

Position Paper

RoHS Pack 15
Stakeholder Consultation –
Position Paper Substance Methodology



1 **Preliminary remarks**

2 Thank you for the possibility to provide comments on the draft substance methodology
3 manual and the substance inventory.

4 We welcome the decision, in contrast to the original timetable, to carry out the consul-
5 tation on substance methodology first and then, based on its outcome, to assess the
6 seven substances already prioritized.

7 In future, for consultations with high volume documents we would appreciate to get
8 more answering time. This offers the opportunity to identify all relevant concerns and
9 to support the argumentation by providing meaningful examples. In order to involve as
10 many stakeholders as possible, we generally recommend that consultations start best
11 at the beginning of the week and end at times when availability is expected to be high.
12 We would like to express the following concerns and improvement proposal.

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15 **Substance assessment**

16 In RoHS Article 6 (1) it is clearly stated that REACH is an essential basis for prioritizing
17 new substances: "The review and amendment of the list of restricted substances in
18 Annex II shall be coherent with other legislation related to chemicals, in particular Reg-
19 ulation (EC) No 1907/2006, and shall take into account, inter alia, Annexes XIV and
20 XVII to that Regulation."

21 In addition to the comprehensive substance knowledge provided from the registration
22 dossiers available at ECHA, RoHS Article 6(1)(a)-(d) defines the crucial criteria for in-
23 cluding a substance in a pre-assessment with special regards on waste treatment. Spe-
24 cial attention should therefore be paid to this aspect.

25 The prioritization of substances under RoHS should focus on actually occurring, un-
26 controllable hazards in the treatment of WEEE that exist despite the application of all
27 available occupational health and safety measures.

28 After all the argument that hazardous substances might be released to the environ-
29 ment, through wrong or illegal treatment of WEEE it is in our view not a criterion to be
30 considered for the evaluation of new substances in EEE.

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33 **Grouping of substances**

34 The grouping for the evaluation of substances should be carried out and reviewed on
35 a case-by-case basis as there are no common rules for the grouping of substances.
36 Substances show different intrinsic properties despite similar molecular structures.
37 Concerns about a substance are an individual characteristic that may differ within a
38 chemical class. When a grouping is performed, it should be evaluated according to the
39 effects of substances, e.g. (eco-)toxicological effects, hazard classification or toxicoki-
40 netic.

41 The statement in the draft on page 17 about chemical structure and characteristics
42 within a substance group is scientifically incorrect, e.g. enantiomers are stereo isomers
43 with identical chemical composition and structure, they just differ in their configuration.
44 Their physical properties are identical, but their biological effects may be completely
45 different, e.g. Thalidomide. Furthermore, a group of phthalates includes short chain
46 phthalate e.g. DEHP, repr. 1B and long-chained phthalate, e.g. DINP not classified.

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49 **Assessment of substitutes**

50 The assessment of possible substitutes for materials in electrical and electronic equip-
51 ment should focus on their benefits in relation to criteria (a) to (c) referred to in Article
52 6(1) and should follow the principle of RoHS Article 5(1)(a) for adapting of Annexes III
53 and IV. Potential substitutes must be evaluated according to identical criteria to the
54 same extent as the substances to be substituted. Otherwise, unknowingly higher risks
55 might be accepted, see also Rec. 18 of the RoHS.

56 Developing the methodology, it should be considered that a replacement of a sub-
57 stance or material needs intensive technically testing for functionality and durability for
58 each specific application. Thus, substitution is often time consuming and expensive.
59 This poses a particular threat to SMEs and leads to undesirable market concentration.

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62 **Socioeconomic Analysis**

63 A socio-economic assessment should be carried out according to clear, established
64 rules and a scientific approach which is in line with the REACH Regulation and is es-
65 tablished/applied by the SEAC.

66 In the event of a lack of data, rules and/or measures for the collection of valid data must
67 be defined. Example "Impact on waste management": For example, if reliable data from
68 waste treatment are not available, studies involving stakeholders may need to be com-
69 missioned before a substance is proposed for inclusion in Annex II. This also applies
70 to other categories. If no substantiated data are available, it must be proved which
71 effects can be expected without restriction of the substance.

72 The balancing of the potential benefits of a restriction against the costs is not to be
73 understood exclusively in monetary terms. Rather, the sum of all negative effects
74 should be considered. Example "Impact on manufacturers": Impact on energy effi-
75 ciency, sustainability, impact on product safety, restrictions on innovation, reduced
76 competitiveness with non-EU countries.

77 Also, the benefits must be quantified using clearly defined criteria, so that a calculation
78 of the cost-benefit ratio becomes transparent and comparable.

79 The entire process of socio-economic evaluation should include technical and eco-
80 nomic feasibility for small and medium-sized enterprises (SMEs) (RoHS 2011/65/EU
81 Rec. 8).

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84 **Burden of proof**

85 The CE procedure must comply with standard EN 50581 (IEC 63000). Experience
86 shows that the market expects a higher verification depth than originally prescribed by
87 the standard. Therefore, a corresponding analysis effort is generated for each addi-
88 tional regulated substance. Even if the use of the substances is not to be expected.

89 Impact assessment per additional substance for an average company:

- 90 • Effort for communication in the supply chain with approx. 20 working days / com-
91 pany including maintenance of product documentation.
- 92 • Analysis e.g. as part of the initial sampling with a cost expenditure of approx. 2.5
93 million Euro / Company
- 94 • Resource requirements, e.g. energy and solvents, within the framework of the
95 analysis at the test institutes

96 The largest part of the supply chain is composed of SMEs (ZVEI members: > 80 %),
97 with a significant higher burden on the verification process in relation to other compa-
98 nies.

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101 **Inventory**

102 The objective of the review of Annex II is to prioritize future substances of potential
103 concern, in particular with regard to their effects on the waste treatment process (Article
104 6.1) and the amount of their use in EEE. For this purpose, the identification of each
105 substance or material used in EEE is irrelevant. Therefore, a survey on the use of sub-
106 stances is inefficient and ineffective regardless of their (potential) concerns, e.g. see
107 required information on palladium, platinum, copper, citric acid, magnesium carbonate,
108 EVAC etc.

109 The inventory to be compiled should be limited to those substances whose risk has
110 already been identified in coherent legislation, Rec. 10 and 16. The inventory should
111 focus on substances which remain in the final product and have been identified to pose
112 a potential risk in the recycling or waste phase of the product.

113 Therefore, the given inventory information is related to other regulated substances
114 (REACH Annex XVII and Candidate List, POP V), already examined by the IEC 62474
115 Validation Team. Regarding the quantities used for EEE, there are extensive data avail-
116 able from the registration process at the ECHA.

117 Using the registration data guarantees to avoid wrong interpretation because collecting
118 information of different supply chain members causes multiplying of amounts.
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138 **About ZVEI**

139 The ZVEI - German Electrical and Electronic Manufacturers' Association promotes the industry's joint economic, tech-
140 nological and environmental policy interests on a national, European and global level. The ZVEI represents more than
141 1,600 companies, mostly SMEs.
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143 The industry has round about 868,000 employees in Germany plus 736,000 employees all over the world. In 2017 the
144 turnover was Euro 191 billion.
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146 The electrical and electronics industry is the most innovative industry sector in Germany. One-third of the industry's
147 sales are based on new products. The industry spends Euro 17.2 billion in R&D every year, Euro 6.1 billion in invest-
148 ments and Euro 2 billion in training and further training. Every third innovation in Germany's manufacturing sector
149 stems from solutions of this industry.

150	Annex 1 - Consultation questionnaire
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152	1. Please specify additional lists of relevance for specifying substances identified or
153	suspected of having hazardous properties. (Section 1.1., pg. 24)
154	
155	See RoHS Pack 15 Stakeholder Consultation – ZVEI Position Paper Substance
156	Methodology lines 15 to 30 (substance assessment)
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158	2. Please specify additional lists of relevance for specifying substances used or sus-
159	pected of being used in EEE. (Section 1.2., pg. 27)
160	
161	See RoHS Pack 15 Stakeholder Consultation – ZVEI Position Paper Substance
162	Methodology lines 15 to 30 (substance assessment)
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164	6. For the purpose of specifying an exhaustive list of socio-economic impacts to be
165	considered, please specify categories that should be taken into consideration. (Sec-
166	tion 3., pg. 43)
167	
168	See RoHS Pack 15 Stakeholder Consultation – ZVEI Position Paper Substance
169	Methodology lines 62 to 81 (socioeconomic analysis)
170	
171	7. If you have any further comments, where relevant, please note the section/page to
172	which they refer or quote the text of relevance from the manual to support under-
173	standing.
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175	See RoHS Pack 15 Stakeholder Consultation – ZVEI Position Paper Substance
176	Methodology:
177	• lines 33 to 46 (grouping of substances)
178	• lines 49 to 59 (assessment of substitutes)
179	• lines 84 to 98 (burden of proof)
180	• lines 101 to 118 (inventory)