### Patient experience with ambulatory care

Delivering health care that is responsive and patient-centered is playing a greater role in health care policy across OECD countries. Measuring and monitoring patient experience empowers patients and the public, involves them in decisions on health care delivery and governance, and provides insight into the extent to which they are health-literate and have control over the treatment they receive. Across countries, using the health care user as a direct source of information is becoming more prevalent for health system monitoring, planning and decision making, and efforts to measure and monitor patient experiences have actually led to health care quality improvements (Fujisawa and Klazinga, forthcoming).

Since the mid-1990s, there have been efforts to institutionalise measurement and monitoring of patient experiences. In many countries, responsible organisations have been established or existing institutions have been taking charge of measuring and reporting patient experiences. They developed survey instruments for regular collection of patient experience data and standardised procedures for analysis and reporting. An increasing number of countries collect not only Patient-Reported Experience Measures (PREMs) but also Patient-Reported Outcome Measures (PROMs) which collect patients' perception on their specific medical conditions and general health, including mobility, pain/discomfort and anxiety/depression, before and after a specific medical intervention such as hip and knee replacement.

A growing number of countries are using patient-reported data to drive quality improvements in health systems. Patient experience data are reported in periodic national health system reports or on public websites, showing differences across providers, regions and over time. Korea, Norway, Sweden and the United Kingdom use patient experience measures in payment mechanisms or for fund allocations to promote quality improvement and patientcentred care, and Australia, Canada, the Czech Republic, Denmark and France use them to inform health care regulators for inspection, regulation and/or accreditation. Patient-reported measures are also used in some Canadian jurisdictions, Denmark, France and the Netherlands to provide specific feedback for providers' quality improvement. In England, PROMs and patients' feedback about their experience are used to inform patient choice and to incentivise service improvement. For example, PROMs data for patients undergoing some procedures such as hip and knee replacement are used for benchmarking hospitals. The use of PROMs can also enable the potential shift from a volume-based to a value-based model of health system resource management (Canadian Institute for Health Information, 2015).

Patients generally report positive experiences when it comes to communication and autonomy in the ambulatory health care system. Across countries, the majority of patients report positive experiences with regards to time spent with the doctor (Figure 8.39), easy-to-understand explanations (Figure 8.40), opportunities to ask questions or raise concerns (Figure 8.41), as well as involvement in care and treatment decisions (Figure 8.42). For all four aspects of patient experience, Belgium and Luxembourg score high at above 95% of patients reporting positive experiences. Poland has lower rates with fewer than one in two patients reporting having been given the opportunity to ask questions or been involved in their care and treatment during consultation. The proportion of patients with positive experience has decreased since 2010 in Australia, France, the Netherlands and Switzerland, but countries with lower rates such as Sweden and Poland have improved some aspects of patient experiences in recent years (Commonwealth Fund, 2010).

#### Definition and comparability

In order to measure and monitor general patient experience in the health system, the OECD recommends collecting data on patient experience with any doctor in ambulatory settings. An increasing number of countries have been collecting patient experience data based on this recommendation through nationally representative population surveys while Japan and Portugal collect them through nationally-representative service user surveys. Some countries, however, collect data on patient experience with a regular doctor. For about half the countries presented, the Commonwealth Fund's International Health Policy Surveys 2010 and 2013 were used, even though there are critiques relating to the sample size and response rates. Data from this survey refer to patient experience with a regular doctor rather than any doctor.

Rates are age-sex standardised to the 2010 OECD population, to remove the effect of different population structures across countries.

#### References

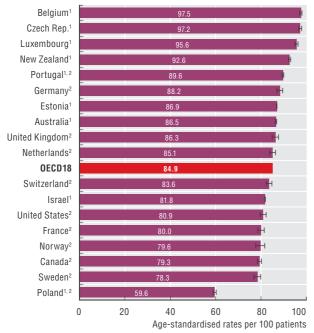
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Fujisawa, R. and N. Klazinga (forthcoming), "Measuring Patient Experiences (PREMs): Progress Made by the OECD and its Member Countries 2006-2015", OECD Health Working Papers, Paris.

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## 8.39. Doctor spending enough time with patient in consultation, 2013 (or nearest year)

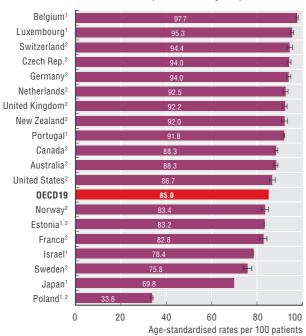


Note: 95% confidence intervals represented by H.

 National sources. 2. Data refer to patient experiences with regular doctor. Source: Commonwealth Fund International Health Policy Survey 2013 and other national sources.

StatLink http://dx.doi.org/10.1787/888933281241

## 8.41. Doctor giving opportunity to ask questions or raise concerns, 2013 (or nearest year)



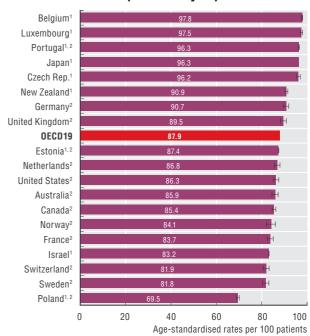
Note: 95% confidence intervals represented by H.

 National sources. 2. Data refer to patient experiences with regular doctor. Source: Commonwealth Fund International Health Policy Survey 2010 and other national sources.

StatLink http://dx.doi.org/10.1787/888933281241

Information on data for Israel: http://oe.cd/israel-disclaimer

## 8.40. Doctor providing easy-to-understand explanations, 2013 (or nearest year)

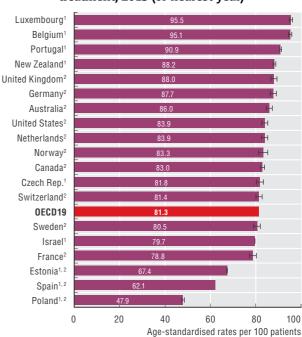


Note: 95% confidence intervals represented by H.

1. National sources. 2. Data refer to patient experiences with regular doctor. Source: Commonwealth Fund International Health Policy Survey 2013 and other national sources.

StatLink http://dx.doi.org/10.1787/888933281241

## 8.42. Doctor involving patient in decisions about care and treatment, 2013 (or nearest year)



Note: 95% confidence intervals represented by H.

1. National sources. 2. Data refer to patient experiences with regular doctor. Source: Commonwealth Fund International Health Policy Survey 2013 and other national sources.

StatLink http://dx.doi.org/10.1787/888933281241



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