UN/SCEGHS/28/INF.29

Committee of Experts on the Transport of Dangerous Goods and on the Globally Harmonized System of Classification and Labelling of Chemicals

4 December 2014

Sub-Committee of Experts on the Transport of Dangerous Goods

Forty-sixth session

Geneva,1–9 December 2014

Item 8 (g) of the provisional agenda

Issues relating to the Globally Harmonized System of Classification and Labelling of Chemicals: corrosivity criteria

Sub-Committee of Experts on the Globally Harmonized System of Classification and Labelling of Chemicals

Twenty-eighth session

Geneva, 10–12 (a.m.) December 2014 Item 2 (d) of the provisional agenda

Classification criteria and related hazard communication: Work of the TDG-GHS working group on corrosivity criteria

Amended proposal for revision of Chapter 2.8 of the Model Regulations

Transmitted by the expert of the Netherlands and CEFIC

Introduction

- 1. Since the Netherlands presented proposal ST/SG/AC.10/C.3/2014/69 ST/SG/AC.10/C.4/2014/12, the United Kingdom, the United States of America, Canada and CEFIC have also presented modifications or different proposals for a new Chapter 2.8 with the aim to align the Model Regulations with GHS, retaining the level of safety and not promoting the use of animal testing.
- 2. Based on the papers submitted by the Netherlands and CEFIC for this session, a joint proposal has been developed which is presented in this current paper.
- 3. The primary modifications as compared to working paper ST/SG/AC.10/C.3/2014/69 ST/SG/AC.10/C.4/2014/12 are as follows:
 - The additivity approach (section 2.8.2.3.3.3) has been simplified. Mixtures classified using the additivity approach will in the amended proposal be classified as Class 8 without sub-classification.
 - As a consequence of the simplified additivity approach, Figure 2.8.3 (flow-scheme for assignment of packing group using the additivity approach) and Table 2.8.2 (concentration limits for use by Figure 2.8.3) are no longer needed and have been deleted.
 - Figure 2.8.4 (flow-scheme for mixtures in Class 8 without sub-classification) has been simplified. Box 6 for calculations of PG III was superfluous and has been deleted. Mixtures classified as Class 8 and which are not assigned to packing group I or II will per definition be in packing group III.
 - As a consequence of a simplified flow-scheme for mixtures classified as Class 8 without sub-classification, generic concentration limits for PG III are no longer needed.
- 4. One of the primary advantages of the proposal is the simplification of the additivity approach. Another advantage is that only two (2) instead of four (4) generic concentration limits are needed.



- 5. The GHS text is more or less unchanged copied into the proposed text for the Model Regulations. The GHS text is included with the aim of optimal global harmonization of criteria now and in the future. Despite the non-legislative style of the GHS text, several examples of successful implementation in jurisdictions exist.
- 6. As compared to the proposal in ST/SG/AC.10/C.3/2014/69 ST/SG/AC.10/C.4/2014/12 from the Netherlands, the proposal in this document contains additions in **bold underlined** font and deletions as double strike through.
- 7. There are square brackets in notes to section 2.8.2.2.4 and Table 2.8.3. The notes to 2.8.2.2.4 refer to the states of the revision of the OECD test guidelines. The square bracket in Table 2.8.3 propose two concentration limits for packing group I and two for packing group II to reflect the current state of the discussion. The sub-committee is invited to take a decision on the most appropriate version of the OECD test guidelines and the concentration limits.

Proposal

8. Replace current Chapter 2.8 in the Model Regulations with the following text:

"CHAPTER 2.8

CLASS 8 – CORROSIVE SUBSTANCES

2.8.1 Definitions and general provisions

- 2.8.1.1 Class 8 (corrosive) substances are substances which, by chemical action, lead to the production of irreversible damage to the skin; namely, visible necrosis through the epidermis and into the dermis, following the application of a test substance for up to 4 hours and observation periods of up to 14 days, or, in the case of leakage, will materially damage, or even destroy, other goods or the means of transport.
- 2.8.1.2 For substances and mixtures that are corrosive to skin, hazard classification is determined using criteria in section 2.8.2. Substances or mixtures shall be classified in one of the three sub-classifications 8A, 8B or 8C. Where the available data do not allow sub-classification, substances and mixtures shall be assigned to Class 8 without sub-classification. Substances and mixtures corrosive to skin are assigned to a packing group using criteria in section 2.8.3. Assignment of sub-classification is based on in vivo results, in vitro results where the methods allows sub-classification, validated Structure Activity Relationship (SAR) methods or by applying the bridging principles. The additivity approach, non additivity approach and extreme pH-value lead to Class 8 without sub-classification.

NOTE: The sub-classifications 8A, 8B and 8C do not constitute divisions in Class 8.

- 2.8.1.3 Liquids and solids which may become liquid during transport, which are judged not to be skin corrosive shall still be considered for their potential to cause corrosion to certain metal surfaces in accordance with the criteria in 2.8.4.
- 2.8.1.4 A substance or a mixture meeting the criteria of Class 8 having an inhalation toxicity of dusts and mists (LC_{50}) in the range of packing group I, but toxicity through oral ingestion or dermal contact only in the range of packing group III or less, shall be allocated to Class 8 (see Note under 2.6.2.2.4.1).

2.8.2 Criteria for hazard classification of substances or mixtures as corrosive to skin

For hazard classification of a substance or a mixture into Class 8, all available information on corrosive properties of a substance or a mixture shall be taken into account in a tiered approach (see 2.8.2.2). Emphasis shall be placed upon existing human data, followed by existing animal data, followed by *in vitro* data and then other sources of information. Classification results directly when the data satisfy the criteria. In some cases, classification of a substance or a mixture is made on the basis of the weight of evidence within a tier. In a total weight of evidence approach all available information bearing on the determination of skin corrosion is considered together, including the results of appropriate validated *in vitro* tests, relevant animal data, and human data such as epidemiological and clinical studies and well-documented case reports and observations.

2.8.2.1 Hazard classification corrosive to skin based on standard animal test data

- 2.8.2.1.1 A substance is corrosive to skin when it produces destruction of skin tissue, namely, visible necrosis through the epidermis and into the dermis, in at least one tested animal after exposure for up to 4 hours. An example of an internationally accepted validated test method for skin corrosion is OECD Test Guideline 404¹.
- 2.8.2.1.2 Three sub-categories are provided within the corrosion Class (Class 8, see Table 2.8.1): Class 8A, where corrosive responses are noted following up to 3 minutes exposure and up to 1 hour observation; Class 8B, where corrosive responses are described following exposure greater than 3 minutes and up to 1 hour and observations up to 14 days; and Class 8C, where corrosive responses occur after exposures greater than 1 hour and up to 4 hours and observations up to 14 days.

CriteriaClass 8Destruction of skin tissue, namely, visible necrosis through the epidermis and into the dermis, in at least one tested animal after exposure $\leq 4 \text{ h}$ Class 8ACorrosive responses in at least one animal following exposure $\leq 3 \text{ min}$ during an observation period $\leq 1 \text{ h}$ Class 8BCorrosive responses in at least one animal following exposure > 3 min and $\leq 1 \text{ h}$ and observations $\leq 14 \text{ days}$ Class 8CCorrosive responses in at least one animal after exposures > 1 h and $\leq 4 \text{ h}$ and observations $\leq 14 \text{ days}$

Table 2.8.1: Skin corrosion hazard classification^a

2.8.2.2 Hazard classification in a tiered approach

2.8.2.2.1 A *tiered approach* to the evaluation of initial information shall be considered, where applicable (Figure 2.8.1), recognizing that not all elements may be relevant.

a The use of human data is addressed in GHS 3.2.2.2 and in GHS chapters 1.1 (par. 1.1.2.5 (c)) and 1.3 (par. 1.3.2.4.7).

¹ OECD Guideline for the testing of chemicals No. 404 "Acute Dermal Irritation/Corrosion" 2002.

- 2.8.2.2.2 Existing human and animal data including information from single or repeated exposure shall be the first line of evaluation, as they give information directly relevant to effects on the skin.
- 2.8.2.2.3 Acute dermal toxicity data may be used for classification. If a substance is highly toxic by the dermal route, a skin corrosion/irritation study may not be practicable since the amount of test substance to be applied would considerably exceed the toxic dose and, consequently, would result in the death of the animals. When observations are made of skin corrosion in acute toxicity studies and are observed up through the limit dose, these data shall be used for classification provided that the dilutions used and species tested are equivalent. Solid substances (powders) may become corrosive or irritant when moistened or in contact with moist skin or mucous membranes.
- 2.8.2.2.4 In vitro alternatives that have been validated and accepted can be used to make classification decisions. Examples of internationally accepted validated test methods for skin corrosion include OECD Test Guidelines 430 ¹ (Transcutaneous Electrical Resistance Test (TER)), 431² (Human Skin Model Test) and 435 (Membrane Barrier Test Method)³. Some in vitro tests are suitable to sub-classify. A substance **or mixture** which is determined not to be corrosive in accordance with OECD Test Guideline 430 or 431 may be considered not to be corrosive to skin for the purposes of these Regulations.
- 2.8.2.2.5 Likewise, pH extremes like ≤ 2 and ≥ 11.5 may indicate skin effects, especially when associated with significant acid/alkaline reserve (buffering capacity). Generally, such substances are expected to produce significant effects on the skin. In the absence of any other information, a substance is considered corrosive (Class 8) if it has a pH ≤ 2 or a pH ≥ 11.5 . However, if consideration of acid/alkaline reserve⁴ suggests the substance may not be corrosive despite the low or high pH value, this needs to be confirmed by other data, preferably by data from an appropriate validated in vitro test.
- 2.8.2.2.6 In some cases sufficient information may be available from structurally related substances to make classification decisions.
- 2.8.2.2.7 The tiered approach provides guidance on how to organize existing information on a substance and to make a weight of evidence decision about hazard assessment and hazard classification (ideally without conducting new animal tests). Although information might be gained from the evaluation of single parameters within a tier (see 2.8.2.2.1), consideration shall be given to the totality of existing information and making an overall weight of evidence determination. This is especially true when there is conflict in information available on some parameters.

OECD Guideline for the testing of chemicals No. 430 "In Vitro Skin Corrosion: Transcutaneous Electrical Resistance Test (TER)" [2004][2013].

OECD Guideline for the testing of chemicals No. 431 "In Vitro Skin Corrosion: Human Skin Model Test" [2004][2013].

OECD Guideline for the testing of chemicals No. 435 "Membrane Barrier Test Method" 2006.

Acid/Alkaline reserve may be determined e.g. by the methodology detailed in Young J. R., How M.J., Walker A.P., Worth W.M.H. (1988): Classification as corrosive or irritant to skin of preparations containing acidic or alkaline substances, without testing on animals. Toxicology in Vitro 2, 19-26 and Young J.R., How M.J. (1994): Product classification as corrosive or irritant by measuring pH and acid / alkali reserve. In Alternative Methods in Toxicology vol. 10 - In Vitro Skin Toxicology: Irritation, Phototoxicity, Sensitization, eds. A.Rougier, A.M. Goldberg and H.I.Maibach, Mary Ann Liebert, Inc. 23-27.

Figure 2.8.1: Tiered evaluation for skin corrosion					
Step	<u>Parameter</u>		Finding		Conclusion
1a:	Existing human or animal skin corrosion data ^a Not corrosive/No data	→	Skin corrosive	→	Classify as skin corrosive ^b
1b:	Existing human or animal skin corrosion data ^a V No/Insufficient data	→	Not a skin corrosive	→	Not classified
2:	Other existing skin data in animals ^c Wo/Insufficient data	→	Yes; other existing data showing that substance may cause skin corrosion	→	May be deemed to be a skin corrosive b
3:	Existing ex vivo/in vitro data d No/Insufficient dataNegative response	→	Positive on corrosivity: Skin corrosive	→	Classify as skin corrosive ^b
4:	pH-Based assessment (with consideration of acid/alkaline reserve of the chemical) e Not pH extreme, no pH data or extreme pH with data showing low/no acid/alkaline reserve	→	$pH \le 2$ or ≥ 11.5 with high acid/alkaline reserve or no data for acid/alkaline reserve	→	Classify as skin corrosive
5:	Validated Structure Activity Relationship (SAR) methods	→	Skin corrosive	→	Deemed to be skin corrosive b
6:	Consideration of the total weight of evidence ^f	→	Skin corrosive	→	Deemed to be skin corrosive b
7:	Not classified				

Existing human or animal data could be derived from single or repeated exposure(s), for example in occupational, consumer, transport, or emergency response scenarios; or from purposely-generated data from animal studies conducted according to validated and internationally accepted test methods. Although human data from accident or poison centre databases can provide evidence for classification, absence of incidents is not itself evidence for no classification as exposures are generally unknown or uncertain;

b Classify in Class 8/sub-classification, as applicable;

- ^c All existing animal data shall be carefully reviewed to determine if sufficient skin corrosion evidence is available. In evaluating such data, however, the reviewer shall bear in mind that the reporting of dermal lesions may be incomplete, testing and observations may be made on a species other than the rabbit, and species may differ in sensitivity in their responses;
- Evidence from studies using validated protocols with isolated human/animal tissues or other, non-tissue-based, though validated, protocols shall be assessed. Examples of internationally accepted, validated test methods for skin corrosion include OECD Test Guideline 430 (Transcutaneous Electrical Resistance Test (TER)), 431 (Human Skin Model Test), and 435 (Membrane Barrier Test Method).
- ^e Measurement of pH alone may be adequate, but assessment of acid or alkali reserve (buffering capacity) would be preferable. Presently, there is no validated and internationally accepted method for assessing this parameter;
- All information that is available shall be considered and an overall determination made on the total weight of evidence. This is especially true when there is conflict in information available on some parameters. Expert judgment shall be exercised prior to making such a determination. Negative results from applicable validated skin corrosion/irritation in vitro tests are considered in the total weight of evidence evaluation.

2.8.2.3 Hazard classification criteria for mixtures

- 2.8.2.3.1 Hazard classification of mixtures when data are available for the complete mixture
- 2.8.2.3.1.1 The mixture shall be classified using the criteria for substances, taking into account the tiered approach to evaluate data for Class 8 (as illustrated in Figure 2.8.1).
- 2.8.2.3.1.2 When considering testing of the mixture, classifiers are encouraged to use a tiered weight of evidence approach as included in the criteria for classification of substances for skin corrosion to help ensure an accurate classification, as well as to avoid unnecessary animal testing. In the absence of any other information, a mixture is considered corrosive (Class 8) if it has a pH \leq 2 or a pH \geq 11.5. However, if consideration of acid/alkaline reserve⁵ suggests the mixture may not be corrosive despite the low or high pH value, this needs to be confirmed by other data, preferably by data from an appropriate validated *in vitro* test.
- 2.8.2.3.2 Hazard classification of mixtures when data are not available for the complete mixture: bridging principles
- 2.8.2.3.2.1 Where the mixture itself has not been tested to determine its skin corrosion potential, but there are sufficient data on both the individual ingredients and similar tested mixtures to adequately characterize the hazards of the mixture, these data will be used in accordance with the following agreed bridging principles. This ensures that the

⁵ Acid/Alkaline reserve may be determined e.g. by the methodology detailed in Young J.R., How M.J., Walker A.P., Worth W.M.H. (1988): Classification as corrosive or irritant to skin of preparations containing acidic or alkaline substances, without testing on animals. Toxicology in Vitro 2, 19-26 and Young J.R., How M.J. (1994): Product classification as corrosive or irritant by measuring pH and acid/alkali reserve. In Alternative Methods in Toxicology vol. 10 - In Vitro Skin Toxicology: Irritation, Phototoxicity, Sensitization, eds. A.Rougier, A.M. Goldberg and H.I Maibach, Mary Ann Liebert, Inc. 23-27.

classification process uses the available data to the greatest extent possible in characterizing the hazards of the mixture without the necessity for additional testing in animals.

2.8.2.3.2.2 Dilution

If a tested mixture is diluted with a diluent which has an equivalent or lower corrosivity classification than the least corrosive original ingredient and which is not expected to affect the corrosivity of other ingredients, then the new diluted mixture may be classified as equivalent to the original tested mixture. Alternatively, the method explained in 2.8.2.3.3 could be applied.

2.8.2.3.2.3 Batching

The skin corrosion potential of a tested production batch of a mixture can be assumed to be substantially equivalent to that of another untested production batch of the same commercial product when produced by or under the control of the same manufacturer, unless there is reason to believe there is significant variation such that the skin corrosion potential of the untested batch has changed. If the latter occurs, a new classification is necessary.

2.8.2.3.2.4 Concentration of mixtures of the highest corrosion sub-classification

If a tested mixture classified in the highest sub-classification for skin corrosion is concentrated, the more concentrated untested mixture shall be classified in the highest corrosion sub-classification without additional testing.

2.8.2.3.2.5 Interpolation within one sub-classification

For three mixtures (X, Y and Z) with identical ingredients, where mixtures X and Y have been tested and are in the same skin corrosion sub-classification, and where untested mixture Z has the same toxicologically active ingredients as mixtures X and Y but has concentrations of toxicologically active ingredients intermediate to the concentrations in mixtures X and Y, then mixture Z is assumed to be in the same skin corrosion subclassification as X and Y.

2.8.2.3.2.6 Substantially similar mixtures

Given the following:

- (a) Two mixtures: (i) X + Y; (ii) Z + Y;
- (b) The concentration of ingredient Y is essentially the same in both mixtures:
- (c) The concentration of ingredient X in mixture (i) equals that of ingredient Z in mixture (ii);
- (d) Data on skin corrosion for X and Z are available and substantially equivalent, i.e. they are in the same sub-classification and are not expected to affect the skin corrosion potential of Y.

If mixture (i) or (ii) is already classified based on test data, then the other mixture can be classified in the same sub-classification.

- 2.8.2.3.3 Hazard classification of mixtures when data are available for all ingredients or only for some ingredients of the mixture
- 2.8.2.3.3.1 In order to make use of all available data for purposes of classifying the skin corrosion hazards of mixtures, the following assumption has been made and is applied where appropriate in the tiered approach:

The "relevant ingredients" of a mixture are those which are present in concentrations $\geq 1\%$ (w/w for solids, liquids, dusts, mists and vapours and v/v for gases), unless there is a presumption that an ingredient present at a concentration < 1% can still be relevant for classifying the mixture for skin corrosion.

2.8.2.3.3.2 Additivity

In general, the approach to classification of mixtures as corrosive to skin when data are available on the ingredients, but not on the mixture as a whole, is based on the theory of additivity, such that each skin corrosive ingredient contributes to the overall corrosive properties of the mixture in proportion to its potency and concentration. The mixture is classified as corrosive when the sum of the concentrations of such ingredients exceeds a cut-off value/concentration limit.

2.8.2.3.3.3 Where the sum of all ingredients <u>corrosive to skin</u> of a mixture <u>is > 5% the mixture shall be classified Class 8 without sub-classification.</u> <u>sub-classified 8A, 8B or 8C is each \geq 5% the mixture shall be classified as skin sub-classification 8A, 8B or 8C, respectively. Where the sum of 8A ingredients is < 5% but the sum of 8A + 8B ingredients is \geq 5%, the mixture shall be classified as sub-classification 8B. Similarly, where the sum of 8A + 8B ingredients is < 5% but the sum of 8A + 8B + 8C ingredients is \geq 5% the mixture shall be classified as sub-classification 8C. Where at least one relevant ingredient in a mixture is classified as Class 8 without sub-classification, the mixture shall be classified as Class 8 without sub-classification if the sum of all ingredients corrosive to skin is \geq 5%.</u>

2.8.2.3.3.4 Non-additivity

Particular care must be taken when classifying certain types of chemicals such as acids and bases, inorganic salts, aldehydes, phenols, and surfactants. The approach explained in 2.8.2.3.3.2 and 2.8.2.3.3.3 might not work given that many such substances are corrosive at concentrations < 1%. For mixtures containing strong acids or bases the pH shall be used as classification criterion (see 2.8.2.3.1.2) since pH will be a better indicator of corrosion than the concentration limits in 2.8.2.3.3.3. A mixture containing corrosive ingredients that cannot be classified based on the additivity approach due to chemical characteristics that make this approach unworkable, shall be classified as Class 8 if it contains $\geq 1\%$ of a corrosive ingredient. Classification of mixtures with ingredients for which the approach in 2.8.2.3.3.3 does not apply is summarized in Table 2.8.2 below.

2.8.2.3.3.5 Exemptions

On occasion, reliable data may show that the skin corrosion of an ingredient will not be evident when present at a level above the generic concentration limits/cut-off values mentioned in 2.8.2.3.3.3 and Table 2.8.2. In these cases the mixture may be

classified according to those data. On occasion, when it is expected that the skin corrosion of an ingredient will not be evident when present at a level above the generic concentration cut-off values mentioned in 2.8.2.3.3.3 and Table 2.8.2, testing of the mixture may be considered. In those cases the tiered weight of evidence approach shall be applied as described in 2.8.2.2 and illustrated in Figure 2.8.1.

2.8.2.3.3.6 If there are data showing that (an) ingredient(s) may be corrosive to skin at a concentration of < 1% (corrosive) the mixture shall be classified accordingly.

Table 2.8.2: Concentration of ingredients of a mixture when the additivity approach does not apply, that would trigger classification of the mixture as corrosive to skin

Ingredient	Concentration	Mixture classified as
Acid with pH ≤ 2	≥ 1%	Class 8
Base with pH ≥ 11.5	≥ 1%	Class 8
Other skin corrosive (Class 8) ingredient	≥ 1%	Class 8

2.8.3 Assignment of packing group

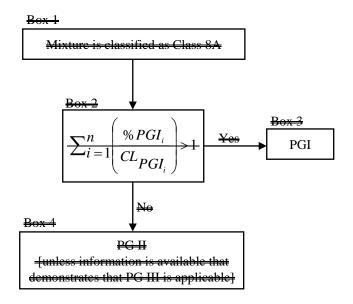
- 2.8.3.1 Substances and mixtures of Class 8 are divided among three packing groups according to their degree of danger in transport as follows:
 - (a) Packing group I: Very dangerous substances and mixtures;
 - (b) Packing group II: Substances and mixtures presenting medium danger;
 - (c) Packing group III: Substances and mixtures presenting minor danger.
 - 2.8.3.2 Allocation of substances and mixtures listed in the Dangerous Goods List in Chapter 3.2 to packing groups in Class 8 has been made on the basis of experience taking into account such additional factors as inhalation risk (see 2.8.1.4) and reactivity with water (including the formation of dangerous decomposition products).
 - 2.8.3.3 Unless otherwise specified in section 2.8.3.4 to 2.8.3.5, substances and mixtures not listed by name in the Dangerous Goods List shall be assigned to packing groups as follows (for assignment of sub-classifications see 2.8.2.1.2):
 - (a) Substances and mixtures classified as Class 8A are assigned to packing group I
 - (b) Substances and mixtures classified as Class 8B are assigned to packing group II
 - (c) Substances and mixtures classified as Class 8C are assigned to packing group III
 - (d) Substances and mixtures classified as Class 8 without sub-classification are assigned to packing group I.
 - 2.8.3.4 Notwithstanding 2.8.3.3, the packing group of mixtures classified as Class 8A based on additivity calculations (see 2.8.2.3.3.2 and 2.8.2.3.3.3) may be assigned using the following method:
 - (a) Derive the packing group for each individual ingredient. For substances listed by name in the Dangerous Goods List, the packing group shall be

- taken directly from the list. For substances not listed by name, the packing group from the most appropriate n.o.s entry shall be used;
- (b) Identify the specific or generic concentration threshold for each individual ingredient. For some substances listed by name on the Dangerous Goods List, the concentration threshold can be taken directly from the list. If no specific concentration threshold is available, generic concentration threshold listed in Table 2.8.3 shall be used;
- (e) Assign the packing group for the mixture in accordance with Figure 2.8.2 [unless information is available that demonstrates that packing group III is applicable].

Table 2.8.3: Generic concentration limit for determination of the packing group of mixtures classified as Class 8A based on additivity calculations

Generic Concentration Limit	Concentration
CL PG I	[5%-]

Figure 2.8.2: Flow chart scheme for assignment of packing group for mixtures with hazard-classification 8.4 based on additivity calculations



Notes to Figure 2.8.2:

% PG I, is the concentration of ingredient i assigned to packing group I.

CL_{PG-li} is the concentration limit for ingredient i with packing group I. This concentration limit can be either a specific concentration limit from the Dangerous Goods List or generic concentration limit from Table 2.8.3.

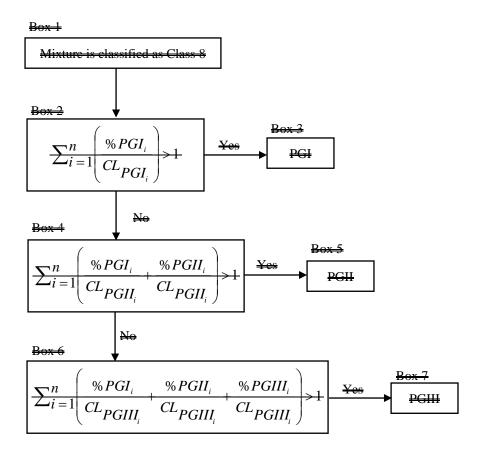
- 2.8.3.<u>5.4</u> Notwithstanding 2.8.3.3 (d), the packing group of mixtures classified as Class 8 without sub-classification may be assigned using the following method:
 - (a) Derive the packing group for each individual ingredient. For substances listed by name in the Dangerous Goods List, the packing

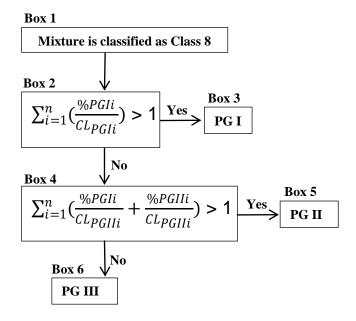
- group shall be taken directly from the list. For substances not listed by name, the packing group from the most appropriate n.o.s entry shall be used;
- (b) Identify the specific or generic concentration threshold for each individual ingredient. For some substances listed by name on the Dangerous Goods List, the concentration threshold can be taken directly from the list. If no specific concentration threshold is available, generic concentration threshold listed in Table 2.8.4 shall be used:
- (c) Assign the packing group for the mixture in accordance with Figure 2.8.3.2.

Table 2.8.4.3: Generic concentration limit for determination of the packing group of mixtures classified as Class 8 without sub-classification

Generic Concentration Limit	Concentration
CL PG I	[5%][<u>50%]</u>
CL PG II	[3%][<u>5%]</u>
CL PG III	[1%]

Figure 2.8.3.2: Flow chart scheme for assignment of packing group for mixtures classified as Class 8 without sub-classification based on non-additivity





Notes to Figure 2.8.3:

% PGI_i is the concentration of ingredient i assigned to packing group I.

% PG II_i is the concentration of ingredient i assigned to packing group II.

% PG III; is the concentration of ingredient i assigned to packing group III.

 CL_{PGIi} is the concentration limit for ingredient i in PG I. This concentration limit can be either a specific concentration limit taken from the Dangerous Goods List or the generic concentration limit from Table 2.8.4.

 CL_{PGII} is the concentration limit for ingredient i in PG II. This concentration limit can be either a specific concentration limit taken from the Dangerous Goods List or the generic concentration limit from Table 2.8.4.

CL_{PG-III} is the concentration limit for ingredient i in PG-III. This concentration limit can be either a specific concentration limit taken from the Dangerous Goods List or the generic concentration limit from Table 2.8.4.

2.8.4 Corrosive to metals

- 2.8.4.1 Liquids, and solids which may become liquid during transport, which are judged not to be corrosive to skin, but which exhibit a corrosion rate on either steel or aluminium surfaces exceeding 6.25 mm a year at a test temperature of 55 °C when tested on both materials are assigned to Class 8.
- 2.8.4.2 For the purposes of testing steel, type S235JR+CR (1.0037 resp. St 37-2), S275J2G3+CR (1.0144 resp. St 44-3), ISO 3574 or Unified Numbering System (UNS) G10200 or a similar type or SAE 1020, and for testing aluminium, non-clad, types 7075–T6 or AZ5GU-T6 shall be used. An acceptable test is prescribed in the Manual of Tests and Criteria, Part III, Section 37.

NOTE: Where an initial test on either steel or aluminium indicates the substance being tested is corrosive the follow up test on the other metal is not required.

2.8.4.3 Packing group III is assigned in accordance with Table 2.8.5.

Table 2.8.5

Packing Group	Effect
III	Corrosion rate on either steel or aluminium surfaces exceeding 6.25 mm a year
	at a test temperature of 55 C when tested on both materials

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