Data to be submitted jointly or separately for registration under REACH

Joint submission	Separate submission	Joint or separate submission: free decision
Classification and labelling of the substance.	Identity of the manufacturer or the importer of the substance.	Guidance of safe use of the substance.
Study summaries[1] of the information derived from the application of annexes VII to XI.	Identity of the substance.	Chemical Safety Report (CSR), when required.
Robust study summaries[2] of the information derived from the application of annexes VII to XI, for all key data used in the hazard assessment.	Information on the manufacture and use(s) of the substance. This information shall represent all the registrant's identified use(s).	
Indication as to which of the above information has been reviewed by an assessor chosen by the manufacturers or importers and having appropriate experience.	Indication as to which of the information on the manufacture and use(s) has been reviewed by an assessor chosen by the manufacturer or importer and having appropriate experience.	Indication as to which of the information given in the CSR has been reviewed by an assessor chosen by the manufacturer(s) or importer(s) and having appropriate experience.
Proposals for testing where listed in annexes IX and X.	For substances in quantities of 1 to 10 tonnes, exposure information.	

1. Study summary: a summary of the objectives, methods, results and conclusions of a full study report providing sufficient information to make an assessment of the relevance of the study.

2. Robust study summary: a detailed summary of the objectives, methods, results and conclusions of a full study report providing sufficient information to make an independent assessment of the study minimising the need to consult the full study report.