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amending Regulation (EC) No 1907/2006 of the European Parlia-
ment and of the Council on the Registration, Evaluation, Authorisa-
tion and...~~

Without prejudice to Article 4 of Directive 98/24/EC, a chemical safety assessment shall be performed and a chemical safety report completed for all substances subject to registration in quantities of 10 tonnes or more per year per registrant. A chemical safety assessment of a substance shall include the following steps:

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~~REACH Legislation - ECHA~~

~~Evaluation of Regulation (EC) No 1107/2009 on the placing ...~~

~~REGULATION (EC) No 1907/2006 OF THE EUROPEAN PARLIAMENT
28 European Commission. No data on emergency authorisations
pre-2007. 29 Ecorys (2018), Study supporting the REFIT Evaluation
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cide residues (Regulation (EC) No 1107/2009 and Regulation (EC)
No 396/2005). 30 Mattaar, H. (2010).~~

Evaluation of Regulation (EC) No 1831/2003 FEFANA welcomed the launch of the EU evaluation of the Feed Additives (FAs) Regulation (read more) and is determined to provide its full support and expertise for the proper development of this process.

Reg. (EC) No 1272/2008. Current legislation. Registration, Evaluation, Authorisation and Restriction of Chemicals (REACH) is a European Union regulation dating from 18 December 2006. REACH addresses the production and use of chemical substances, and their potential impacts on both human health and the environment.

~~Commission Regulation (EU) 2020/878 of 18 June 2020 ...~~

~~Regulation (EC) No 1907/2006 of the European Parliament and of
the Council of 18 December 2006 concerning the Registration,
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~~Evaluation of Regulation (EC) No 1831/2003 - FEFANA~~

~~Regulation (EC) No 1907/2006 - Registration, Evaluation ...~~

~~Regulation (EC) No 1907/2006 of the European Parliament and of
the Council of 18 December 2006 concerning the Registration,
Evaluation, Authorisation and Restriction of Chemicals (REACH),
establishing a European Chemicals Agency, amending Directive
1999/45/EC and repealing Council Regulation (EEC) No 793/93
and Commission Regulation (EC) No 1488/94 as well as Council Di-
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93/67/EEC, 93/105/EC and 2000/21/EC.~~

~~Regulation (EC) No 1333/2008 requires the Commission to set up
a programme for the re-evaluation, by the European Food Safety
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additives that were already permitted in the Union before 20 Janu-
ary 2009. (2)~~

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~~EUROPEAN COMMISSION Brussels, 9.7.2019 SWD(2019) 295 final
COMMISSION STAFF WORKING DOCUMENT EVALUATION of the
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Council of 12 December 2006 on medicinal products for paediatric
use and Regulation (EC) No 141/2000 of the European Parliament
and of the Council of 16 December 1999 on orphan medicinal
products~~

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of the European Parliament and Council on additives for use in animal nutrition.

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