informed consent may have the advantages of providing feedback on whether the treatment plan for patients is well established and impel in terms of adequacy and effectiveness, as well as the right to self-determine patient. However, given the differences in medical systems and environments The nature of informed consent inevitably varies between institutions.

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