19	CAS No.: 129-00-0	Substance: Pyrene						
Chem	Chemical Substances Control Law Reference No.: 4-782							
PRTR	PRTR Law Cabinet Order No.:							
	Structural Formula:							
Molec	Molecular Formula: C <sub>16</sub> H <sub>10</sub>							
Molec	Molecular Weight: 202.25							
		$\sum_{i=1}^{n} \sum_{j=1}^{n} \sum_{i=1}^{n} \sum_{i$						

### 1. General information

The aqueous solubility of this substance is 0.135 mg/L (25°C), the partition coefficient (1-octanol/water) (log  $K_{ow}$ ) is 4.88, and the vapor pressure is  $2.45 \times 10^{-6}$  mmHg (= $3.27 \times 10^{-4}$  Pa) (25°C, extrapolated value). The mean biodegradability (aerobic degradation) as determined by BOD, TOC, and GC is 71% (test substance concentration of 5 mg/L) and 11% (test substance concentration of 10 mg/L). The substance does not have any hydrolyzable groups.

This substance is contained in coal tar, and the main applications of coal tar are as a raw material for tar products, rustproof coating, fishing net dyestuff, lamp black, fuel, road paving, roof coating, cast iron pipe coating, waterproof coating, and as an electrode binder. Polycyclic aromatic hydrocarbons (PAHs) containing this substance are unintentionally formed and released to the environment. Release sources of PAHs to the environment are classed as either combustion-derived or non-combustion-derived, and combustion-derived sources are believed to account for more than 90%.

#### 2. Exposure assessment

Because this substance is not a Class 1 Designated Chemical Substance under the Law Concerning Reporting, etc. of Releases to the Environment of Specific Chemical Substances and Promoting Improvements in Their Management (PRTR Law), release and transfer quantities could not be obtained. Predictions of distribution by medium using a Mackay-type level III fugacity model indicated that if equal quantities were released to the atmosphere, water bodies, and soil, the proportions distributed to soil would be higher.

The predicted maximum exposure to humans via inhalation, based on general environmental atmospheric data, was approximately 0.006  $\mu$ g/m<sup>3</sup>. The predicted maximum oral exposure was estimated to be less than around 0.00024  $\mu$ g/kg/day based on calculations from data for groundwater. Further, there is a report of around 0.04  $\mu$ g/kg/day calculated from food data, albeit from a limited area. In addition, while detailed survey results are not clear, a daily intake quantity of 0.03  $\mu$ g/kg/day has been reported based on the findings of a survey that measured quantities contained in samples of various groups of food purchased in seven cities throughout Japan.

The predicted environmental concentration (PEC), which indicates exposure to aquatic organisms, is less than around 0.006  $\mu$ g/L for public freshwater bodies and about 0.010  $\mu$ g/L for seawater.

------

## 3. Initial assessment of health risk

Exposure to sunlight may induce the irritant action of this substance on the skin, causing chronic skin discoloration. Redness of the skin or eyes is caused by contact with this substance. Acute oral and inhalation exposure studies in rats have reported persistent self-stimulation, delirium, myotonia or spasticity.

Sufficient information could not be obtained on its carcinogenicity, and its initial assessment was conducted on the basis of data on its non-carcinogenic effects.

As for its oral exposure, its no-observed-adverse-effect-level (NOAEL) of 75 mg/kg/day (for increase in liver and

kidney weights, nephropathy in females) obtained from its mid-term and long-term toxicity tests for rats was divided by 10, due to their short test periods, to produce 10 mg/kg/day as its 'non-toxic level\*'. As for inhalation exposure, its 'non-toxic level\*' could not be identified.

As for its oral exposure, the predicted maximum exposure was estimated to be around 0.00024  $\mu$ g/kg/day, when intakes of groundwater were assumed. Its margin of exposure (MOE) would be more than 3,100,000 when calculated from its 'non-toxic level\*' of 7.5 mg/kg/day and the predicted maximum exposure, and then divided by 10 due to the fact that 'non-toxic level\*' was obtained from animal experiments. When its exposure through food intakes, as reported for some location, are considered, the predicted maximum exposure will be around 0.04  $\mu$ g/kg/day to provide MOE of 19,000. No further action will be required at the moment to assess health risk from oral exposure to this substance.

As for inhalation exposure to this substance, its 'non-toxic level' could not be identified, and its health risk could not be assessed. The 'non-toxic level' for its oral exposure, if 100% absorption is assumed for it, turns to be the 'non-toxic level' of 25 mg/m<sup>3</sup> for its inhalation exposure. When combined with the predicted maximum concentration in the ambient air, MOE will be calculated to be 420,000. Collection of information on its inhalation exposure to assess health risk associated with exposure to it in the ambient air would not be required.

	I	nformation of toxic	ity		Exposure assessment						
Exposure Path	Criteria for r	isk assessment	Animal	Criteria for diagnoses (endpoint)	Exposure medium	Predicted exposure q concer	maximum uantity and utration	Result of risk assessment			Judgment
Oral	'Non-toxic level <sup>*,</sup>	7.5 mg/kg/day	Mice	Increase in liver and kidney weights and nephropathy in females	Drinking water	_	µg/kg/day	MOE	-	×	0
Ofai					groundwater	< 0.00024	µg/kg/day	MOE	> 3,100,000	0	
Inhalation	'Non-toxic level <sup>*,</sup>	— mg/m <sup>3</sup>	_	_	Ambient air	0.006	µg/m³	MOE	-	×	(O)
					Indoor air	-	µg/m³	MOE	-	×	×

Non-toxic level \*

• When a LOAEL is available, it is divided by 10 to obtain a level equivalent to NOAEL.

• When an adverse effect level is available for the short-term exposure, it is divided by 10 to obtain a level equivalent to an adverse effect level for the long-term exposure.

\_\_\_\_\_

## 4. Initial assessment of ecological risk

With regard to acute toxicity, the following reliable data were obtained: a 48-h median effective concentration ( $EC_{50}$ ) of 4.33 µg/L for swimming inhibition in the crustacean *Daphnia magna*, and a 96-h median lethal concentration ( $LC_{50}$ ) of 30 µg/L in the fish species *Pagrus major* (red sea bream). No value for algae was obtained that could be used, but the acute toxicity to *Pseudokirchneriella subcapitata* was considered to exceed the degree of solubility. Accordingly, an assessment factor of 100 was applied and a predicted no effect concentration (PNEC) of 0.04 µg/L was obtained based on the acute toxicity values.

With regard to chronic toxicity, the following reliable data were obtained: a 21-d no observed effect concentration (NOEC) of 20  $\mu$ g/L for reproductive inhibition in the crustacean *D. magna*; and a 39-d NOEC of 5  $\mu$ g/L was obtained for developmental inhibition in the fish species *Oryzias latipes* (medaka). No value for algae was obtained that could be used, but the chronic toxicity to *P. subcapitata* was considered to be approximately the degree of solubility. Accordingly, an assessment factor of 10 was applied and a predicted no effect concentration (PNEC) of 0.5  $\mu$ g/L was obtained based on the chronic toxicity values. The 0.04  $\mu$ g/L obtained from the acute toxicity to the crustacean was used as the PNEC for this substance.

The PEC/PNEC ratio was less than 0.2 for freshwater bodies and was 0.3 for seawater. Accordingly, further efforts

to collect data are considered necessary. Further, 0.0099  $\mu$ g/L was detected in public freshwater bodies in fiscal 1999, and the ratio between this and PNEC is 0.2.

	Hazard assessment (basis for PNEC)				Predicted no	Expos	ure assessment			
	Species	Acute/ chronic	Endpoint	Assessment factor	effect concentration PNEC (µg/L)	Water body	Predicted environmental concentration PEC (µg/L)	PEC/ PNEC ratio	Result of assessment	
	Crustacean	Acute	EC <sub>50</sub> Swimming	100	0.04	Freshwater	<0.006	<0.2		
(water f	(water flea)	neute	inhibition	100		Seawater	0.010	0.3		

#### 

# 5. Conclusions

		Conclusions					
	Oral exposure	No need for further work.	0				
Health risk	Inhalation exposure	Though a risk characterization cannot be determined, there would be little necessity of collecting information.	(())				
Ecological risk	Requiring information	iring information collection.					
[Risk judgments] O: No need for further work							
■: Candidates for further work ×: Impossibility of risk characterization							
$(\bigcirc)$ : Though a risk characterization cannot be determined, there would be little necessity of							
collecting information.							
$(\blacktriangle)$ : Further information collection would be required for risk characterization.							