

Commission Implementing Decision (EU) 2020/1807 of 27 November 2020 concerning the extension of the action taken by the United Kingdom Health and Safety Executive permitting the making available on the market and use of the biocidal product Biobor JF in accordance with Article 55(1) of Regulation (EU) No 528/2012 of the European Parliament and of the Council (notified under document C(2020) 8158) (Only the English text is authentic)

- Article 1 The United Kingdom Health and Safety Executive may extend the...
- Article 2 This Decision is addressed to the United Kingdom Health and...
Signature

Changes to legislation: There are currently no known outstanding effects for the
Commission Implementing Decision (EU) 2020/1807. (See end of Document for details)

- (1) [OJ L 167, 27.6.2012, p. 1.](#)
- (2) [OJ L 29, 31.1.2020, p. 7.](#)
- (3) Annex II to Commission Delegated Regulation (EU) No 1062/2014 of 4 August 2014 on the work programme for the systematic examination of all existing active substances contained in biocidal products referred to in Regulation (EU) No 528/2012 of the European Parliament and of the Council ([OJ L 294, 10.10.2014, p. 1.](#)).

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There are currently no known outstanding effects for the Commission Implementing Decision (EU) 2020/1807.