1. General information

The aqueous solubility of this substance is 1.9 mg/L (25°C), and the partition coefficient (1-octanol / water) (log Kow) is 4.59. The vapor pressure is 6.28 x 10^{-6} mmHg (= 8.37 x 10^{-4} Pa) (25°C, extrapolated value). Degradability is judged to be good. In terms of hydrolyzability, the half-life at 27°C is 30 - 300 days (pH = 8 - 7). The half-life at 21 ± 2°C is 7.5 days (pH = 8.2) and 1.3 days (pH = 9.5).

The primary applications for this substance are as a plasticizer to provide flame retardant capability, transparency, water resistance, softness and non-adhesive properties to cellulose acetate film, as a plasticizer to provide tensile strength to nitrate cellulose film, as a plasticizer to provide softness, gasoline resistance and mineral oil resistance to synthetic rubber, as a flame retardant plasticizer for phenol resin laminate, and as a flame retardant and plasticizing agent for engineering plastics. Production and import quantities in FY2001 came to 1,000 - 10,000 tons as triphenyl (or monomethylphenyl, dimethylphenyl, nonylphenyl) phosphate.

2. Exposure assessment

As triphenyl phosphate 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 to the environment could not be obtained. When predictions of distribution ratios by medium were made using the Mackay-Type Level III Fugacity Model, in the event of equal release to the atmosphere, water and soil, the distribution ratio was highest for soil.

It was not possible to obtain the data needed to establish a predicted maximum exposure concentration for inhalation exposure to human beings. However, it has been reported that when the data for a limited area (Kawasaki City) was used, the predicted maximum exposure concentration was approximately 0.0035 µg/m³. In addition, it has been reported that the value based on data for indoor air was approximately 0.0151 µg/m³. When data for groundwater and food are used, the predicted maximum oral exposure was estimated to be more than 0.0016 µg/kg/day but less than 0.022 µg/kg/day.

The predicted environmental concentration (PEC) that indicates exposure to aquatic organisms was estimated to be 0.06 µg/L for freshwater and less than 0.01 µg/L for seawater public water bodies.

3. Initial assessment of health risk

No information could be obtained with regard to acute symptoms in humans. However, it has been reported that, when administered orally to rat, mouse and cat, the substance caused diarrhea, tremors, drowsiness, hyperactivity, changes in righting reflex, muscle weakness and other symptoms.

There is insufficient information regarding the carcinogenicity of the substance, and it is not possible to make a judgment as to whether it causes cancer in human beings. For this reason, an initial assessment of the substance was not possible.
conducted based on information of non-carcinogenic effects. As the level at which no adverse effect, etc. was observed, used to estimate the margin of exposure (MOE), a no observed adverse effect level (NOAEL) of 161 mg/kg/day (suppression of weight increase), obtained from rat medium- and long-term toxicity testings, was obtained for oral exposure. As the test period was short, this value was divided by 10 to establish a value of 16 mg/kg/day. For inhalation exposure, a NOAEL value for humans of 3.5 mg/m³ (concentration at which no health effects were observed) was obtained. This value was corrected to match the exposure circumstances, with the result that a value of 0.7 mg/m³ was established.

With regard to oral exposure, when intake of groundwater and food was postulated, the maximum predicted exposure was estimated to be more than 0.0016 µg/kg/day but less than 0.022 µg/kg/day. As the ‘Non-toxic level’ of 16 mg/kg/day and the maximum predicted exposure were established by means of animal testing, the value was divided by 10 to derive an MOE of more than 73,000 but less than 1,000,000. Accordingly, assessment of the health risk from oral exposure to this substance is thought to be unnecessary at this time.

With regard to inhalation exposure, data on a national level could not be obtained, and so the health risk could not be determined. When calculations were performed for reference purposes using general environmental data that had been reported for local areas, the predicted maximum value was estimated at 0.015 µg/m³, and the MOE was 47,000. Although there was a high incidence of detection of the substance in local areas in both the ambient air and indoor air, as the MOE derived for reference purposes is sufficiently high, there is thought to be comparatively little need to determine the concentrations of the substance in ambient air and indoor air.

<table>
<thead>
<tr>
<th>Knowledge of toxicity</th>
<th>Exposure assessment</th>
<th>Result of risk assessment</th>
<th>Judgment</th>
</tr>
</thead>
<tbody>
<tr>
<td>Exposure path</td>
<td>Animal</td>
<td>Exposure medium</td>
<td>Predicted maximum exposure quantity and concentration</td>
</tr>
<tr>
<td>Oral</td>
<td>No observed adverse effect level 16 mg/kg/day Rat Suppression of weight increase</td>
<td>Drinking water / food</td>
<td>– µg/kg/day</td>
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<tr>
<td></td>
<td>Groundwater / food 0.0016 ~ 0.022 µg/kg/day</td>
<td>MOE</td>
<td>73,000 ~ 1,000,000</td>
</tr>
<tr>
<td>Inhalation</td>
<td>No observed adverse effect level 0.7 mg/m³ Human Concentration at which no health effects were observed</td>
<td>Ambient air</td>
<td>– µg/m³</td>
</tr>
<tr>
<td></td>
<td>Indoor air</td>
<td>MOE</td>
<td>–</td>
</tr>
</tbody>
</table>

4. Initial assessment of ecological risk

With regard to acute toxicity, reliable information of a 96-hour EC⁵₀ growth inhibition value of 2,000 µg/L was found for the algae *Pseudokirchneriella subcapitata*, a 96-hour EC⁵₀ value of 250 µg/L was found for the crustacea *Gammarus pseudolimnaeus* (amphipod), a 96-hour LC⁵₀ value of 400 µg/L was found for the fish *Oncorhynchus mykiss* (rainbow trout), and a 48-hour EC⁵₀ growth inhibition value of 360 µg/L was found for the chironomid *Chironomus riparius*. Accordingly, an assessment factor of 100 was used, and a predicted no effect concentration (PNEC) of 3 µg/L was obtained based on the acute toxicity values. With regard to chronic toxicity, reliable information of a 72-hour no observed effect concentration (NOEC) growth inhibition value of 980 µg/L was found for the algae *P. subcapitata*, a 21-day NOEC reproduction value of 254 µg/L was found for the crustacea *D. magna*, and a 30-day NOEC of 87 µg/L at death was found for the fish *Pimephales promelas* (fathead minnow). Accordingly, an assessment factor of 10 was used, and a PNEC value of 9 µg/L was obtained based on the chronic toxicity values. As the PNEC for the substance, a value of 3 µg/L obtained from the acute toxicity for the crustacea was used. The PEC/PNEC ratio was 0.02 for freshwater bodies and less than 0.003 for seawater bodies. Accordingly,
further work is thought to be unnecessary at this time.

<table>
<thead>
<tr>
<th>Species</th>
<th>Acute / chronic</th>
<th>Endpoint</th>
<th>Assessment factor</th>
<th>Predicted no effect concentration PNEC (µg/L)</th>
<th>Exposure assessment</th>
<th>PEC/PNEC ratio</th>
<th>Result of assessment</th>
</tr>
</thead>
<tbody>
<tr>
<td>Crustacea</td>
<td>Acute</td>
<td>EC₅₀</td>
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<td>3</td>
<td>Freshwater</td>
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<td>0.02</td>
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<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Seawater</td>
<td>&lt; 0.01</td>
<td>&lt; 0.003</td>
</tr>
</tbody>
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5. Conclusions

<table>
<thead>
<tr>
<th>Conclusions</th>
<th>Judgment</th>
</tr>
</thead>
<tbody>
<tr>
<td>Health risk</td>
<td></td>
</tr>
<tr>
<td>Oral exposure</td>
<td>○</td>
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<tr>
<td>Assessment is thought to be unnecessary at this time.</td>
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<tr>
<td>Inhalation exposure</td>
<td>×</td>
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<tr>
<td>Risk could not be determined. There is thought to be comparatively little</td>
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</tr>
<tr>
<td>need to determine concentrations for this substance.</td>
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</tr>
<tr>
<td>Ecological risk</td>
<td>○</td>
</tr>
<tr>
<td>No need of further work.</td>
<td></td>
</tr>
</tbody>
</table>

[Risk judgments] ○: No need of further work  ▲: Requiring information collection  ■: Candidates for further work  ×: Impossible of risk characterization