(2) In addition to the receptors postulated by the above theory, there are either yellow receptors, white receptors, or both. In this respect the results so far obtained agree with the results obtained by Prof. Ragnar Granit who, as stated above, used the micro-electrode technique on the eyes of animals.

(3) Fixation can be extremely precise, since the effects of eye-movements do not show themselves.

(4) It is possible to stimulate by light, either single cones, or very small groups of cones indeed.

(5) It has been possible to identify, with the precision of at least half 'the cone intercentre distance', the position of some of the receptors which possess specific colour properties.

No evidence has so far been obtained that the green and blue sensa-tions of human vision are due to the combined responses of several different kinds of receptor operating in narrow regions of the spectrum. It is hoped that further research with the microstimulator will help to elucidate this point. I should like to thank Dr. John D'Silva, who acted as recorder for many of the above observations.

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Carcinogenic Substance from Human Cancer

J. F. MENKE⁴ obtained lipid extracts from human breast cancer which, when injected into white mice, induced, in seven of thirty-six animals so treated, the development of sarcomas at the site of in-jection. In our experiments analogous extracts were prepared from gastric carcinoma, three extracts from breast carcinoma, and two extracts from fibrosarcoma. Each of the extracts was tested separately. Four-months-old white mice of our own breeding were used in the experiments. Our strain of mice has a negligible incidence (less than 2 pro mille) of spontaneous tumours. The animals received 10, 20 or 30 mgm. of the lipid extract suspended in sweet almond oil, as a single subcutaneous injection. No differences were noted in the effect with variation of the doses within these limits. Of ninety-four mice injected with the extracts, twenty died within the first four months of the experiment. Of the remaining seventy-four mice, twenty-one animals (28 4 per cent) developed malignant tumours. The tumours developed chiefly in organs at a distance from the site of injection, and exhibited various histological types including carcinoma and sarcoma. Gastric carcinoma extracts provoked two breast carcinomas (in two females), two lung lymphosarcoma (in one funded, and one female), one lung lymphosarcoma (in one fome founde), one liver carcinoma (in one male) and two lymphosarcomas at the site of injection (in two females). Fibrosarcoma at the site of injection (in one female), one lung carcinoma extracts provoked four breast carcinomas (in four females). Fibrosarcoma at the site of injection (in one female), one lung carcinoma at the site of injection (in one female), one lung carcinoma at the site of injection (in one female), one lung carcinoma (in one male), one kidney carcinoma (in one male), and one lymphosarcoma at the site of injection (in one female). All seven extracts tested Induced approximately the same percentage of tumours in the animals treated. The average, period of time

Cartendonia (in one male), and one sympnosarcona at the site of m-jection (in one female). All seven extracts tested induced approximately the same percentage of tumours in the animals treated. The average period of time necessary for the development of tumours was 6 months for the gastric carcinoma extracts, 11 months for the breast carcinoma extracts, and 7.6 months for the fibrosarcoma extracts. Of the fifty-three animals which died without developing tumours, the individuals survived as follows: four for 5 months, eight for 6 months, five for 7 months, seven for 8 months, four for 9 months, two for 10 months, two for 11 months, three for 12 months, and eighteen longer than 12 months. Attempts undertaken with the aim of separating the active factor from the extracts resulted in a marked diminution of the number of malgnant tumours provoked. Of fifty-seven mice injected with the chemically modified extracts, only six animals developed cancer. The average time for the development of the tumour after the single injection was approximately twice the time observed with the non-modified extracts. Lipid carcinogenic extracts have been obtained from human livers²⁻⁸ and from beef pituitary glands⁴. These experimental findings indicate

and from beef pituitary glands⁶. These experimental findings indicate that a lipid carcinogenic substance, probably of hormonal character, can be extracted from certain organs. Our experiments demonstrate that an analogous substance is present in human cancers. For this substance we propose the name 'boardin', which is accepted in our laboratories. Attempts are in progress to separate boardin from the extracts. extracts

The histological diagnosis of the tumours was verified by Dr. Francis Carter Wood, to whom we are indebted for his co-operation. HENRY K. WACHTEL

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Transmission of Litomosoides carinii to Mice and Hamsters

Litomosoides carinii is a filariid parasite of the cotton rat, Sigmodon

Litansmission of Litamosoides calinii to Mice and Hamsters Litamosoides carinii is a filaridi parasite of the cotton rat, Sigmodon Mispidus. It has been used extensively in the United States to investig-ate the chemotherapy of filarial infections. R. W. Williams and H. W. Brown' and J. A. Scott (private communication) have recently shown that infection was transmitted from one animal to another by means of the tropical rat mite Liponyssus bacoti. These workers kindly showed their results and methods to one of us and provided us with a colony of the mites and some infected cotton rats. Further infected cotton rats were kindly lent us by Prof. R. M. Gordon. The transmission of Litamosoides to clean cotton rats and to lab-oratory (piebald) rats has been confirmed in this Institute, micro-filarize being found in the blood of the rats 63 days after the first exposure to infected mites. The blood of some of these rats has con-tained as many as 450,000 microfilarize per c.c. In addition, the attempt was made to transmit the infection to other laboratory animals. Nine albino mice were exposed to infected mites for 40-70 days. After 42 days, one mouse was killed and nine worms, measuring 5-14 mm. long, were found in the pleural cavity. The blood of the other mice was examined at somewhat irregular intervals. Microfilariæ were found in the blood of two mice, each on a single occasion, on the eighty-second and ninety-first days re-spectively after the beginning of the exposure to infection. No microfilariæ have been seen in the blood of the other six mice up to the ninetieth day. Three hamsters (Cricetus (Mesorriceus) auratus) were exposed to infection for periods of 26-44 days. One hamster was killed after 44 days; the pleural cavities contained thirty-four worms, 1-3 cm. long, both sexes being present. In the case of the third hamster, microfilariæ were found at the first examination of the blood made on the seventieth day and on subsequent occasions, the number present being small.

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Distribution of Number of Segments in Earthworms and its Significance

DURING the course of an investigation into the relations between

and its Significance



