

## Chapter 11 Section 4 The Implied Powers

OECD Guidelines for the Testing of Chemicals, Section 4 Test No. 442E: In Vitro Skin Sensitisation In Vitro Skin Sensitisation assays addressing the Key Event on activation of dendritic cells on the Adverse Outcome Pathway for Skin Sensitisation OECD Guidelines for the Testing of Chemicals, Section 4 Test No. 417: Toxicokinetics OECD Guidelines for the Testing of Chemicals, Section 4 Test No. 467: Defined Approaches for Serious Eye Damage and Eye Irritation OECD Guidelines for the Testing of Chemicals, Section 4 Test No. 442D: In Vitro Skin Sensitisation Assays addressing the Adverse Outcome Pathway Key Event on Keratinocyte activation OECD Guidelines for the Testing of Chemicals, Section 4 Test No. 493: Performance-Based Test Guideline for Human Recombinant Estrogen Receptor (hrER) In Vitro Assays to Detect Chemicals with ER Binding Affinity OECD Guidelines for the Testing of Chemicals, Section 4 Test No. 426: Developmental Neurotoxicity Study OECD Guidelines for the Testing of Chemicals, Section 4 Test No. 443: Extended One-Generation Reproductive Toxicity Study SCS National Engineering Handbook, Section 4: Hydrology Final Environmental/section 4(f) Statement Middle River Employment Center Access Study, Section 4(f) Evaluation, Baltimore County OECD OECD OECD OECD OECD OECD OECD United States. Soil Conservation Service

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the present key event based test guideline tg addresses the human health hazard endpoint skin sensitisation following exposure to a test chemical skin sensitisation refers to an allergic response following skin contact with the tested chemical as defined by the united nations globally harmonized system of classification and labelling of chemicals un ghs this tg is proposed to address the activation of dendritic cells which is one key event on the adverse outcome pathway aop for skin sensitisation it provides four in vitro test methods addressing the same key event on the aop i the human cell line activation test or h clat method ii the u937 cell line activation test or u sens iii the interleukin 8 reporter gene assay or il 8 luc assay and iv the genomic allergen rapid detection for assessment of skin sensitisers gardtmskin all of them are used for supporting the discrimination between skin sensitisers and non sensitisers in accordance with the un ghs the test methods described in this tg either quantify the change in the expression of cell surface marker s cd54 and cd86 the cytokine il 8 or a series of genes genomic biomarker signature that are associated with the process of activation of monocytes and dc following exposure to sensitisers

this test guideline describes in vivo studies that provide information on mass balance absorption bioavailability tissue distribution metabolism excretion and basic toxicokinetic parameters e g auc as well as supplemental approaches that

this test guideline proposes defined approaches da combining data generated in vitro methods with information sources such as physicochemical properties

this test guideline tg provides an in vitro procedure the are nrf2 luciferase test method used for supporting the discrimination between skin sensitisers and non sensitisers in accordance with the un ghs

this performance based test guideline pbtg describes in vitro assays which provide the methodology for human recombinant in vitro assays to detect substances with estrogen receptor binding affinity hrer binding assays

a developmental neurotoxicity study will provide information on the effects of repeated exposure to a substance during in utero and early postnatal development the test substance is administered daily generally orally to mated females rats are

this test guideline is designed to provide an evaluation of reproductive and developmental effects that may occur as a result of pre and postnatal chemical exposure as well as an evaluation of systemic toxicity in pregnant and lactating females and

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