

Figures 1-8. 1. C-metaphase × 1300 2. Sticky metaphase showing unequal segregation × 1200 3. A polyploid cell with gaps × 1350 4. Metaphase with breaks × 1000 5. Sticky anaphase bridge × 950 6. Diagonal anaphase × 850 7. Cell with micronuclei × 1000 8. A binucleate cell × 1050.

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THIOACETAMIDE CAUSES INCREASE IN LEUCOCYTES IN CHANNA PUNCTATUS (BL.).

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THIOACETAMIDE is a known carcinogen. Its main target organs are thyroid and in some cases liver¹⁻³, Its carcinogenic effect on blood has not been reported so far.

The fish Channa punctatus collected from local resources, after acclimatization in the laboratory for ten days, were transferred to aquaria. A sub-lethal concentration of thioacetamide 50 mg/l was dissolved in unchlorinated water (pH 7.5; total solids 14.7 mg/l; alkalinity as CO₃ 57 mg/l; alkalinity as OH 4.5 mg/l; hardness 60-70 mg/l; dissolved oxygen 6-7 mg/l). The blood parameters were studied after the interval of 15, 30 and 45 days.

The observations (Table 1) reveal that thioacetamide sub-lethal concentration 50 mg/l causes decrease in the total erythrocytes count and haemoglobin percentage. On the other hand a rapid increase in the total number of leucocytes count was observed. The increase in leucocytes count is upto approximately 40 times in 45 days. A considerable increase in the percentage of immature crythrocytes and in erythrocytes sedimentation rate (ISR) also suggest leukaemogenic effects in *Channa punctatus*.

The present observations reveal that thioacetamide probably interferes with the development of erythro-

Time (days)	Erythrocytes count × 10 ⁶ /cmm.	НЬ%	Leucocytes count × 10 ³ /cmm.	% of immature erythrocytes	Erythrocytes sedimentation rate
Control	3.95 ± 0.26	14.1 ± 0.7	0.06 ± 0.01	3.9	7.0 ± 1
15	3.80 ± 0.21	13.9 ± 0.8	1.05 ± 0.11	17.0	11.3 ± 1.2
30	3.40 ± 0.19	13.3 ± 0.9	1.16 ± 0.11	26.3	14.5 ± 1
45	2.35 ± 0.23	12.1 ± 0.3	2.35 ± 0.16	65.2	18.0 ± 2

TABLE 1

Treatment of Thioacetamide in days and different blood parameters

cytes in haemopoetic tissues thereby creating leukaemogenic conditions.

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SYNTHESIS OF QUEEN BEE PHROMONE

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THE mandibular glands of the queen honey-bee, Apis mellifera, secrete the queen substance which principally contains 9-0x0- Δ^2 -decenoic acid. The queen substance inhibits the development of ovaries and prevents queen rearing in workers. It also acts as sex attractant in mating¹.

9-oxo- Δ^2 -decenoic acid (VI) has been synthesised from a number of starting materials $^{2-10}$. We report here its synthesis from 7-hydroxyheptanal, one of the periodate oxidation products of aleuritic acid, the major constituent acid of shellac.

7-Hydroxyheptanal (II), on condensation with malonic acid in the presence of pyridine gave an α , β -unsaturated hydroxy acid (III), which on oxidation with pyridiniumchlorochromate resulted in an unsaturated aldehydic acid (IV).

The carbinol (V) obtained by the condensation of IV with CH₃MgI on further oxidation with aluminium tert. butoxide yielded 9-oxo- Δ^2 -decenoic acid (VI).

7-Hydroxyheptanal (II)

Threo-aleuritic acid (I, m.p. 99-100°, 8g) in methanol-water (400 ml, 1:1) at 40°C on sodium periodate oxidation¹¹ for 10 min and on usual workup afforded 7-hydroxyheptanal as liquid (3.2 g). It was purified through a column of neutral alumina by eluting with ether. I.R.(Neat);3250, 1720 cm⁻¹ (Found:C, 64.80; H, 10.72.Calcd. for:C₇H₁₄O₂:C, 64.70; H, 10.80%).

9-Hydroxy- Δ^2 -nonenoic acid (III)

The above hydroxyaldehyde (II, 3 g) was heated on a steam bath for 4 hr with malonic acid (3 g) in dry pyridine (5 ml). Extraction with ether yielded the unsaturated hydroxyacid as thick liquid (2.8 g), which was purified over a column of neutral alumina in ether. I.R.(Neat):3250, 1700, 970 cm⁻¹ (Found:C, 62.72; H, 9.24. Calcd. for C₉H₁₆O₃:C, 62.80; H, 9.30%).

Δ^2 -Noneldehydic acid (IV)

A solution of III (2g) in dry methylene chloride(10 ml) was added with stirring to a suspension of pyridinium chlorochromate (3.28g) and anhy. sodium acetate (0.25g) in dry methylene chloride. After 2 hr, dry ether was added and the supernatant decanted from the black gummy mass. The ethereal extract was then passed through a column of neutral alumina to remove the impurities and the solvent was