EHS NEWS FLASH

（情報区分; b グリーン調達）

（取得区分;1 委員会報告情報）

1）件名

・RoHS 指令除外更新の官報公布

2）内容

・2018 年 5 月 18 日、RoHS 指令の除外更新 Pack9 で検討されていた除外項目の内、
7 件の除外項目更新に関する官報が公布された。

・以下除外項目内容

(EU) 2018/736

'7(c)-I	Electrical and electronic components containing lead in a glass or ceramic other than dielectric ceramic in capacitors, e.g. piezoelectronic devices, or in a glass or ceramic matrix compound	Applies to categories 1-7 and 10 (except applications covered under point 34) and expires on 21 July 2021. For categories 8 and 9 other than *in vitro* diagnostic medical devices and industrial monitoring and control instruments expires on 21 July 2021. For category 8 *in vitro* diagnostic medical devices expires on 21 July 2023. For category 9 industrial monitoring and control instruments, and for category 11 expires on 21 July 2024.'

(EU) 2018/737

'24	Lead in solders for the soldering to machined through hole discoidal and planar array ceramic multilayer capacitors	Expires on: — 21 July 2021 for categories 1-7 and 10, — 21 July 2021 for categories 8 and 9 other than *in vitro* diagnostic medical devices and industrial monitoring and control instruments, — 21 July 2023 for category 8 *in vitro* diagnostic medical devices, — 21 July 2024 for category 9 industrial monitoring and control instruments, and for category 11.'

(EU) 2018/738

'34	Lead in cermet-based trimmer potentiometer elements	Applies to all categories; expires on: — 21 July 2021 for categories 1-7 and 10, — 21 July 2021 for categories 8 and 9 other than *in vitro* diagnostic medical devices and industrial monitoring and control instruments, — 21 July 2023 for category 8 *in vitro* diagnostic medical devices, — 21 July 2024 for category 9 industrial monitoring and control instruments, and for category 11.'

(EU) 2018/739

'6(a)	Lead as an alloying element in steel for machining purposes and in galvanised steel containing up to 0,35 % lead by weight	Expires on: — 21 July 2021 for categories 8 and 9 other than in vitro diagnostic medical devices and industrial monitoring and control instruments; — 21 July 2023 for category 8 in vitro diagnostic medical devices; — 21 July 2024 for category 9 industrial monitoring and control instruments, and for category 11.
6(a)-I	Lead as an alloying element in steel for machining purposes containing up to 0,35 % lead by weight and in batch hot dip galvanised steel components containing up to 0,2 % lead by weight	Expires on 21 July 2021 for categories 1-7 and 10.'

(EU) 2018/740

'6(b)	Lead as an alloying element in aluminium containing up to 0,4 % lead by weight	Expires on: — 21 July 2021 for categories 8 and 9 other than *in vitro* diagnostic medical devices and industrial monitoring and control instruments, — 21 July 2023 for category 8 *in vitro* diagnostic medical devices, — 21 July 2024 for category 9 industrial monitoring and control instruments, and for category 11.
6(b)-I	Lead as an alloying element in aluminium containing up to 0,4 % lead by weight, provided it stems from lead-bearing aluminium scrap recycling	Expires on 21 July 2021 for categories 1-7 and 10.
6(b)-II	Lead as an alloying element in aluminium for machining purposes with a lead content up to 0,4 % by weight	Expires on 18 May 2021 for categories 1-7 and 10.'

(EU) 2018/741

'6(c)	Copper alloy containing up to 4 % lead by weight	Expires on: — 21 July 2021 for categories 1-7 and 10, — 21 July 2021 for categories 8 and 9 other than *in vitro* diagnostic medical devices and industrial monitoring and control instruments, — 21 July 2023 for category 8 *in vitro* diagnostic medical devices, — 21 July 2024 for category 9 industrial monitoring and control instruments, and for category 11.'

(EU)2018/742

'7(a)	Lead in high melting temperature type solders (i.e. lead-based alloys containing 85 % by weight or more lead)	Applies to categories 1-7 and 10 (except applications covered by point 24 of this Annex) and expires on 21 July 2021. For categories 8 and 9 other than in vitro diagnostic medical devices and industrial monitoring and control instruments expires on 21 July 2021. For category 8 in vitro diagnostic medical devices expires on 21 July 2023. For category 9 industrial monitoring and control instruments, and for category 11 expires on 21 July 2024.'

・適用開始日：2019 年 7 月 1 日

3)SEAJ コメント

・今回の官報公布で、6(a)と 6(b)の除外項目は細分化された。
従来からある除外項目には、カテゴリ 89 製品の有効期限が記載された。
2017 年 9 月 25 日の WTO/TBT 通報時には、18(b)（日焼け用ランプの鉛）も意見募集にかけられていたが、今回 18(b)の官報は公布されなかった。

・RoHS 指令の除外項目に対する対応は各社の判断で行って下さい。

4)添付情報・資料

・なし

5)関連情報

・官報の URL（上から 7(c)-I、24、34、6(a)、6(b)、6(c)、7(a)）
https://eur-lex.europa.eu/legal-content/EN/TXT/PDF/?uri=CELEX:32018L0736&from=EN
https://eur-lex.europa.eu/legal-content/EN/TXT/PDF/?uri=CELEX:32018L0737&from=EN
https://eur-lex.europa.eu/legal-content/EN/TXT/PDF/?uri=CELEX:32018L0738&from=EN
https://eur-lex.europa.eu/legal-content/EN/TXT/PDF/?uri=CELEX:32018L0739&from=EN
https://eur-lex.europa.eu/legal-content/EN/TXT/PDF/?uri=CELEX:32018L0740&from=EN
https://eur-lex.europa.eu/legal-content/EN/TXT/PDF/?uri=CELEX:32018L0741&from=EN
https://eur-lex.europa.eu/legal-content/EN/TXT/PDF/?uri=CELEX:32018L0742&from=EN

6)その他

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