

Commission Delegated Directive 2014/15/EU of 18 October 2013 amending, for the purposes of adapting to technical progress, the Annex IV to Directive 2011/65/EU of the European Parliament and of the Council as regards an exemption for lead, cadmium and hexavalent chromium in reused spare parts, recovered from medical devices placed on the market before 22 July 2014 and used in category 8 equipment placed on the market before 22 July 2021, provided that reuse takes place in auditable closed-loop business-to-business return systems, and that the reuse of parts is notified to the consumer (Text with EEA relevance)

- Article 1 Annex IV to Directive 2011/65/EU is amended as set out...
Article 2 (1) Member States shall bring into force the laws, regulations...
Article 3 This Directive shall enter into force on the twentieth day...
Article 4 This Directive is addressed to the Member States.
Signature

ANNEX

In Annex IV to Directive 2011/65/EU the following point 31...
Lead, cadmium and hexavalent chromium in reused spare parts,
recovered...

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 174, 1.7.2011, p. 88.](#)