ERYTHROCYTES


The maturation of rabbit reticulocytes was studied in vitro and in vivo. The RNA and Mg++ contents of the red cells were compared in vitro and in vivo. The maturation time of reticulocytes in vivo was 60-80 hr; in contrast, the decline in RNA content of incubated reticulocytes suggested maturation times of 36-48 hr. P.F.


Kininase activity of plasma, whole blood, erythrocytes, and hemolysates was examined in patients with hypochromic anemia and in healthy subjects. Kininase activity was found to be highest in erythrocyte hemolysates. It was demonstrated that kininases do not penetrate through undamaged erythrocyte membrane. Trasylol (200 U/ml) and epsilon-amino-caproic acid (6 x 10^-3 M) did not inhibit the kininase activity of plasma, whole blood and of hemolysates. 11-orthophenanthroline (3 x 10^-3 M) was found to inhibit the kininases present in plasma or hemolysates while EDTA (4 x 10^-3 M) induced an inhibitory effect on plasma kininases only. No significant differences in kininase activity could be detected in hemolysates from anemic patients and from controls. M.A.

Erythropoietin in Human Milk. M. Bielecki, F. Praca, and M. Jazewska. Second Department of Obstetrics and Gynecology, School of Medicine, Białystok and Institute of Physiology and Biochemistry of Animals, School of Agriculture, Olsztyn, Poland. Acta Physiol Pol 23:497 502, 1972
The content of erythropoietin in human milk during the first week of lactation was examined using a biological test. The effect of extracts of deproteiniized human milk on $^{59}$Fe incorporation into erythrocytes of polycythemic mice was determined and compared with the results obtained in two control groups treated either with saline or with preparations of standard erythropoietin. It was found that human milk contained circa 0.015 IU of erythropoietin per milliliter. M.K.


The results of investigations on changes occurring in complexes of iron with nucleotides during blood storage are reported. The blood was collected into ACD solution and stored at 4°C. Free nucleotides of erythrocytes were separated by column chromatography on the Dowex 1 x 8 (200-400 mesh) using continuous formate gradient. No changes of total content of iron in the acid-soluble fraction of erythrocytes were observed during storage of the blood for 21 days. During this period of time, disintegration of the complex of ATP with Fe was demonstrated. The iron released from this complex was eluted with the fractions containing NADP, IMP, and ADP. In the samples of blood stored for 21 days, the total content of iron in the erythrocyte extracts decreased and the concentration of Pi, ADP, NADP, NAD, and ATP diminished probably due to impairment of cell membrane permeability. M.K.


Incorporation of $^{59}$Fe into erythrocytes and its retention in the liver and spleen were compared in control rats and in rats with experimental polyarthritis induced by intradermal injections of complete Freund adjuvant. $^{59}$Fe-citrate, $^{59}$Fe-hemoglobin isolated from rat erythrocytes and erythrocytes labeled in vivo with $^{59}$Fe and stored for 16 days before injection were used as radioiron preparations. In the rats with postadjuvant disease but without pronounced anemia, incorporation of $^{59}$Fe from old erythrocytes into red blood cells was significantly lower while retention of $^{59}$Fe from $^{59}$Hb and old erythrocytes in the liver was markedly greater than in controls. In the rats with postadjuvant disease associated with severe anemia incorporation of $^{59}$Fe into erythrocytes was significantly greater than in controls independently on the form in which $^{59}$Fe was applied. Retention of $^{59}$Fe from old erythrocytes in the liver in experimental groups with marked anemia was greater than in controls. The results indicate that reutilization of iron from effete erythrocytes is impaired in postadjuvant disease but considerable anemia may diminish this impairment. M.K.


The level of serum haptoglobin was determined in patients with polycythemia vera and with polycythemia secondary to the syndrome of cor pulmonale, cyanotic cardiac diseases and kidney diseases. Highly significant increase of haptoglobin level was observed only in patients with the cor pulmonale syndrome. The authors conclude that determination of serum haptoglobin level may be helpful in differentiating polycythemia associated with cor pulmonale syndrome from polycythemia vera and other forms of polycythemia. The second conclusion is that "central" hypoxia has no effect on the development of hyperhaptoglobinema. M.K.


The appearance of symptoms of permanent intravascular hemolysis (hemoglobinemia, hemosiderinuria) together with positive Ham's and Hartmann-Jenkins tests on a background of hypoplastic bone marrow is not infrequently the first clinical symptom of the Strübing-
Marchiafava disease, i.e. PNH. When this is disregarded, the disease may remain unrecognized for a long time, until the typical syndrome of paroxysmal nocturnal hemoglobinuria becomes apparent. In some cases the disorder begins with thrombocytopenic purpura (TP), while the typical PIH appears several years (in one female patient 12 yr) after the removal of the spleen for TP. In other cases, transformation of the hypoplastic bone marrow into a partially hyperplastic one with predominance of erythroid cells, imitating erythremia developed spontaneously. Because of the permanent character of the intravascular hemolysis, we suggest that the title PIH (permanent intravascular hemolysis) is more correct than that of PNH which only reflects a moment of the clinical moving picture. The development of symptoms of PIH on the background of a hypoplastic bone marrow is considered as the second phase of the Marchiafava disease. The myeloid metaplasia of the spleen is considered as the third phase. P.F.


In view of the relationship between irradiation of the fetus, chromosomal damage and leukemia, a study was made of the lymphocytes of 20 neonates whose mothers had been injected with 113mIndium for placental scanning. Cord blood was cultured with PHA and 100 metaphases examined. No difference was found from control infants, and it was concluded that the isotope did not cause microscopically evident chromosomal damage. A.A.M.


Obstructive prostatic symptoms were encountered in 14 patients with leukemia, in six of whom there was leukemic infiltration of the gland; these six patients showed no characteristic symptoms or physical findings. J.M.B.


Twenty-one patients with Hodgkin’s disease who were in relapse following extensive radiation therapy were treated with a combination chemotherapy program (MOPP), which included nitrogen mustard, vincristine, prednisone, and procarbazine for 6 monthly cycles. Sixteen patients (76%) achieved complete re-
Activity of Nonspecific s-naphthylacetate mission. In a comparable group of patients with extensive disease but no previous radiotherapy, the overall remission rate and the degree of myelosuppression were similar. The interval between the end of radiotherapy and the onset of chemotherapy did not correlate with the extent of subsequent drug toxicity. Relapse of Hodgkin's disease in the patients following intensive radiotherapy with curative intent does not preclude a subsequent excellent response to combination chemotherapy and a prolonged disease-free interval. J.E.U.


Case records of 425 patients with Hodgkin's disease treated at the NIH were reviewed. Note of all biopsy-proved malignant tumors other than Hodgkin's disease was made. Cases were divided into subgroups on the basis of treatment received, and expected incidences of malignant tumors were calculated for each subgroup on the basis of age, sex, and mean follow-up period from the time of diagnosis of Hodgkin's disease. Significantly increased risks of development of second malignant tumors were found in the entire 425 patients (ratio of observed to expected, 3.5) and in the subgroups treated with both radiotherapy and chemotherapy (ratio, 3.3) and with intensive radiotherapy without intensive chemotherapy (ratio, 3.8). The greatest increase in risk was observed in 35 patients who received both intensive radiotherapy and intensive chemotherapy (ratio, 29). J.E.U.


The HL-A phenotypes of 127 patients with Hodgkin's disease have been determined. A very significant association has been found between Hodgkin's disease and two HL-A antigens, HL-All (p < 0.009), and W5 (p < 0.005). The families of 40 of these patients were genotyped for HL-A antigens. A normal mendelian segregation of the relevant antigen was found in all 12 families of HL-All positive patients and in six of eight families of W5 positive patients. These findings suggest that certain Hodgkin's patients have a genetically determined susceptibility to their disease. It is postulated that this susceptibility could be due to linkage between HL-A genes and genes controlling immune responsiveness. Analysis of subgroups of Hodgkin's patients based on age, sex, and pathology suggests that these HL-A associations are most marked in certain subgroups. J.E.U.

Characteristics of Lymphoid Cells of Acute Leukemia in Children. N. S. Kislak, R. V. Lenskaya, and E. A. Morozova. Second Institute of Medicine, Moscow, USSR. Probl Gematol Pereliv Krovi 1:11-14, 1973

Cytochemical and clinical investigations in 49 children (ages 3-15 yr) with acute lymphocytic leukemia are presented. Two types of lymphoid cells (lymphoblasts and lymphoreticular cells) are described. These two types of cells are morphologically similar, but we can differentiate one type from the other by cytochemical properties (especially by the high activity of acid phosphatase revealed by the method of Goldberg and Barka). High activity of hydrolytic enzymes in the microsomal fraction was observed in the lymphoreticular cells. Because of this peculiarity, the lymphoreticular cells look like the cells of a monocytoid line. A marked tendency to tumor growth of lymph nodes and thymus in children with lymphoreticular cell infiltration in the bone marrow was noted. G.A.

Differentiation of Bone Marrow Cells by Examination of Acid Phosphatase in Children With Diseases of the Blood System. I. S. Peterson. Institute of Pediatrics, Moscow, USSR. Probl Gematol Pereliv Krovi 1:15-16, 1973

The activity of acid phosphatase was studied by the method of Goldberg and Barka in the bone marrow of 19 children (ages 5-14 yr) with various hematological diseases. Three children with acute lymphoblastic and a child with acute myeloblastic leukemia were studied during complete remission. In all these cases a high activity of acid phosphatase was observed in the cells of the granulocytic line. G.A.

Activity of Nonspecific α-naphthylacetate
Esterase in the Bone Marrow Cells of Children With Acute Leukemia. N. A. Rybakova, and R. V. Lenskaya. Second Institute of Medicine, Moscow, USSR. Probl Gematol Pereliv Krovi 10/9 14, 1972

The activity of nonspecific α-naphthylacetate esterase was studied in the cells of blood and bone marrow of 130 children with acute leukemia. A method based on the use of α-naphthylacetate was utilized. A parallel test with an inhibitor of the reaction (sodium fluoride in concentration of 1.5 mg/ml) was carried out. A precise identification of leukemic cells was possible by this method. This reaction is of particular significance for the differentiation of monocytic leukemia. In monocytic cells the activity of the enzyme which can be completely blocked by the inhibitor is high. In promyelocytic acute leukemia, activity of nonspecific esterase is also high, but it can only little be affected by sodium fluoride. Particularly high activity of the enzyme in 100% of cells, which was not influenced by the inhibitor, was observed in patients with plasma-cell leukemia. In cases of erythroleukemia, a characteristic picture was observed. On the background of an intensified diffuse staining in the perinuclear zone of the erythroblasts one could see "spots" corresponding to high activity of the enzyme. The inhibitor suppressed the activity of the enzyme by one half. Besides the identification of cell forms, of great importance is the determination of cell maturity. In lymphoid and monocytic cell lines the enzyme activity increased together with the increase of cell maturity. In the neutrophilic cell line, the highest activity was in the promyelocytes. In the myeloblasts and segmented neutrophils, the enzyme activity was absent. — G.A.

Mitotic Indices in Various Forms of Acute Leukemia. I. Ursiniński. Second Department of Internal Medicine, School of Medicine, Szczecin, Poland. Acta Haematol Pol 4:9–26, 1973

Mitotic indices were determined in populations of bone marrow paraleukoblasts in 51 patients with various forms of acute leukemia and compared with these indices calculated for granulocyte precursors in bone marrow of five healthy subjects. Mitotic indices in patients with acute leukemias were significantly lowered and showed a very wide scatter. The lowest values of mitotic indices were observed in lymphoblastic leukemia, undifferentiated cell-leukemias and in patients with the blastic crisis during the course of chronic granulocytic leukemia. Higher values were usually found in pararoblasts, monocytic and particularly parapromyelocytic leukemias. The results are discussed in respect to pathogenesis and treatment of acute leukemias. — M.K.


Cytophotometric determinations of DNA and histone proteins in leukocytes of peripheral blood were carried out in 22 patients with various types of leukemia. Paraleukoblasts from patients with paramyeloblastic and parapromyelocytic leukemias as well as with myeloblastic crisis in the course of chronic granulocytic leukemia were found to contain decreased amounts of histone proteins as related to DNA content. Therefore, the DNA/histone ratio was increased in the leukemic cells. On the contrary, leukemic lymphoblasts showed the DNA/histone ratio approximating the values obtained for normal lymphocytes. The authors suggest that the changes observed may be connected with heterochromatinization of the nuclei of parameyloblasts. — M.K.

Storage of Leukocyte Concentrates at a Temperature of –196°C. J. Dąszyński and M. Karelus. Blood Bank, Institute of Hematol-

A technique for storage of leukocyte concentrates at low temperatures is reported and a special device enabling freezing at slow rates is described. Viability of leukocytes suspended in four cryoprotective fluids (containing either various concentrations of glycerol or dimethylsulfoxide or polyvinylpyrrolidone) was compared after freezing and thawing. On the basis of the results obtained the authors recommend a fluid of the following composition: 26.0 g of glycerol, 9.0 g of sucrose, 2.0 g of glucose, 0.1 g of Na2 EDTA, bidestilled water to 100.0 ml. — M.K.


The ability of leukocytes to phagocytize latex particles and to reduce NBT was investigated in diabetic children by the method of Park et al., and compared with the results obtained in controls. Both groups were divided into two subgroups depending on the presence or absence of bacterial infections. A significant suppression of the leukocyte functions was observed in diabetic patients without bacterial infections if compared with appropriate controls. The ability of NBT reduction by leukocytes of diabetic children during bacterial infections approached that observed in the nondiabetic children. The author suggests that disturbances in phagocytosis and NBT reduction by leukocytes may be of some significance for the course of bacterial infections in diabetic children. — M.K.


Investigations were carried out in seven patients with chronic lymphatic leukemia (CLL), eight with chronic myeloid leukemia (CML), and five with acute myeloid leukemia (AML). Phagocytic ability of leukocytes was found to be significantly increased in untreated CLL, normal in CML, and markedly decreased in AML. Repeated examinations showed no fluctuations in phagocytic activity of leukocytes in CML and AML, but slowly decreasing values in CLL. The bacteriostatic activity of sera depended on the type of antibodies administered during the observation. Complement activity did not differ from normal values in all cases while properdin level was found to be significantly lowered in the majority of the patients. — M.K.


Thirty-four patients with burns of various severity were examined. Differential leukocyte counts and selected morphological, cytochemical, and functional parameters were investigated. Peripheral leukocytes were found to be increased with higher proportion of young forms of granulocytes and absolute lymphopenia. The presence of persisting granules in granulocytes was found to correlate with increased alkaline phosphatase activity. Significantly increased was the percentage of PAS positive lymphocytes (in some cases above 50%). Some impairment of blastic transformation was observed in lymphocytes cultured in the presence of the patients own plasma and stimulated with PHA. — M.K.

HEMOSTASIS


Specific antibodies were prepared against human prothrombin, thrombin, and serum derivatives of prothrombin, respectively. These allowed to tentatively analyze the changes in the prothrombin molecule during its activation into thrombin. Thrombin and serum prothrombin derivatives had antigenic determinants in common with prothrombin, but none in common with each other. It appears that the thrombin moiety of prothrombin can acquire
some new antigenic determinants during prothrombin fragmentation. *J.C.*


Analysis of the technical conditions for a better use of the fibrin plate method is described. It is recommended to use always the same substrate, with a 100 mg/100 ml fibrinogen concentration. *P.d.N.*

**Immunologic Evaluation of Plasminogen in Patients With Liver Cirrhosis.** S. Pisarri and F. Salsano. Institute of Semeiotica Medica University, Rome, Italy. Progr Med (Rome) 28:24 26, 1972

The study includes 35 patients. In 65% of the cases the plasminogen values were lower than in normal conditions. A good correlation was found between the results obtained with the immunologic method and those obtained with the enzymatic method. The authors assume that in liver cirrhosis there is a reduced synthesis of inhibitors of fibrinolysis which may influence the enzymatic evaluation method. *P.d.N.*


This is a study of 18 cases. Low values of plasminogen were found in 18%, and related to a decreased synthesis in the liver. No hemorrhagic symptoms were observed. These investigations are limited to a particular aspect of the problem but deserve to be taken into consideration because of the proper technique used and of the number of cases studied. *P.d.N.*


Prostaglandins E2 and F2α are known to be formed and released from platelets by thrombin. In this study the authors investigate the effect of ADP, epinephrine and collagen on the production and release of prostaglandins by platelets. The prostaglandins were identified either by biological assay or by a radioimmuno assay. Nonaggregated platelets showed exceedingly low prostaglandin levels (less than 1 pmol PGF2α/ml). When platelets were aggregated by ADP (40 μM) and epinephrine (40 μM) production and release of prostaglandins from platelets was increased to 3 and 4.4 pmoles/ml, respectively. A proportional increase in prostaglandin E2 and F2α was noted with increasing concentrations of collagen added to the platelet suspension. In experiments using low concentrations of ADP (1.4 μM) prostaglandin formation could not be demonstrated. Prostaglandin production was always highest in thrombin aggregated platelets (9.8 pmoles/ml). Almost 100% of the prostaglandin produced by the platelets was released during the aggregation process. Prostaglandins were synthesized apparently as aggregation began. Synthesis stopped after 5 min of incubation with collagen. The authors suggest that formation and release of prostaglandins is associated with secretion of endogenous ADP and 5-hydroxytryptamine in the release reaction. *M.S.*

**Thrombokinetics in Dietary Induced Folate Deficiency in Human Subjects.** D. H. Cowan and J. D. Hines. Department of Medicine, Case Western Reserve University, School of Medicine and the Cleveland Metropolitan General Hospital, Cleveland, O. J Lab Clin Med 81:577, 1973

Platelet production and survival were performed in human subjects with dietary induced folate deficiency. The mean number of megakaryocytes in seven nonthrombocytopenic folate deficient chronic alcoholic subjects studied during periods of abstinence was three times that observed in normals. A similar increase was also noted in the total mass of megakaryocytes but the mean volume of megakaryocytes in the folate deficient group differed only insignificantly from that in the normal. In three chronic alcoholic subjects provided with folate supplementation the values of megakaryocyte number, volume and total mass were normal. 51Cr survival of autologous platelets was normal in both groups and the shape of the
defined in folate deficiency before thrombocyto-
ceeded effective cu mm/day). In patients with mild to moderate
peniasetsin. MS.

normal pattern of thrornbokinetics can be
determined from the total megakaryocyte mass ex-
cceeded effective thrombopoiesis determined from the platelet turnover by 2.5-fold. An ab-
normal pattern of thrombokinetics can be
defined in folate deficiency before thrombocyto-
penia sets in. M S.

The Kinetics of Antiheparin Platelet Factor Re-
lease. II. The Role of Epinephrine and
Norepinephrine. G. M. Gandolfo, P. Bongio-
anni, and R. Ottavianii. Institute of Semei-
otica Medica, University, Rome, Italy. Progr
Med (Rome) 28:431 437, 1972

The release of platelet antiheparin factor is
proportional to the concentration of catechol-
amines added to platelet-rich plasma. It is
early in occurrence and takes place at the time
of the second wave of aggregation. It only oc-
curs at 37°C (not at 25°C). Abstractor's com-
ment: The results are interesting and ac-
curate. P.d.N.

Cellular Control of Blood Coagulation. III.
The Response of Platelets to Adrenaline.
H. Kowarz, L. Czerkawski, K. Krcmenieh,
and R. Janda. Department of General and
Experimental Pathology, School of Medi-
cine, Wroclaw, Poland. Bull Acad Pol Sci
(Biol) 20:755 759, 1972

The authors demonstrated that activation of
clotting induced in rats by infusion of adrena-
line (4 μg/kg body weight) is accompanied by
significant drop in platelet counts. These two re-
sponses to adrenaline could be abolished by
splenectomy and restored in splenectomized
animals by intravenous infusion of 5.5 x 10⁸
autologous spleen cells. It was found that
Trasylol inhibits blood clotting activation and
drop in platelet counts induced by adrenaline.
10 times lower doses of Trasylol being re-
quired for inhibition of platelet changes. The
authors conclude that activation of clotting and
decrease in platelet counts induced by
adrenaline depend on spleen cell and are
mediated by a factor sensitive to Trasylol
which is known to inhibit kallikreine and
trypsin. It is assumed that this factor may be a
proteolytic enzyme. M K.

Results of Long-term Administration of Anti-
coagulants in Patients With Myocardial
Infarction and Coronary Disease Without
Infarction. A. Korolko. Department of
Cardiology, School of Medicine, Lublin,
Poland. Pol Tyg Lek 27:1897 1900, 1972

Long-term treatment with anticoagulants
(Sintrom, Synumar) was carried out in 300
patients with myocardial infarction and in 106
patients with coronary disease without infar-
tion, aged 31 71 yr. The duration of treatment
ranged from 1 to 6 yr and the prothrombin
level was maintained in the range of 35°, 45°.
The control groups of patients nontreated with
anticoagulants comprised 347 cases with myo-
cardial infarction and 195 with coronary dis-
 ease without infarction. Treatment with anti-
coagulants was started on the first day of
hospitalization and was continued after dis-
charge under control of the outpatient service.
In the group of patients with myocardial in-
farction treated with anticoagulants the death
rate was 8.0°, while in the nontreated patients
it was 20.5°. In the treated group, repeated
infarctions occurred in 12.7°, in the controls
these were 33.1°. On the contrary, no differ-
eence in death rate was observed in patients
with chronic coronary disease treated with
anticoagulants and in the respective control
group. M.K.

IMMUNOHEMATOLOGY

Serum Immunoglobulin Patterns in the First
Year of Life in Normal and Low Birth
Weight Infants. I. Relationship to Birth
Weight. C. J. Lewis, L. H. Stevens and J.
Vivian Wells. University of New South
Wales, Sydney, Australia. Med J Aust
1:282 288, 1973

A longitudinal study of immunoglobulin
levels was performed in 103 infants of low birth
weight and in their mothers. The immunoglobu-
lin levels of the mothers were not related to the
birth weight of their infants but the level of
cord IgG in the infants was significantly related
to birth weight. The rate of decay of the ma-
ternally acquired IgG was not related to weight
and the ultimate nadir of IgG reached between
12 and 24 wk reflected the initial IgG level.
Thus of infants weighing less than 2000 g, 50°,
showed a serum IgG level below 200 mg/100 ml
and this was associated with a high incidence of
infection. A.A.M.
ABSTRACTS


Mice were rendered tolerant to injection of "remission" bone marrow cells from acute leukemia patients by simultaneous injection of cyclophosphamide. Antisera were then raised to blast cells from two patients, one with acute myeloblastic and the other with acute lymphoblastic leukemia. In both cases the antisera produced were more toxic to blast cells than to remission cells and were believed to be directed against leukemia-specific antigens. A.A.M.


PHA transformation of lymphocytes from patients with AML before therapy was reduced only if cultures contained a significant number of primitive cells; modification of the cultural conditions could be associated with significant PHA response in such cultures. Serum from the majority of patients with AML inhibited "in vitro" transformation of normal lymphocytes. —J. M. B.


In two patients presenting as chronic lymphocytic leukemia, immunofluorescence studies have demonstrated the presence of an accumulated intracytoplasmic material reacting with antisera to \( \mu \) chains and only to one type of light chains, whereas their serum contained neither monoclonal IgM nor free \( \mu \) chains. In the first case, the IgM \( \lambda \) inclusion bodies were crystals detected in the cytoplasm of small lymphocytes and were not clearly lined by rough endoplasmic reticulum. IgM \( \lambda \) molecules were present at the surface of the lymphocytes. In the other patient, who elaborated free \( \kappa \) light chains, the inclusions found in marrow plasma cells and in peripheral blood lymphocytes were Russell bodies. The practical and theoretical importance of nonsecretory immunoproliferative disorders, characterized by an unreleased monoclonal immunoglobulin marker, is outlined. —J. E. U.


Incorporation of radioactive precursors into protein, RNA and DNA was examined in PHA stimulated cultures of lymphocytes isolated from dairy cattle blood. Significant synthesis of protein and RNA was observed after 12 hr contact of PHA with lymphocyte cultures. Induction of DNA synthesis required 30-40 hr of PHA treatment. Experimental conditions for preparation and culture of bovine lymphocytes as well as for protein, RNA, and DNA synthesis determination are reported. —M. K.


An analysis of the distribution of blood groups in ABO system in 739 patients with mammary carcinoma showed deviations related to age of the patients and advancement of the neoplastic process. In women with advanced mammary carcinoma over 50 yr of age the A group was significantly more frequent and the O group was less frequent than in a group of lower age and in the general population of South Poland. The authors consider the possibility that the biological factor which controls the dynamics of the neoplastic process is the reason for blood group selection. —M. K.

The sera of 17 patients with Addison-Biernier disease containing antibodies type I (blocking) or type II (binding) or both against the intrinsic factor were examined. In 12 sera (70%) antiparietal cells activity was detected by an indirect immunofluorescent technique. In all cases antibodies against parietal cells were identified as IgG. Thyroid antibodies were found in sera of four (23%) patients. Antibodies against nuclear, mitochondrial, and smooth muscle antigens were not detected. M.K.


The effect of chronic poisoning with carbon disulfide on the peripheral blood was investigated in rats. Hemoglobin, hematocrit, erythrocyte, leukocyte and platelet counts were examined. It was found that changes in peripheral blood resulting from chronic CS₂ poisoning consisted in normocytic and normochromic anemia, eosinopenia, some increase in the percentage of reticulum cells and monocytes. No differences in white cell and platelet counts were observed. M.K.

The third annual meeting of the International Society for Experimental Hematology will be held at the Holiday Inn (Medical Center), Houston, Texas, from March 31 to April 3, 1974. The meeting is open to nonmembers. The opening ceremony will be held the evening of Sunday, March 31, the day after the close of the annual meeting of the American Association for Cancer Research, also being held in Houston. The program of the Experimental Hematology meeting will be organized from abstracts submitted by both members and nonmembers. Subjects include the control of differentiation and proliferation of both normal and malignant stem cells, bone marrow transplantation, prevention and control of graft-versus-host reactions, immunohematology, immunogenetics, cancer immunity, and applications of the former to the treatment of marrow aplasia, leukemia, lymphoma, and immune deficiency diseases.

Deadline for receipt of abstracts and pre-registration is December 20, 1973. Abstract forms and further information may be obtained from the meeting organizer and chairman, Dr. John J. Trentin, Division of Experimental Biology, Baylor College of Medicine, Texas Medical Center, Houston, Texas 77025.

Partial support of the meeting has been obtained from the Leukemia Society of America, and from the Texas Division of the American Cancer Society.

The annual meeting of the International Cooperative Group for Bone Marrow Transplantation will be held on April 4, 1974, also at the Holiday Inn (Medical Center). This year the meeting will consist of a workshop, organized by Dr. Mortimer Bortin, to develop cooperative clinical trial protocols.
ABSTRACTS