

The Cross-border Healthcare Directive: The 2018 Implementation Report



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Overview

- The Cross-border Healthcare Directive
- Main messages of the 2018 Implementation Report to the European Parliament and the Council
- Core conclusions of the Report
- Reception by inter-institutional partners and the media

The Directive 2011/24/EU



 A major change in the EU's involvement in health policy



Main aims of this Directive

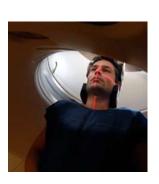
To help patients exercising their rights for healthcare in another EU country.

Therefore the Directive clarifies:

- 1. Information to patients;
- 2. Rules of reimbursement;
- 3. Procedural guarantees;









Triennial Commission report on the operation of the Directive 2011/24/EU on the application of patients' rights in cross-border healthcare

- 2018 Report published and submitted to the European Parliament and the Council on 21 September 2018:
- I. State of play of transposition
- II. Patient mobility
- **III.** Information to patients and National Contact Points
- IV. Cooperation between health systems
- **V.** Conclusions



I. State of play of transposition

Transposition check:

- Completeness check finished
 - 26 infringements launched (+ 21 for Implementing Directive 2012/52/EU)
- Compliance check ongoing

Issues identified:

- Systems of reimbursement (unreasonably low reimbursement tariffs or restriction on reimbursement);
- Use of prior authorisation (lack of transparency or incorrect use of PA);
- 3) Unreasonable administrative requirements;
- 4) Charging of incoming patients.



I. State of play of transposition

- Based on the systematic assessment of all notified measures by all Member States, 11 own-initiative investigations gathering information were launched:
 - 4 structured dialogues have been closed already since Member States changed their legislation;
 - 1 infringement is almost at the level of referral to the next instance;
- Overall, this work strand confirmed that solutions can be found for the benefit of EU citizens through structured bilateral dialogues.

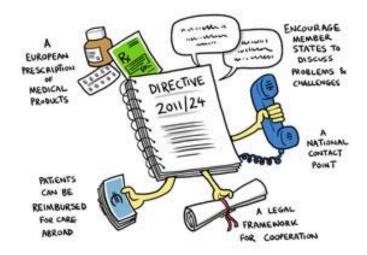


Photo credit: https://drawnalism.com



I. i) Systems of reimbursement of costs

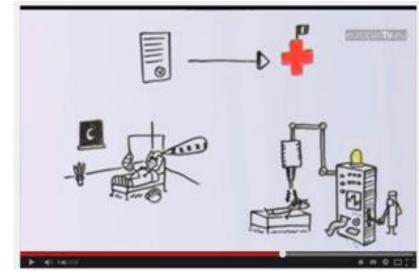
- Reimbursement tariffs based on cost of treatment at home from public / contracted provider;
- ➤ No specific notifications received under Article 7(9), allowing Member States to limit application of the rules on reimbursement of cross-border healthcare for overriding reasons of general interest.



I. ii) Prior authorisation

Prior authorisation possible for

- a) overnight stay; or
- b) highly specialised and cost-intensive healthcare



- Presently, 6 MSs and Norway have no prior autorisation system in place at all;
- If prior autorisation is considered necessary, a detailed and sufficiently defined shortlist should be publically available.



I. iii) Administrative procedures regarding cross-border healthcare

- Administrative procedures for cross-border reimbursement are based on objective, nondiscriminatory criteria which are necessary to the objective to be achieved;
- The 2018 Report offers examples of administrative procedures that were lifted in the interest of patients following discussions with the Member States on the proportionality and necessity thereof;
- The prior notification option under Art 9(5): a mechanism worth upscaling.

Health



I. iv) Fees for patients from other Member States

- Non-discrimination of patients from other Member States with respect to access and *pricing*;
- Same scale of fees to patients from other Member States as for domestic patients in a comparable medical situation;
- If no comparable price for domestic patients, obligation on providers to charge a price calculated according to objective, non-discriminatory criteria;
- The establishment of a cost-based pricing system may well have implications for reimbursement obligations of Member States to outgoing patients.

Health



II. Key figures on patient mobility

1. Coordination on social security schemes

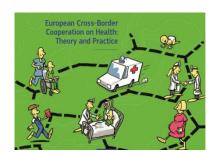
- Necessary (unplanned) healthcare: ±2 million cases/year;
- Planned healthcare: ±55,000 PA/year;
- Living outside of the competent MS: ± 1.4 million people;
 - → 0.1% of the EU-wide annual healthcare budget

2. <u>Directive 2011/24/EU</u>

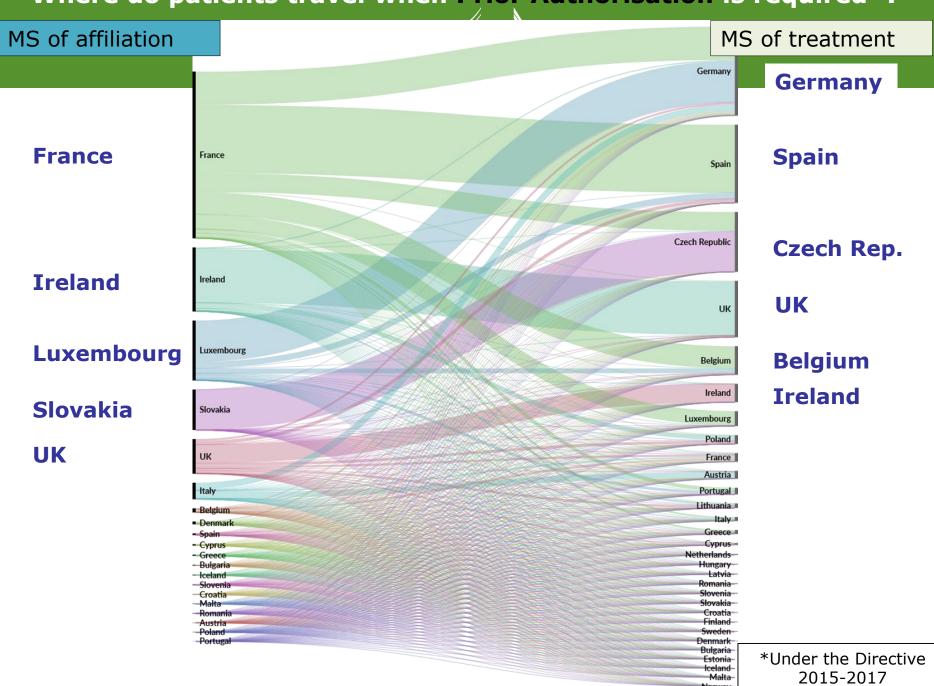
- CB healthcare without prior authorisation: ±200,000 reimbursement/year
- CB healthcare with prior authorisation: ±3500 PA/year
 - → 0.004% of the EU-wide annual healthcare budget
- 3. Bilateral agreements for cross-border healthcare
 - No data available



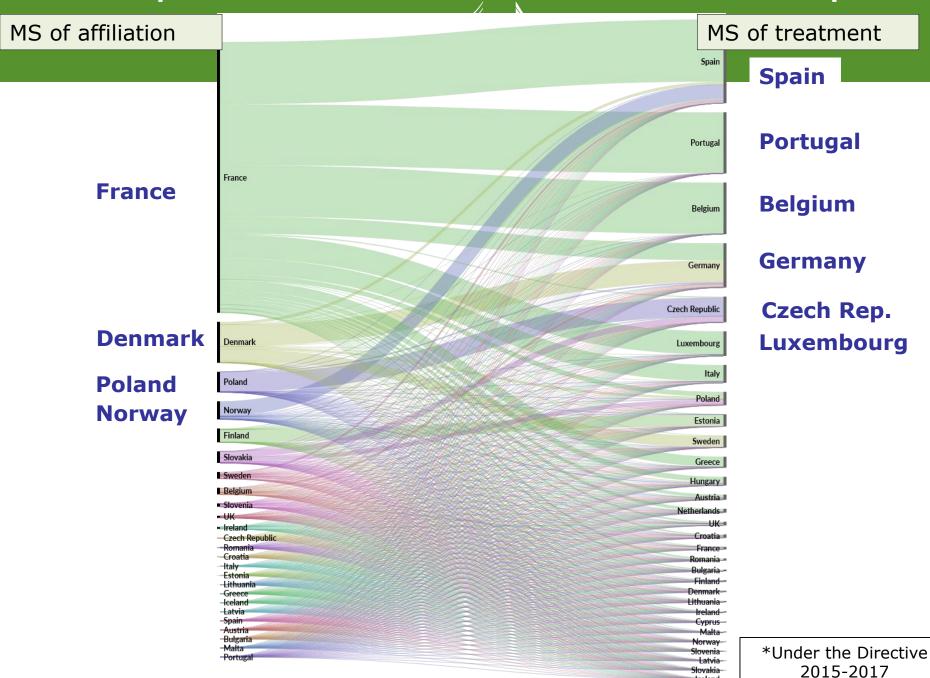




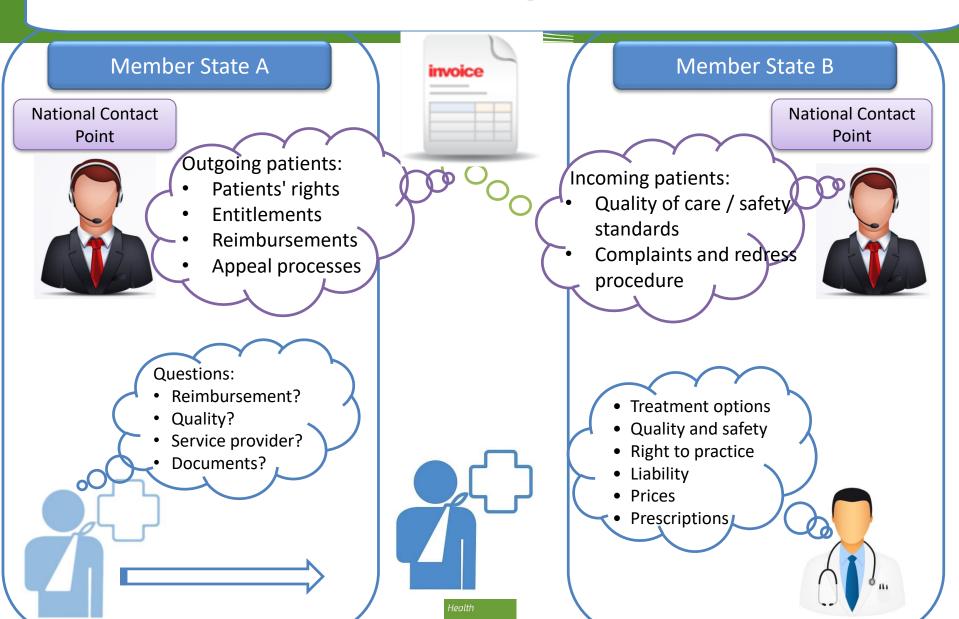
Where do patients travel when Prior Authorisation is required*?



Where do patients travel when Prior Authorisation is not required?

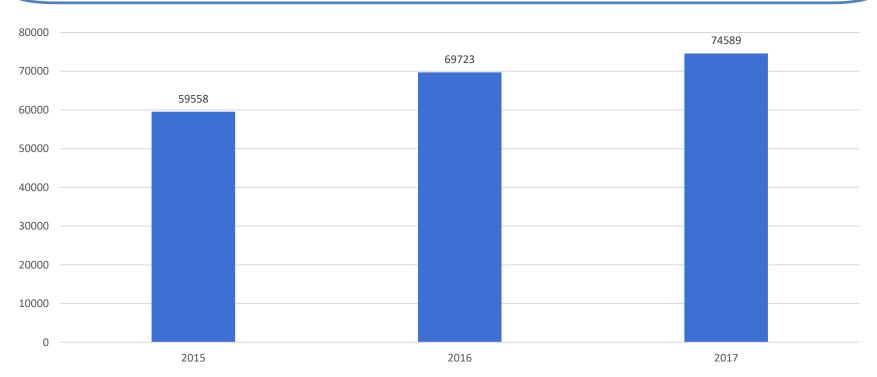


III. Information to patients and NCPs





III. Requests for information made to NCPs – a slow yet steady increase, due to websites, doctors?



Patients have right to receive healthcare abroad (Directive 2011/24/EU)

- ✓ Main rule: No prior authorisation (overnight stay and highly specialised and cost intensive care);
- ✓ Direct payment to providers;
- Reimbursement based on tariffs and rights in the MS of affiliation
- ✓ Public / private providers and medicines are available;

Coordination of social security schemes (Regulation (EC) No 883/2004)

Necessary treatment

✓ Medically necessary care;

<u>Planned treatment</u>

 ✓ Prior authorization in case of undue delay;



- ✓ Reimbursement between institutions based on the tariffs of treatment, (No co-payment);
- ✓ Public (contracted) providers only





IV. Cooperation between Health Systems

- the ERNs
- European Reference Networks
- eHealth
- Health Technology Assessment
- Encouraging cooperation between MS to improve complementarity of their health systems in border regions – priority for the EU



Commission Communication on growth and cohesion in EU border regions September 2017



Conference on Enhancing Healthcare Cooperation in Cross-Border Regions 4th December 2018 — Brussels, Centre de Conférences Albert Borschette



Conclusions

- Patients' mobility shows a slight increasing trend;
- Information provided by the NCPs has been enhanced over the reporting period + websites have been improved;
- The Directive has proven to clarify and guarantee patients' rights to receive healthcare in another MS;
- Voluntary cooperation between health systems developed further – framework and momentum provided by the Directive (HTA, eHealth, ERN);
- The Directive has not resulted in a major budgetary impact on the sustainability of national health systems.



Reception by media and interinstitutional partners thus far

- EP IMCO Committee Opinion favourable;
- EP ENVI Committee Draft Report for a Motion for a Resolution vote in Plenary planned for February 2019;
 - Calls on MSs to provide sufficient funding for their NCPs to be able to develop comprehensive information;
 - Recommends that the Commission develops guidelines on the functioning of NCPs.
- Awaiting possible Council uptake next year;
- Awaiting Court of Auditors Performance Audit Q2 2019;
- Overall positive reaction from stakeholders and the media.

Thank you for your attention!

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Health and Food Safety Directorate-General
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