

Commission Implementing Directive 2012/52/EU of 20 December
2012 laying down measures to facilitate the recognition of medical
prescriptions issued in another Member State (Text with EEA relevance)

Article 1	Subject matter
Article 2	Scope
Article 3	Content of prescriptions
Article 4	Information requirements
Article 5	Transposition
Article 6	Entry into force
Article 7	Addressees
	Signature

ANNEX

Non-exhaustive list of elements to be included in medical prescriptions

Headings appearing in bold in this Annex are not required...

Identification of the patient

Authentication of the prescription

Identification of the prescribing health professional

Identification of the prescribed product, where applicable

***Status:** EU Directives are being published on this site to aid cross referencing from UK legislation. After IP completion day (31 December 2020 11pm) no further amendments will be applied to this version.*

- (1) [OJ L 88, 4.4.2011, p. 45–65](#)
- (2) [OJ L 311, 28.11.2001, p. 67.](#)