Considerations On Cross-Border Healthcare Services In The Context Of The Standardization Of The Legislation At European Level And The Implications On The Romanian Healthcare System

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Abstract
In the context in which EU citizens were recognized the right to get treatment abroad by the Court of Justice of the European Union in several specific cases, starting with decisions dating back more than a decade, and these decisions became part of the European acquis, this Article reviews the way in which the judgments of the Court of Justice of the European Union were refined within Directive 2011/24/EU. In the field of cross-border medical services, a certain overlap is somehow reached between the law of the Union and the national law, so that European law in many cases is essentially limited to indicating a binding objective, i.e. the achievement of the free movement of citizens patients and their equal treatment, regardless of nationality, in relation to national authorities, while preserving the competence of the member states. Against this overlap, the article aims to analyse how Romania obliged to submit to the regulatory framework imposed by primary and secondary legislation, manages to ensure the sustainability of the current model of the healthcare system, in order to increase its efficiency and effectiveness, all the more since the European Commission has established the role of healthcare as part of the Europe 2020 Policy. Keywords: European Union, patients’ rights, cross-border healthcare, the case law of the Court of Justice of the European Union, the standardization of European law, European health strategy, the Romanian healthcare system.

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INTRODUCTION
In exercising the powers which have been conferred in the interpretation of EU law, the Court of Justice of the European Union has developed over time a rich case law which has been refined with each application for a preliminary judgment submitted to the Court by the courts of the Member States. Thanks to the very rich case law in this area, the right of EU citizens to use free of constraints cross-border healthcare services, which is generally known by the term “patient mobility”, could be clearly outlined.

The Court has paved the way for the implementation of the right recognized at Article 35 of the Charter of Fundamental Rights of the European Union [1] for every person to have access to preventive healthcare and to benefit from medical treatment.
Through the case law of the Court, restrictions could be eliminated in the form of national regulations, which stood in the way of creating an internal market in healthcare delivery.

In our opinion, from the constant case law of the Court has arisen certain important principles for the conditions in which, in accordance with the provisions on the freedom to provide services, patients are entitled to receiving medical care in other Member States and to the reimbursement of these treatment by the health insurance system to which they belong.

The principles developed in that case law were considered components of the acquis of the EU, which the European legislator has taken into account in the development of Directive 2011/24/EU of the European Parliament and of the Council of 9 March 2011 on the application of patients' rights in cross-border healthcare [2].

THE STANDARDIZATION OF THE EUROPEAN LEGISLATION IN THE CROSS-BORDER HEALTHCARE FIELD

Given that healthcare was excluded from the Directive 2006/123/EC [3] on services in the internal market, it has become imperative that these aspects be addressed through a legal instrument in the European legal context, through which the principles established by the Court of Justice be applied generally and effectively.

The right of EU citizens to get treated abroad was recognized by the Court of Justice of the European Union in several specific cases, starting with decisions dating back more than a decade, these decisions becoming part of the European acquis. However, individual decisions of the Court have not been assimilated coherently by national legislations, practically existing many situations in which patients were forced to solve problems of access to treatment abroad on their own by taking the entire legal route to the Court of Justice. Unfortunately, often the Court's decision came only after the patient's death, although a saving treatment would have been possible in a Member State other than that of residence.

Given the fact that at European level, planned and emergency treatment costs abroad represent only about 1% of public expenditure for healthcare [4], the simplification of the access procedure to cross-border medical services has become a moral imperative.
The problem of the legislative gap between Member States has become more visible with the adoption of the Treaty of Lisbon [5]. Treaty requires common standards at the level of social and medical assistance (Articles 34 and 35) and explicitly encourages, in particular, „cooperation between the Member States to improve the complementarity of their health services in cross-border areas.” (article 152).

Complementary to the rights of citizens under the Treaty, has emerged the need for a European law that clarifies the responsibilities of Member States towards the patients.

The road towards the harmonization of medical services in Europe, opened by the introduction of the European Health Insurance Card, ought to continue with a pan-European development of patients’ rights. From the earliest days of European integration and to date, the European institutions have actively promoted intra-European movement. The initiative to introduce a European Health Insurance Card to replace the prior necessary documents to access medical treatment during a temporary stay in another country falls within the same general phenomenon.

Since around the values expressed by the case law of the Court of Justice of the European Union there have been a number of uncertainties, which made them difficult to apply in practice, this development of patients’ rights did not occur ab initio, requiring the intervention of the European legislator to clarify the situation through a directive to support the provision of cross-border healthcare, both for the benefit of patients and of the national health service.

In the field of cross-border medical services, a certain overlap is somehow reached between the law of the Union and the national law, so that European law in many cases is essentially limited to indicating a binding objective, i.e. the achievement of the free movement of citizens patients and their equal treatment, regardless of nationality, in relation to national authorities, while preserving the competence of the member states.

Against this overlap member states are obliged to submit to the regulatory framework imposed by primary and secondary legislation, to the extent that they are not allowed to violate EU law when exercising their powers.

Important institution of the European Union, the Court of Justice is the one that assesses the scope of the EU legal framework established by Article 49 EC for the
exercise of the competences of the Member States. It is also incumbent on the Court, assigned by the founding treaties, that by the interpretation given to a provision of European law, to clarify and specify its meaning and scope, such as to be understood and applied from the time of its entry into force.

THE REFINEMENT OF THE CASE LAW OF THE COURT WITHIN DIRECTIVE 2011/24/EC

The main principles proclaimed by Directive 2011/24/EU law have their legal source in a long series of cases in which the Court identified the limits imposed by the EU legislation on the restrictions in Member States of the right of patients to use medical services across national borders within the European internal market.

Most of the provisions of Directive 2011/24/EU aim to improve the functioning of the internal market and the free movement of goods, persons and services in the healthcare field. Given this aspect, the legal basis for the adoption of Directive 2011/24/EU is found in the provisions of Article 114 TFEU. The Union’s legislation is based on this legal basis even when public health protection is a decisive factor in the choices made, Article 114 TFEU expressly stipulating that in this regard, a high level of human health protection has to be ensured, taking account in particular of any new development based on scientific facts.

Thus, patients citizens are free to choose the Member State of the European Union and the preferred institution for medical treatment, social insurance offices in the State of residence assuming treatment costs in the same proportion as in at national level.

The European regulatory framework, aiming a new scheme of monitoring the services provided, was created precisely to enhance the quality and safety of healthcare services. The free movement of patients, without the legal force of European regulation, would have produced a competition between the health systems of the Member States in order to attract more patients. There is the risk that the free access to cross-border medical services may produce a drop in the price of medical services throughout the European Union to the detriment of the quality of health services.

The new cross-border healthcare system favours rare disease patients, whose treatment requires costly investments in research. The existence on the European
internal market of health services of specialized hospitals on these diseases prevents the waste of resources due to the parallel investment in equipment and research and also provides for closer cooperation between Member States in terms of health.

The Directive clarifies the rights of citizens to access safe and good quality treatment across the EU and its reimbursement. Europeans prefer to receive healthcare close to home: no one wants to travel further than they should when they are ill. However, sometimes people have to go abroad, because experience or the medical care they need is not available within the national borders. Or simply because the nearest hospital is across the border.

However, from the application of the provisions of the Directive are exempted certain health care services such as, for example, long-term services, whose purpose is to support people who need help with daily routine tasks.

For OECD, long-term care is "a political issue of confluence, which brings together a range of services for people who are dependent on help in basic activities of daily living over an extended period of time". National definitions on long-term care vary within the European Union, and reflect the differences in the length of stay, range of beneficiaries and the often unclear boundary between health (health care) services and non-medical (social) services. Some countries prefer, for example, to focus on early rehabilitation outpatient treatment, while others focus more on providing care in hospitals or similar institutions. Long-term care can include rehabilitation, basic medical treatment, home health care, social care, housing and services such as transportation, food, occupational assistance and help in managing daily activities [6].

In the field of cross-border medical services, a certain overlap is somehow reached between the law of the Union and the national law, so that European law in many cases is essentially limited to indicating a binding objective, i.e. the achievement of the free movement of citizens patients and their equal treatment, regardless of nationality, in relation to national authorities, while preserving the competence of the member states. Directive 2011/24/EU preserves the competences of the Member States, which are obliged to submit to the regulatory framework imposed by primary law and secondary legislation, to the extent that Member States must not violate EU law when exercising their powers.
As argued in the Watts judgment, Member States are obliged to adapt their national healthcare and social security systems [7]. Moreover, the Court emphasized since its previous decisions that Member States must comply with EU law, in particular with the provisions on the freedom to provide services [8].

Those provisions prohibit Member States from introducing or maintaining unjustified restrictions on the freedom to provide medical care services [9].

In addition, the Court case law expressly emphasized that the mandatory adaptations of national social security systems aiming to achieve the fundamental freedoms guaranteed by the Treaty should not be considered by Member States as interference in their sovereign competence in the field of public health [10].

We believe that should not remain unmentioned the fact that the European Union can exert considerable influence on the health systems of Member States, for example, by measures designed to achieve fundamental freedoms [11].

The Directive is without prejudice to the laws, regulations and administrative provisions of the Member States relating to the organization and financing of healthcare in situations not related to cross-border healthcare. In particular, nothing in this Directive obliges a Member State to reimburse the costs of healthcare provided by healthcare providers established on its territory if those providers are not part of the social security system or national health system of that State Member State.

From the interpretation of the text, in conjunction with Article 3, paragraph 1 letters (a) - (c) of Directive 2005/36/EC, follows that it does not matter whether the work performed by a qualified person (as is the sanitary field, such as the analysed case) has a temporary or occasional basis. As the promotion of the provision of services must be ensured in the context of the strict compliance with public health and safety and the protection of the consumer, Member States have special provisions in the national legislation for professions regulated at sector level with implications in terms of health.

Given the different systems established on the one hand, for the provision of temporary and occasional cross-border services and, on the other hand, for establishment, it is necessary to specify criteria for distinguishing between these two concepts in the case of the movement of the service provider on the territory of the host Member State.
THE PROTECTION OF PERSONAL DATA IN THE CASE OF PUBLIC HEALTH

The fundamental right to privacy with regard to the processing of personal data is protected in conformity with Member States' national measures for implementing Union provisions on the protection of personal data, in particular Directives 95/46/EC [12] and 2002/58/EC [13].

To analyse how these rules apply to public health in general and, in particular, on cross-border healthcare, we shall refer to the Romanian legislation relating to the activity of personal data processing and the free movement of such data, Law no. 677/2001 [14].

Thus, the processing of personal data related to racial or ethnic origin, political, religious, philosophical or similar nature opinions, the union membership, as well as personal data concerning health or sex life is prohibited. This provision shall not apply where the subject has given their express consent to such processing.

Regarding the prior express consent we believe that, regarding healthcare, the mere presentation of the patient to a health service provider, amounts to a tacit consent, so we cannot discuss express consent. It is inevitable that the supplier request personal data, even for an appointment for diagnosis, (name, address, telephone number, affection suspected or confirmed by someone else etc.).

The National Law, Law no. 677/2001, provides for special rules on the processing of personal data concerning health. According to this regulation, healthcare professionals, medical care institutions and their staff may process personal data on health status, without the authorization of the supervisory authority, only if the processing is necessary to protect the life, physical integrity or health of the concerned person.

To detail how personal data concerning health can be processed by service providers, the law provides that this operation can be performed only by a health professional or under its supervision, subject to professional secrecy.

We believe that the competent national authorities in public health should regulate more differentiated all aspects of the patient's right to confidentiality, to reduce the risks of disclosing personal data on the health of citizens. In our opinion and in the
absence of these legislative differentiations, information that normally would not be provided to the public appears in the mass media.

CONCLUSIONS

Within health systems throughout the European Union there are a number of common principles of operation [15], which have been affirmed by the case law of the Court of Justice of the European Union. These principles must be applied uniformly in national health systems, both to strengthen the confidence of patients in cross-border healthcare, a prerequisite for achieving patient mobility, and to ensure a high level of health protection.

Referring to the decisions of national authorities on market mechanisms and the pressure of competition to manage health systems, the Council was of the view that decisions about the health care package which citizens are entitled to and the respective mechanisms used to finance and provide healthcare, must be placed in the national context of the Member States.

Under the Treaty on the Functioning of the European Union [16], at the basis of all European policies lies the aim to ensure a high level of human health protection, a major goal of the whole Union. This goal is also considered when the European legislator adopts acts under other Treaty provisions.

From the case law of the Court of Justice of the European Union unequivocally results that people normally resident in a Member State operating a national health service, are entitled to receiving hospital treatment in another Member State at the expense of the national health service.

Member States may condition this right by the requirement that the person concerned should have obtained prior authorization, only if such authorization is based on objective, non-discriminatory and transparent criteria within a procedure system. In addition, applications for the authorization of treatment abroad must be analysed objectively and impartially, within a reasonable time, and the national health authority's refusal to grant such authorization can be challenged in court or out of court. The absence of such criteria and the lack of easily accessible and transparent procedures cannot deprive a person of this right. Also, if the conditions for authorization (form E112) are designed to safeguard the financial stability of the national health system,
considerations of a purely budgetary or economic nature not being able to justify the refusal to grant such authorization.

To determine whether the treatment is available without undue delay might be considered the waiting time and the priority to treatment granted by the national health authority, only on condition that they are based on concrete indications relating to the patient’s condition at the time of evaluation, as well as its medical history and the prognosis for the patient seeking treatment.

Under European law, the affiliate Member State is obliged to fund the hospital treatment carried out in another Member State and the reimbursement of this treatment is based on national legislation. In the absence of tariffs or rates for calculating the amount of reimbursement, the reimbursement must be calculated at the actual cost of the treatment. Travel and accommodation costs related to hospital treatment received in another Member State are reimbursable only where this is provided for by national law for treatment on national territory.

Regarding the obligation of a Member State to reimburse the cost of hospital treatment provided in another Member State of the European Union, Article 49 EC does not allow to take into account budgetary reasons, unless it is demonstrated that compliance with this obligation on a more general scale would threaten the financial balance of the respective national health system. Moreover, in accordance with Article 22 (2) of Regulation EEC No. 1408/71 [22], budgetary considerations cannot be taken into account in decisions refusing prior authorization for treatment abroad.

References
[7] Judgment of the Court (Grand Chamber) of 16 May 2006. The Queen, on the application of Yvonne Watts v Bedford Primary Care Trust and Secretary of State for Health. Reference for a preliminary ruling: Court of Appeal (England & Wales) (Civil Division) - United Kingdom. Social security - National health
system funded by the State - Medical expenses incurred in another Member State - Articles 48 EC to 50 EC and 152(5) EC - Article 22 of Regulation (EEC) No 1408/71. Case C-372/04 Watts, paragraph 147.  
[10] This idea is clearly expressed in Judgment Commission/Luxembourg (C-490/09, Rec., 2011, p. I-247, paragraph 32)  
[14] Law no. 677/2001 on the protection of persons with regard to the processing of personal data and the free movement of such data, published in the Official Monitor of Romania, Part I, no. of 12 December 2001, with the subsequent modifications and completions.  