

This is the **accepted version** of the journal article:

Agathokleous, Evgenios; Barceló, Damià; Aschner, Michael; [et al.]. «Rethinking subthreshold effects in regulatory chemical risk assessments». *Environmental Science & Technology*, Vol. 56, issue 26 (2022), p. 11095–11099. DOI 10.1021/acs.est.2c02896

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Rethinking subthreshold effects in regulatory chemical risk assessments

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4 99 A great number of dose-response studies indicate that hormesis is a common phenomenon,
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7 100 occurring in numerous organisms exposed to singular or combined environmental stressors, such
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9 101 as pharmaceuticals, heavy metals, micro/nanoplastics, organic flame retardants, pesticides, and
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11 102 rare earths¹⁻⁶. While biological responses to low exposure levels are often beneficial, exposure
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14 103 to doses below the no-observed-adverse-effect-level (NOAEL; hereafter subthreshold doses)
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16 104 does not always translate to beneficial responses^{2,4}. For example, subthreshold contaminant
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18 105 doses can enhance the virulence of phytopathogenic microbes and promote the resistance of crop
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20 106 pests with significant implications for crop production^{2,7,8}. Subthreshold contaminant exposures
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23 107 can also stimulate infectious animal/human pathogens and promote their resistance to antibiotics
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25 108 and other drugs, threatening long term sustainability. Importantly, the hormetic function of
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27 109 common pathways that regulate cancer progress indicate that current regulatory standards may
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30 110 not protect adequately against cancer risks⁹⁻¹¹.
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32 111 Current risk assessment frameworks used around the world to assess exposure and effects
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34 112 are largely based on scientific developments from the mid-to-late 20th century, which frequently
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37 113 included only very high (often environmentally unrealistic) doses and the broad assumption of
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39 114 linearity in the response in the absence of evidence of alternative dose-response relationships
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41 115 (Fig. 1)¹²⁻¹⁶. How representative and realistic this approach is increasingly being challenged, in
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43 116 a modern era of analytical advances enabling measurement of low doses and hormetic responses.
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46 117 An expanding scientific literature provides evidence of significant effects of subthreshold
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48 118 contaminant doses on numerous animals, plants, and microbes¹⁻⁶. We opine that regulatory risk
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50 119 assessments on exposure and effects should not be based upon outdated science and biologically-
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53 120 unsupported assumptions regarding linearity. Instead, subthreshold effects and dose-response
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55 121 behavior should be included in the regulatory risk assessment. We urge for this approach to be
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3 122 adopted as part of a more real-life risk simulation approach ¹⁷, especially in the light of the
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5 123 growing evidence of genotoxicity of chemicals such as fluoride and arsenic ^{18,19}.

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7 124 Currently, subthreshold responses/effects in regulatory frameworks are largely not
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10 125 considered in worldwide risk assessments, impeding their identification and evaluation
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12 126 (Supporting Information). In the USA, the US Environmental Protection Agency (EPA) does
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14 127 permit non-linear approaches where adequate evidence is provided to prove divergence from the
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16 128 default linear assumption. However, a recent proposal for the inclusion of subthreshold responses
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19 129 and non-default dose-response models in the risk assessment was not implemented ²⁰. In 2017
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21 130 the National Institute for Occupational Safety and Health (NIOSH) acknowledged the dilemma
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23 131 regarding linear extrapolation and endorsed the consideration of non-linear responses for
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26 132 carcinogens in recent new guidelines ²¹. The US Food and Drug Administration (FDA) also
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28 133 recognized non-linear responses in 2018 in its guidance document on the assessment and control
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30 134 of mutagenic substances, and permits deviation from the linear-no-threshold (LNT) dose-
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33 135 response model if protective mechanisms exist ²².

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35 136 In Europe, the European Food Safety Authority (EFSA) has made efforts to evaluate the
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37 137 relevance of subthreshold effects and non-linear responses in recent years ²³. For example,
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40 138 EFSA's scientific committees recently acknowledged subthreshold effects and non-linear
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42 139 responses for bisphenol A and bis(2-ethylhexyl phthalate) and called for internationally-
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44 140 coordinated efforts to identify and address such responses as part of the risk assessment process
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46 141 ²⁴. The European Chemicals Agency (ECHA) also focuses on threshold and non-threshold
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49 142 events, but does not clearly acknowledge or consider subthreshold effects in its guidelines. It
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51 143 does, however, allow the best-fit dose-response model to be used instead of enforcing default
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53 144 dose-response models ²⁵. In 2019, China's Ministry of Ecology and Environment published its
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56 145 trial 'Framework Guide to the Technology Methods of Environmental Risk Assessment for

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3 146 Chemical Substances’²⁰. The framework is based on either threshold or linear no-threshold
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5 147 dose-response models, and does not allow for subthreshold responses/effects or more relevant
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7 148 dose-response modeling based on best fit to specific data sets²⁰.
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10 149 We strongly advocate the consideration of potential subthreshold effects in chemical risk
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12 150 assessment should no longer be postponed. We opine there is an urgent need for regulatory
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14 151 authorities around the world to be inclusive of the most up-to-date science by (re)considering (i)
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16 152 potential subthreshold responses, (ii) non-linear dose-response models able to detect
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19 153 subthreshold responses, and (iii) abandoning the default use of linear dose-response models for
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21 154 all risk assessments. The current lack of subthreshold responses inclusion in the risk assessment
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23 155 of chemicals undermines the accuracy of the risk assessment process, and consequent
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26 156 remediation practices and actions applied. As a recent example, the hormetic model can predict
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28 157 potential subthreshold effects of disinfectants widely introduced into the environment during the
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30 158 COVID-19 pandemic, unlike the linear-no-threshold and threshold models⁵.
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33 159 This article does not suggest that toxicity thresholds are overly conservative and that risk
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35 160 necessarily exists below current limits, but that subthreshold positive or negative effects exist
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37 161 that are not captured by current threshold and LNT models and need to be part of the evaluation
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39 162 and assessment process. Hence, instead of assuming a specific dose-response model *a priori*, the
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42 163 most suitable/effective model to fit or describe the actual data would be selected *ad hoc*. Such a
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44 164 policy would prevent enforcing the exclusion of subthreshold doses and would allow
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46 165 identification of subthreshold effects, as applicable. Furthermore, as lead regulatory agencies
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49 166 increasingly acknowledge subthreshold responses/effects and non-linear dose responses,
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51 167 scientific research should shift the focus to the effects of lower and environmentally realistic
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53 168 doses to facilitate the development of more accurate risk assessments in the future.
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3 170 **Acknowledgments:**
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6 171 The authors are grateful to Dr. Patrick H. Brown, Distinguished Professor of Plant Science at the
7
8 172 University of California, Davis, USA, and Dr. Adrian Covaci, Professor of Environmental
9
10 173 Toxicology and Chemistry at the University of Antwerp, Belgium, for comments and
11
12 174 suggestions on an early draft of the paper.
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17
18 176 **Conflict of Interest Disclosure:** This study did not receive a specific grant from funding
19
20 177 agencies in the public, commercial, or not-for-profit sectors. E.A. acknowledges support from
21
22
23 178 the National Natural Science Foundation of China (No. 4210070867), The Startup Foundation
24
25 179 for Introducing Talent of Nanjing University of Information Science & Technology (NUIST),
26
27 180 Nanjing, China (No. 003080), and the Jiangsu Distinguished Professor program of the People's
28
29
30 181 Government of Jiangsu Province. E.J.C. acknowledges longtime support from the US Air Force
31
32 182 (AFOSR FA9550-13-1-0047) and ExxonMobil Foundation (S18200000000256). The sponsors
33
34 183 were not involved in the study design; the collection, analysis or interpretation of the data; the
35
36 184 preparation of the manuscript or the decision where to submit the manuscript for publication. All
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39 185 authors hold senior editorial positions in various scientific journals. The views presented herein
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41 186 are those of the authors and do not represent views of journals' editorial board as a unit, journals'
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43 187 editorial office, journals themselves or their publishers, authors' institutions, or scientific
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46 188 societies where authors hold senior positions. The authors declare no competing financial
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48 189 interest.
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3 191 **Supporting Information:** Additional text detailing regulatory risk assessment situations in the
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5 192 US (Supporting Text 1), the EU (Supporting Text 2), and China (Supporting Text 3).
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8 9 10 194 **Biography**

11 12 195 **Evgenios Agathokleous⁴**

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14 25 197 Evgenios Agathokleous is a professor at Nanjing University of Information Science &
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25 36
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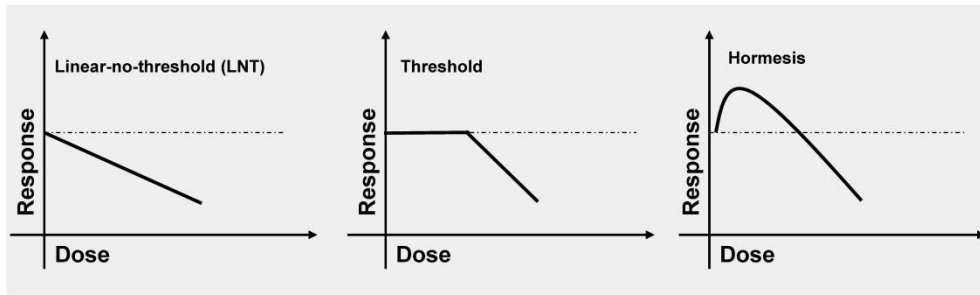
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28 292 **Figure & caption**

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30 293 **Figure 1. Common dose-response relationships.** Linear-no-threshold (LNT) excludes
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32 294 biological repair mechanisms, toxicological threshold, and significant sub-NOAEL (no-
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34 295 observed-adverse-effect-level) responses. Threshold excludes significant sub-NOAEL responses,
35 296 while after NOAEL predicting effects similarly to LNT. Hormesis acknowledges significant sub-
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37 297 and super-NOAEL effects. The dashed line indicates the control response. The relationship's
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39 298 direction is endpoint-specific.
40 299



Common dose-response relationships. Linear-no-threshold (LNT) excludes biological repair mechanisms, toxicological threshold, and significant sub-NOAEL (no-observed-adverse-effect-level) responses. Threshold excludes significant sub-NOAEL responses, while after NOAEL predicting effects similarly to LNT. Hormesis acknowledges significant sub- and super-NOAEL effects. The dashed line indicates the control response. The relationship's direction is endpoint-specific.

248x74mm (600 x 600 DPI)