

The American Association for Thoracic Surgery 1972-1973

<i>President</i>	Frank Gerbode, <i>San Francisco</i>
<i>Vice-president</i>	Lyman A. Brewer, III, <i>Los Angeles</i>
<i>Secretary</i>	Myron W. Wheat, Jr., <i>Louisville, Ky.</i>
<i>Treasurer</i>	Paul C. Adkins, <i>Washington, D.C.</i>
<i>Editor</i>	Brian Blades, <i>Washington, D.C.</i>

<i>Council</i>	John W. Strieder (1973), <i>Newton Lower Falls, Mass.</i> Donald L. Paulson (1973), <i>Dallas</i> Wilfred G. Bigelow (1974), <i>Toronto</i> John K. Kirklin (1975), <i>Birmingham, Ala.</i> William S. Blakemore (1976), <i>Philadelphia</i>
----------------	--

<i>Membership Committee</i>	Thomas F. Nealon, Jr., <i>Chairman, New York</i> Frederick G. Pearson, <i>Ontario</i> Milton Weinberg, Jr., <i>Chicago</i> Benson B. Roe, <i>San Francisco</i> Harold C. Urschel, Jr., <i>Dallas</i> Bert W. Meyer, <i>Los Angeles</i> David B. Skinner, <i>Chicago</i>
-----------------------------	---

<i>Association representatives</i>	Rollin A. Daniel, Jr., <i>Nashville</i>
<i>The American Board of</i>	Benson B. Roe, <i>San Francisco</i>
<i>Thoracic Surgery</i>	Will C. Sealy, <i>Durham, N.C.</i> Herbert Sloan, <i>Ann Arbor, Mich.</i>

<i>Board of Governors,</i>	G. Rainey Williams (1975), <i>Oklahoma City</i>
<i>American College of Surgeons</i>	Lucius D. Hill (1973), <i>Seattle, Wash.</i>

American Association for Thoracic Surgery

53rd Annual Meeting

Scientific Program

MONDAY MORNING, APRIL 16, 1973

- 8:30 A.M. Business Session (Limited to Members)
Regency Ballroom**
- 8:45 A.M. Scientific Session:
Regency Ballroom**

**1. Tubed Gastric Pedicle for Esophageal Replacement in
placement in Children**

**KATHRYN D. ANDERSON* and JUDSON G. RANDOLPH,
Washington, D.C.**

The standard operation for esophageal replacement in children has been interposition of the colon. A favorable response in a child treated with gastric tube after failure of colon bypass, and the recent report by Stephens and Burrington, led us to adopt this operation as the primary approach for esophageal reconstruction. The tubed gastric pedicle has been evaluated in seven children. The patients include four children with extensive lye strictures and three infants with esophageal atresia without fistula. Age range of the patients at the time of surgery was eight months to four years. Patients have been followed from six months to five years.

Results: There have been no deaths or serious complications. Three patients developed mild strictures at the cervical anastomosis requiring several dilatations. No gastric tube has required revision. One patient developed an ulcer in the tube which responded to conservative therapy. All tubes have functioned satisfactorily to meet the nutritional needs of growing children. On the basis of this experience, a wider clinical trial of this procedure seems justified in children who need esophageal replacement.

*By invitation

2. Intrauterine Production of Coarctation of the Aorta: Operative Technique and Newborn Hemodynamic Studies

J. ALEX HALLER, JR., I. J. SHAKER,* ROBERT GINGELL*
and CHARLES HO,* Baltimore, Maryland

Clinical studies of coarctation of the aorta have contributed greatly to our understanding of this anomaly, but the high mortality of preductal versus postductal coarctation remains unexplained. Creation of these abnormalities in fetal lambs has provided an excellent opportunity to study the factors which affect survival, before and after birth, as well as the comparative differences in hemodynamic alterations.

Thirty ewes between 90-115/145 days gestation were operated upon and the fetal lamb was partially marsupialized to minimize amniotic fluid loss. Through a left thoracotomy, postductal coarctation was created in 13 lambs, and preductal in 17. Sixteen lambs were born alive either spontaneously or by C-section. The majority were studied angiographically to verify the site of coarctation and to demonstrate the extent of collateral circulation. Selected lambs received complete cardiac catheterization to assess the status of the ductus arteriosus and to measure the pressure gradient across the coarcted segment.

Operative techniques and data will be presented to show that intrauterine creation of preductal and postductal coarctation is possible with prolonged survival. Collateral circulation was remarkably similar in both groups and newborn survival was not affected by the location of the coarctation. We believe this is a useful model for definitive studies of altered hemodynamics in pre and postductal coarctation.

3. Surgical Repair of Single Ventricle

RICHARD N. EDIE,* KENT ELLIS,* WELTON GERSONY,*
FREDERICK O. BOWMAN, Jr., and JAMES R. MALM,
New York, New York

Single ventricle may occur with absence of the ventricular septum and two separate atrial ventricular valves associated with rotational variation in the origin of the great vessels. Four patients have successfully undergone repair of such an anomaly, two associated with Tetralogy of Fallot anatomy, one with double outlet in combination with pulmonary stenosis and one with D transposition and a pulmonary stenosis secondary to a pulmonary band. A technique of repair has been utilized to minimize residual shunting within the ventricle. The bundle of His is at risk in repair of this defect and direct mapping of its course has facilitated repair of the anomaly. The technique of His mapping will be presented in relation to single ventricle. Post operative studies are available on all patients. These demonstrate the complete prosthetic replacement of the ventricular septum can be carried out with near normal cardiac function post operatively.

*By invitation

4. The Fate of Reconstruction of the Right Ventricular Outflow Tract

SAMUEL KAPLAN,* JAMES A. HELMSWORTH,
GEORGE BENZING, III,* DAVID C. SCHWARTZ* and
J. TRACY SCHREIBER,* Cincinnati, Ohio

One hundred and thirty-eight patients with tetralogy of Fallot have been followed for 1-15 years after surgical correction. Reconstruction of the right ventricular outflow was accomplished with a pericardial patch in 45 patients and in another 11 instances an aortic homograft was used. In the remaining 82 patients, relief of obstruction was achieved by infundibulectomy alone with or without pulmonary valvotomy. Aneurysms of the right ventricular outflow developed in 19 patients, 18 of whom had pericardial patches. In all patients who developed aneurysms, residual significant defects were present. These consisted of (1) persistence of right ventricular hypertension because of inadequate relief of pulmonic stenosis or significant pulmonary arterial branch stenosis and/or (2) persistence of left to right shunt across the ventricular defect. It is concluded that right ventricular outflow patches or aortic homografts are well tolerated for many years after surgery (even in the presence of pulmonary valve incompetence) provided that the obstruction to right ventricular outflow has been relieved and the ventricular septal defect is closed.

5. Early Correction of Congenital Heart Disease with Surface Induced Deep Hypothermia and Circulatory Arrest

P. VENUGOPAL,* J. OLSZOWKA,* H. WAGNER,* P. VLAD,*
E. LAMBERT* and S. SUBRAMANIAN* Buffalo, New York
Sponsored by John W. Kirklin

In the past three years we have done open correction of congenital heart defects in 110 children using the Kyoto technique which consists of surface cooling with ice packs, a short period of perfusion followed by exsanguination and circulatory arrest. Core rewarming was done after surgical correction. Thirty-seven infants were below six months of age, 29 were between 7 and 12 months, 26 patients between 1 and 2 years and 18 were over 2 years. Eighty-nine weighed less than 10 kg and 43 infants were below 5 kg in weight.

The lesions included:

- a. Transposition—57 patients; 34 uncomplicated with 29 survivors. Twenty-three complex Transpositions with 15 survivors.
- b. Twenty-four children had VSD and associated anomalies and in 15 infants with isolated VSD there were no deaths.
- c. Tetralogy of Fallot in 10 infants with 2 deaths.
- d. Nine infants with Total Anomalous Pulmonary Venous Drainage with 5 survivors.

The use of this technique has permitted us to carry out complete correction successfully as early as 7 days of age. Surgical palliation is performed at this institution only in exceptional circumstances.

*By invitation

6. Surgical Correction of the Transposition Complex in Infancy

JAMES W. KILMAN, THOMAS E. WILLIAMS, JR.,*
GERARD S. KAKOS,* JOSEPHA CRAENEN* and
DON M. HOSIER,* Columbus, Ohio

Twenty-five patients with transposition of the great vessels have had total correction of this anomaly using an intra-atrial baffle and a triangular patch to enlarge the new left atrium. Only one death has occurred in this series resulting in a mortality of 4.0% (1/25). Associated defects included seven ventricular septal defects and five patients with pulmonary stenosis. These twelve defects have all been corrected at the same surgical procedure. Surgery was done using normothermic, high flow cardiopulmonary bypass with a standard disposable bubble oxygenator. The mean body weight of these patients is 8.9 kilograms. Fourteen patients were under 10 kilograms and no deaths have occurred in this group. The mean preop PaO₂ was 42 mm Hg. and the mean postop PaO₂ in room air was 76 mm Hg. Arrhythmias have not been a serious problem with this modified Mustard repair. A new maneuver for the exposure and closure of the ventricular septal defect has been used. It is felt that after this experience that transposition of the great vessels can be surgically corrected with only minimal risk in infancy.

7. Repair of Ventricular Septal Defect with Aortic Insufficiency

G. A. TRUSLER, Toronto, Ontario, Canada

Over the past five years, 16 children with ventricular septal defect (VSD) and aortic insufficiency (AI) have been treated by repair of the VSD and valvuloplasty of the affected aortic valve leaflet. The valvuloplasty transforms the elongated prolapsed leaflet into a competent leaflet with a free margin identical in length to that of the other aortic leaflets. Success hinges on precise measurement of the leaflet margin, secure fixation of the excess prolapsed leaflet to the aortic wall and reconstruction of the adjacent commissure.

Three of the 16 children had only slight to moderate relief of AI because the technique was inappropriate due to a bicuspid aortic valve in two and dilatation of the aortic ring in the third.

The other 13 children had a typical prolapsed leaflet (right 11, left 1 and non-coronary 1). The VSD was subcrystal in 10 children and supracrystal in 3.

Excellent results with no diastolic murmur, normal pulse pressure and decreasing heart size were obtained in 7 children. The other 6 children were improved but with mild to moderate aortic insufficiency.

Late studies, including angiocardiology, up to 4½ years after repair, show that improvement is maintained. AI due to a prolapsed valve leaflet can be safely and effectively repaired by this technique.

*By invitation

8. Progress and Problems in the Surgical Management of Congenital Aortic Stenosis

WILLIAM F. BERNHARD, DONALD C. FYLER,* KENNETH E. FELLOWS* and ROBERT E. GROSS, Boston, Massachusetts

During a fourteen year interval, 194 patients with valvar, subvalvar and supra-valvar aortic stenosis were operated upon. One hundred forty-six underwent valvotomy, including 33 infants (under one year). Although there were 11 infant deaths, twenty of the last 24 babies survived. Two deaths occurred among the remaining 113 children. Among the survivors, 80 had no significant aortic regurgitation (AR); however, AR was severe in ten, and mild to moderate in 43 patients. Follow-up LV-aortic pressures (available in 50 patients) revealed gradients less than 40 mmHg in 35. Of the remaining 15, two died with residual stenosis and five had successful valve replacement.

Forty children underwent partial resection of a subvalvar ring (five deaths). Sixteen survivors were recatheterized: gradients less than 40 mmHg were found in 14; nine had appreciable aortic or mitral incompetence.

Eight patients presented with supra-valvar AS (four with discrete lesions). The latter improved following operation; however, three with a hypoplastic ascending aorta and aortic annulus expired.

Conclusions: Valvotomy provides effective palliation in children with severe stenosis (including infants). Valve replacement was necessary in only five. Finally, left ventricular hypertension can be relieved in most patients with discrete obstructions below or above the valve.

11:15 A.M. Presidential Address

Frank Gerbode, San Francisco, California

COMPUTERIZED MONITORING IN THE
SERIOUSLY ILL PATIENT

MONDAY AFTERNOON, APRIL 16, 1973

2:00 P.M. Scientific Session
Regency Ballroom9. Diaphragm Pacing by Radiofrequency Transmission in
the Treatment of Chronic Ventilatory Insufficiency:
Present Status

W. W. L. GLENN, W. G. HOLCOMB,* J. HOGAN,*
I. MATANO,* J. B. L. GEE,* E. K. MOTOYAMA,* C. S. KIM*
and R. A. POIRIER,* New Haven, Connecticut

The indications for, complications of and long term results of diaphragm pacing in 15 patients with chronic ventilatory insufficiency will be reported. In 12 patients the respiratory center was affected (Ondine's curse); in 3 patients there was partial or complete severance of the upper cervical cord. The diaphragm pacemaker consists of a radiowave generator transmitting programmed signals via an antenna to a subcutaneous receiver connected to bipolar electrodes around one or both phrenic nerves. Receiver failure, initially troublesome, appears to have been corrected. Nerve fatigue is evident after 10-12 hours of continuous unidirectional stimulation but its onset may be delayed by utilizing a bidirectional stimulus.

In one patient pacing failed after 2 weeks. In the remaining patients adequate ventilation of the lungs, as evidenced by spirometry, normal blood gas concentration and Xenon¹³³ diffusion studies, has been achieved up to 44 months without evidence of injury to the phrenic nerve; the phrenic nerve stimulation threshold rose initially but stabilized within 6 months usually at between 1 and 2 milliamperes. Right heart failure and pulmonary hypertension, when present, improved during diaphragm pacing. Evidence to date indicates that pacing of the diaphragm is an effective, practical and reasonably safe technique for long term partial or complete ventilatory support.

10. Development of an Implantable Artificial Lung

ARTHUR PALMER,* JOHN COLLINS* and LOUIS R. HEAD,
Chicago, Illinois

A membrane oxygenator, which can be implanted in the chest, has been developed to function as an artificial lung. It operates at pulmonary artery pressure and requires no external power source for ventilation. Silastic capillary tubes function as the membrane surface. These are manifolded in parallel into modules containing 2288 to 6270 individual tubes. The modules are then sealed in a Silastic ventilating envelope which is connected to an artificial bronchus. Blood flows through the capillary tubes and the gas in the ventilating envelope is continuously changed by the normal motion of the dogs chest.

Ten artificial lungs have been implanted in dogs. Blood flow through the prosthesis ranged from 80-260cc/min at mean pulmonary artery pressures of 10-25mm. of Hg. Oxygen transfer ranged from 15-23cc/min/M² at physiologic levels of pO₂, pCO₂ and ph.

The major problem remaining is prevention of intravascular coagulation in the artificial lung. Regional heparinization is presently used and various antithrombogenic surface treatments are under investigation.

*By invitation

11. Delayed Sequelae of Penetrating Cardiac Wounds

P. N. SYMBAS, DAVID A. DiORIO,* D. H. TYRAS,*
R. E. WARE* and CHARLES R. HATCHER, Jr.,
Atlanta, Georgia

In order to determine the course and sequelae of penetrating cardiac wounds, the cases of 76 patients with such injuries treated at Grady Memorial Hospital from July 1964 through June 1972 were reviewed. Of these 76 patients, 56 survived the immediate postinjury period with no further mortality attributable to their injury during an average 19 month follow-up period. A variety of anatomic and physiologic sequelae of the cardiac wounds were encountered, including 5 ventricular aneurysms (3 pseudo, 1 true, and 1 undefined aneurysm), 3 ventricular septal defects, 3 instances of valvular incompetence (2 mitral, 1 pulmonic), 2 aortopulmonary fistulae and 2 instances of bullets retained within the interventricular septum. With the exception of the 2 patients with ventricular septal defects and both patients with aortopulmonary fistulae, none of these patients were more than mildly symptomatic and most were totally asymptomatic. Operative intervention has been carried out without mortality and with excellent results in the 4 patients with marked symptoms (2 VSD and both AP fistulae) and in 4 patients with ventricular aneurysms (3 pseudo and 1 true).

This experience re-emphasizes the need for close follow-up of patients sustaining penetrating cardiac wounds and repair of the delayed sequelae of such wounds when either hemodynamically significant or potentially dangerous to the patient.

12. Air Embolism Following Penetrating Lung Injury

ARTHUR N. THOMAS,* San Francisco, California
Sponsored by Benson B. Roe

The literature does not mention the occurrence of air embolism after penetrating lung injury. Nevertheless we have documented coronary air embolism after penetrating lung injury in three patients. We believe that this was responsible for difficulty in resuscitating the patient in one instance and resulted in death in two.

In order to evaluate the mechanism and significance of air embolism penetrating lung injuries were created in 14 anesthetized dogs. A polyethylene shunt between the aortic root and femoral vein was used as an air bubble trap and detector. The lung was penetrated with a #22 scalpel blade. Air immediately appeared in the shunt in 10 of 14 dogs. Airway, pulmonary artery, right ventricle, left atrial and femoral arterial pressures were measured in animals receiving controlled ventilation.

Air caused rapid death from coronary embolism in 6, and caused hemodynamic changes that spontaneously ceased in 4. Lung injury produced no detectable air embolism in 4 dogs. A communication between the airway, pulmonary artery and pulmonary vein was shown in all lung wounds. The sequential hemodynamic changes measured were left ventricular failure, left atrial hypertension, pulmonary arterial hypertension and systemic hypotension.

The conclusion is that air embolism is a potential hazard following lung wounds. It is most apt to occur in patients who are in shock and require positive pressure ventilation.

*By invitation

13. The Role of Bronchial Brushing on the Decision for Thoracotomy

J. J. FENNESSY* and C. F. KITTLE, Chicago, Illinois

More than 600 transcatheter or bronchial brush biopsies have been performed at The University of Chicago Hospitals between 1965 and 1972. Tissue thus secured was examined histologically and cytologically, and cultured for fungus and bacteria. In most of these patients the indication for the procedure was the presence in the lung of a lesion suspected to be malignant. Flexible fiber optic bronchoscopy has been done for the past 3 years in these patients.

This presentation is a retrospective analysis of the clinical and radiologic data available on patients subjected to bronchial brush biopsy and flexible bronchoscopy to determine to what extent, if any, the results of the procedure influenced the surgeon's decision to operate.

These patients have been analyzed by reviewing the clinical data pre-brushing to determine if the surgeon would advise thoracotomy. A retrospective analysis was also done by both authors to determine if brushing was helpful in the overall management.

Definite indications for brushing have been established: the possibility of inflammatory disease and a need for culture material, the question of metastatic disease with multiple nodules, and the opportunity of obtaining a tissue diagnosis for x-ray therapy when other conditions precluded operation. In many instances "brushing" has been of supplemental interest only without decision-making importance.

14. Parasternal Mediastinotomy—A Useful Adjunct in the Diagnosis of Chest Disease

PHILIP C. JOLLY,* LUCIUS D. HILL, THOMAS WEST*
and PETER LAWLESS,* Seattle, Washington

One hundred consecutive cases of parasternal mediastinotomy have been compared to 240 cases of mediastinoscopy. Distant metastases were not present in these patients to contraindicate thoracotomy or allow easy tissue diagnosis. There was no mortality and morbidity was low after both procedures. Sixty-eight percent of the patients had carcinoma of the lung. In those patients with lung cancer tissue diagnosis was obtained by mediastinotomy in 69% and by mediastinoscopy in 32%. Undifferentiated tumors yielded a higher diagnostic return in both groups. Mediastinotomy proved superior to mediastinoscopy in evaluating patients for resectability. Resectability rates at thoracotomy correlated accurately with the findings at mediastinotomy and mediastinoscopy. Of the malignant cases, thoracotomy was avoided in 62% by mediastinotomy and in 30% by mediastinoscopy.

Parasternal mediastinotomy is a simple, versatile procedure. Lung biopsies were obtained in 22 patients yielding a diagnosis in a variety of chest diseases. Therapeutic procedures such as pericardial window, excision of pericardial cysts and placement of epicardial electrodes for pacemaking are possible through this incision.

*By invitation

15. Delayed Cutaneous Hypersensitivity Reactions to Tumor Antigens and to Non-Specific Antigens: Prognostic Significance in Patients with Lung Cancer

SAMUEL A. WELLS,* Durham, North Carolina
JAMES F. BURDICK* and CHRISTINE CHRISTIANSEN,*
Bethesda, Maryland, WILLIAM L. JOSEPH,* WALTER G.
WOLFE* and PAUL C. ADKINS, Washington, D.C.

Oncogenesis is favored by an environment of depressed immunity, but there are few studies in humans correlating both general immunological status and reactivity to tumor specific antigens with the patient's clinical course.

Delayed cutaneous hypersensitivity reactions (DCHR) to bacterial antigens (mumps, candida and streptokinase-streptodornase) and to a previously unencountered antigen, dinitrochlorobenzene (DNCB), were evaluated in 100 ambulatory patients, 75 with lung cancer and 25 with benign lung disease. Eighteen cancer patients were also tested with membrane antigen extracts (MAE) of autologous tumor tissue.

Twenty-four patients with benign disease had positive DCHR to both bacterial antigens and DNCB. In the 75 cancer patients 72 developed DCHR to bacterial antigens, but reactivity to DNCB was markedly depressed with only 40 patients reacting and in 12 patients with non-resectable disease only 2 reacted. Eight of 18 patients developed DCHR to autologous MAE of lung tumor, but not to normal lung. Seven of these patients were well at eight months, while only 3 patients with negative DCHR to tumor MAE were alive without recurrent disease.

These data demonstrate that in lung cancer patients, a poor prognosis is associated with a depressed immune recognition of DNCB and negative cutaneous reactivity to autologous tumor MAE.

16. Segmental Resection for Lung Cancer—A 15-Year Experience

ROBERT J. JENSIK, L. PENFIELD FABER, FRANK J.
MILLOY* and DAVID O. MONSON,* Chicago, Illinois

One hundred fourteen patients with primary lung cancer underwent segmental resective surgery over the past 15 years. The patients were placed in the following groups:

- I. In 14 patients, previous contralateral resective surgery had been carried out;
- II. In 26 patients, the procedure was done for palliation;
- III. In 74 patients, the resection was considered as a definitive curative operation.

Four patients of Group I survived more than two years with the longest survivor still alive seven plus years later.

Only five patients survived more than two years in the palliative group, the longest attaining a five-year survival.

A 55% five-year survival calculated by actuarial method was achieved in Group III. This declined to 21% over the 15-year period.

Tumor histology and location, the type of segmental procedures, and factors influencing the decision for limited resection will be discussed.

An operative mortality of 5% to 6% with a 55% five-year survival suggests that segmentectomy may be the procedure of choice when indicated.

*By invitation

TUESDAY MORNING, APRIL 17, 1973**8:30 A.M. Scientific Session
Regency Ballroom****17. A New Technique of Treating Esophageal Varices**

MITSUO SUGIURA* and SHUNJI FUTAGAWA,* Tokyo, Japan
Sponsored by John E. Connolly

The high incidence of encephalopathy and progressive hepatic failure following portosystemic shunts for esophageal varices led us to abandon this technique. From 1964 to 1967 26 patients underwent simple transection of the thoracic esophagus. Recurrence of variceal bleeding was noted in 4 and esophageal varices disappeared radiographically in only 15 patients. The excision of coronary veins plus esophageal transection in 14 patients did not improve the late results.

Our current technique has evolved from these earlier experiences and consists of extensive paraesophageal devascularization up to the tracheal bifurcation, transection of the distal thoracic esophagus, splenectomy, devascularization of the abdominal esophagus and cardia, selective vagotomy and pyloroplasty. Thoracic and abdominal operations are performed through separate incisions and can be done in two stages in poor-risk patients. Since 1967, 74 patients have undergone this new procedure, 14 emergency, 48 elective, and 12 prophylactic. The causes of esophageal varices were: cirrhosis 41, fibrosis 22, extrahepatic portal vein occlusion 7, hepatoma 3, and carcinoma of the pancreas 1. Overall operative mortality was 5.4%. A five-year followup study revealed that six patients died of hepatoma, the esophageal varices disappeared in all cases, and all 60 survivors were free from encephalopathy.

Although a longer follow-up is necessary, our preliminary results are encouraging and warrant further trial.

18. Surgical Management of Scleroderma of the Esophagus

R. D. HENDERSON* and F. G. PEARSON, Toronto,
Ontario, Canada

Misconceptions persist concerning the origin, and treatment of dysphagia in patients with scleroderma. This report of current experience clarifies aspects of pathogenesis, and an approach to treatment is suggested.

In the past four years, we have studied 22 patients with scleroderma, 16 of whom had significant complaints of reflux and dysphagia. In 6 of the 16 symptomatic patients a diagnosis of scleroderma was first made following our clinical assessment and esophageal motility studies. Nine of 16 patients with dysphagia had typical peptic strictures.

Twelve patients were treated surgically: Gastroplasty and Belsey hernia repair—9, Belsey repair—2, Thal esophagogastroplasty—1. Cine barium studies were done in all patients before and after operation. Motility studies were done in all 12 patients pre-operatively, and in 7 patients post-operatively.

Good results were obtained in 10 patients: 5 are completely asymptomatic; 3 have slight pharyngoesophageal motor dysphagia; 1 has slight gastroesophageal

*By invitation

mechanical dysphagia; and 1 patient has symptomatic reflux, but relief of dysphagia. One patient was unimproved. There was one operative death.

There is no evidence that scleroderma results in esophageal stricture, other than by predisposing to reflux. Correction of reflux with dilatation of strictures gives good symptomatic relief. No healing problems were noted.

19. Reappraisal of Adjuncts to Avoid Ischemia in the Treatment of Descending Thoracic Aortic Aneurysms

E. STANLEY CRAWFORD, Houston, Texas

Paraplegia is a dreaded complication associated with excisional therapy of aneurysms involving the descending thoracic aorta. Various methods or adjuncts have been devised to prevent this complication, including bypass shunts of various types and hypothermia. These procedures and improved surgical techniques have reduced this complication to a minimum and it is difficult to determine which is more important. This paper is concerned with a consecutive series of 80 patients in whom descending thoracic aortic aneurysm were removed. Bypass shunts were employed in the first 38 patients. The last 42 patients were treated without special protective mechanisms. Paraplegia was present after operation in 3 of the former and in one of the latter patients. Of the 36 patients undergoing operation electively without shunts, none developed paraplegia and 94.5 per cent survived despite being over sixty years of age in most cases. This experience would tend to require reappraisal of need for shunts, and this paper will deal both with this and other parameters in the conduct of operation felt to be more important.

20. Patterns of Myocardial Metabolism During Cardiopulmonary Bypass (CPB) and Coronary Perfusion

O. WAYNE ISOM*, NEIL D. KUTIN*, EMILY A. FALK*
and FRANK C. SPENCER, New York, New York

Much uncertainty prevails about methods of coronary perfusion, tolerance for ischemia, and fibrillating vs beating heart. Therefore, in 35 patients undergoing CPB (30°C) myocardial metabolism was studied during operation and for up to 70 hrs. Paired samples of arterial and coronary sinus blood, obtained from indwelling catheters, were analyzed for PO₂, PCO₂, pH, lactate, and enzymes—CPK, LDH, SGOT. In 15 pts undergoing aortic valve replacement, coronary flow rates and myocardial oxygen consumption were also measured. The data, statistically analyzed for over 20 variables, were as follows:

Coronary blood flow (CBF) was 200-300 ml/min, oxygen extraction ($\Delta A-V$) 2.5-3.0 vol %; MVO₂ 6-8 ml/min, about 20 per cent of normal. The table shows little difference between fibrillating (F) and non-fibrillating (NF) hearts, except greater enzyme production in F hearts ($p < 0.05$). Lactate extraction occurred in about one-half of each group.

*By invitation

	# pts	MVO ₂ ml/min	A-V O ₂ vol %	E O ₂	% pts ext. Lactate	CPK	LDH	SGOT
NF	9	7.96	3.05	25%	56	206	345	125
F	6	6.13	2.7	24%	44	339	644	127

Marked lactate production (Δ A-V 85 mg%) occurred with 15 min ischemic periods. Metabolic recovery from shorter ischemic periods occurred in 5-10 min, but longer periods caused permanent alterations of oxygen consumption. An "overperfusion" injury was identified, characterized by localized edema, normal lactate metabolism, and severe arrhythmias (1-3 hrs duration). Coronary perfusion exceeding 300 ml/min led to significantly greater enzyme production than did lower perfusion rates (CPK 344 vs 172, $p < 0.01$).

Limitations of metabolic studies were found in one patient with a short left main coronary who developed a fatal myocardial infarction. CPK production occurred (A-V CPK 125 units) but lactate metabolism was near normal. The relationship of these data to the pathogenesis of hemorrhagic subendocardial necrosis will be discussed.

21. Anoxic Cardiac Arrest: An Experimental and Clinical Study of its Effects

S. R. K. IYENGAR,* E. J. P. CHARRETTE* and
R. B. LYNN Kingston, Ontario, Canada

"It is important to know not only the survival but also the quality of life after anoxic cardiac arrest," commented Dr. Gerbode in the Thoracic Surgery Forum of the last meeting of the Association. We have been engaged in the study of this problem for the past two years. During phase I of the investigation a correlation between the incidence of subendocardial hemorrhagic necrosis in dogs and anoxic cardiac arrest was established. Simple flushing of the coronary bed during anoxic arrest of 60-75 minutes with a balanced electrolyte solution, designated as "Beks" solution in our laboratory prevented the subendocardial lesions and there was no intraoperative death from low output. Impressed by these results, we have been routinely flushing the coronary bed with the "Beks" solution during aortic valve replacement. The results are being assessed.

In patients with aortic valvular stenosis, it is believed that in addition to left ventricular hypertrophy, the coronary blood flow is interfered with. In the second phase of our programme a canine model in which sub-coronary aortic stenosis was produced to simulate the situation in humans was produced to study the effects of anoxic arrest. The hypertrophied left ventricle was much more vulnerable to anoxic arrest. Whereas the lesions were subendocardial in the normal canine heart, they were extensive and deep in the hypertrophied left ventricle. At present investigation is in progress to study and improve the quality of myocardial function after anoxic arrest in long term canine survivors.

Myocardial damage incidental to anoxic arrest during open heart surgery adversely affects the quality of life and diminishes the maximum benefit that could otherwise be expected. Pharmacologic manipulation and intermittent flushing of the coronary bed during anoxic arrest offers a simple technique which deserves more extensive clinical trial.

*By invitation

22. Profound Local Hypothermia for Myocardial Protection During Open Heart Surgery

RANDALL B. GRIEPP,* EDWARD B. STINSON* and
NORMAN E. SHUMWAY, Stanford, California

Between 7/71 and 6/72, 329 adults underwent cardiac valve replacement (153), or aortocoronary bypass grafting (133), or both (43). The aorta was crossclamped during valve replacement and during the performance of distal coronary artery anastomoses. Local profound hypothermia during aortic cross-clamping was provided by a continuous infusion of normal saline at 4°C. into the pericardial cavity. Coronary artery perfusion was not used. Cardiopulmonary bypass times ranged from 40 to 200 minutes (average 105 minutes), and aortic crossclamp times ranged from 30 to 140 minutes (average 62 minutes).

Twelve hospital deaths occurred, yielding an operative mortality of 3.6%. Six patients could not be weaned from cardiopulmonary bypass; three died of low cardiac output postoperatively, and three died as the result of factors not related to myocardial function. Average stay in the intensive care unit was 2.5 days, and average hospital stay was 10 days. No correlation is evident between duration of aortic crossclamping and morbidity or mortality.

This experience suggests that profound local hypothermia during aortic crossclamping affords excellent protection of the myocardium during the performance of cardiac valve replacement and aortocoronary bypass grafting.

23. The Hazard of Ventricular Fibrillation in Hypertrophied Ventricles During Cardiopulmonary Bypass

CHRISTOF E. HOTTENROTT,* HENRY J. KURKJI,* JAMES V.
MALONEY and GERALD D. BUCKBERG,* Los Angeles, California

We previously reported left ventricular ischemic damage occurs in most patients dying after cardiopulmonary bypass, and defined pre- and postoperative factors contributing to this injury. This study provides evidence that any form of ventricular fibrillation during bypass impairs coronary flow to the hypertrophied left ventricle and causes ischemia.

Seven dogs with left ventricular hypertrophy (aortic stenosis 3-5 months) were compared to fifteen normal dogs to determine how spontaneous^{*} ventricular fibrillation (60 minutes) during cardiopulmonary bypass affects: (1) myocardial function (Sarnoff curves), (2) regional coronary flow (radioactive microspheres), (3) cardiac biochemistry (pH, lactate, potassium) and (4) histochemistry (acid fuchsin).

In normal ventricles, spontaneous fibrillation raised left ventricular oxygen consumption and subendocardial flow and lowered vascular resistance ($P < .01$). It did not impair myocardial function and biochemistry nor cause histochemical damage. Conversely, when hypertrophied left ventricles fibrillated spontaneously, oxygen consumption failed to rise, vascular resistance progressively increased, and biochemical evidence of severe ischemia occurred (myocardial lactate production, decreased coronary sinus pH, and loss of intracellular potassium, $P < .01$). Post-bypass ventricular function was depressed and histochemical ischemia was demonstrated. Clinical studies during aortic valve replacement confirmed these experimental findings.

*By invitation

This study shows that while spontaneous fibrillation may be safe in normal hearts, it is hazardous in hypertrophied hearts.

24. Evaluation of Functional, Metabolic and Structural Alterations in the Myocardium During Aortic Cross-clamping

EDWARD A. STEMMER, PETER McCART,* WILLIAM E. STANTON,*
WILLIAM THIBAULT* and JOHN E. CONNOLLY, Irvine, California

Inability of the myocardium to support satisfactory circulation is the most frequent cause of death after open heart procedures. Knowledge of the interrelationships between metabolism, structure and function of the myocardium can prevent deterioration of myocardial function and refractory myocardial failure.

The ability to maintain normal myocardial metabolism, function and ultrastructure was evaluated in 70 dogs undergoing cardiopulmonary bypass with sustained fibrillation, ischemic arrest, topical hypothermia, intermittent coronary perfusion or continuous coronary perfusion. Survival, arterial pressure, central venous pressure, left ventricular pressure, cardiac output, dp/dt, urinary output, myocardial oxygen consumption, myocardial potassium loss, lactate utilization, pyruvate metabolism, light microscopy and electronmicroscopy were studied in each animal during a baseline period, two hour test period and two hour recovery period.

The best survival with the least impairment of myocardial function was observed with continuous hypothermic coronary perfusion. The poorest survival with the greatest impairment of function occurred after normothermic anoxic arrest for more than 60 minutes. Topical hypothermic arrest without coronary perfusion produced good survival but significantly compromised myocardial function. Electronmicroscopy demonstrated that damage to the mitochondria was associated with poor survival, elevated coronary sinus lactate and poor recovery of myocardial function. Anoxia aggravated these changes while coronary perfusion minimized them.

11:15 A.M. Address of Honored Speaker

**Sir Thomas Holmes Sellors
President, Royal College of Surgeons**

THE GENERALITY OF SURGERY

*By invitation

TUESDAY AFTERNOON, APRIL 17, 1973

2:00 P.M. Scientific Session
Regency Ballroom

25. Drug Influence on Platelet Loss During Extracorporeal Circulation

C. H. MIELKE, JR.,* M. deLEVAL,* J. D. HILL,*
M. F. MACUR* and F. GERBODE, San Francisco, California

Thrombocytopenia, which develops during prolonged extracorporeal circulation, represents the major hazard to this procedure. The potential for serious hemorrhage during prolonged perfusion has led to investigations of the influence of various drugs on platelet loss.

Using labeled platelets (^{51}Cr) in dogs, we were able to show that the majority of this platelet loss occurs because of storage in the liver during bypass. Once bypass is discontinued, some of the sequestered, labeled platelets return to the circulation with a corresponding reduction in liver radioactivity. Circulating fibrinogen labeled with ^{125}I remains stable.

Using this animal model we have evaluated several drugs with known influences on platelet function in both the Temptrol bubble and Bramson membrane oxygenators. The influences of Persantine, Aspirin, Sudoxicam and Pluronic F68 were compared on platelet loss during and after bypass.

Persantine inhibited platelet adhesion but not aggregation. Platelet levels were only slightly diminished during and after bypass. Pluronic F68 was similar but less effective. Platelet microaggregation during bypass was effectively inhibited by both agents. Sudoxicam inhibited both platelet aggregation and adhesion. However, the post bypass level of platelets was lower than with the two other agents. Platelet consumption was increased by high doses of aspirin.

26. Electroencephalographic Changes and Cerebral Complications in Open-Heart Surgery

M. WITOSZKA,* H. TAMURA,* R. INDEGLIA* and
F. A. SIMEONE, Providence, Rhode Island

Cerebral dysfunction and behavioral disorders are not uncommon after surgical correction of cardiac lesions. In the past four years encephalograms (EEG) were continuously monitored during open heart surgery in 50 randomly selected patients who survived (Group I) and 50 patients who succumbed during (Group II) and after the procedure (Group III). These data were correlated with clinical evidence of neurological disorders and findings in the brain at autopsy.

*By invitation

Results:

Group	Total		Hypotension		Encephalopathy		Motor Changes		Total Neurological Changes	
	No.	Percent	No.	Percent	No.	Percent	No.	Percent	No.	Percent
Group I	50		5	10%	26	52%	12	24%	26	52%
EEG Changes	7	14%	3	42%	5	71%	3	42%	5	71%
No EEG Changes	43	86%	4	9%	22	51%	10	23%	22	51%
Group II	5									
EEG Changes	5	100%	5	100%						
No EEG Changes	0		0							
Group III	45		17	37%	28	62%	23	51%	36	80%
EEG Changes	25	55%	13	52%	20	80%	14	56%	23	92%
No EEG Changes	20	44%	4	20%	8	40%	9	45%	13	65%

Postmortem examinations of the brain were performed in 20 patients. Among 18 of these patients with significant EEG changes fifteen had abnormal findings upon histologic examination of the brain. Evidence of cerebral embolism was found in 60% of autopsy specimens. No correlation between hypotension and histologic abnormalities was observed.

Conclusions:

1. Neurological complications occurred in 52% of patients who survived open heart surgery and in 80% of patients who died.
2. These complications followed intraoperative EEG abnormalities in 28% cases and 35% in the absence of intraoperative EEG changes, postoperative encephalopathy or motor disorders occurred in 62% of the patients.
3. Neuronal degeneration was the most common pathological manifestation of the encephalopathic syndrome, while frank cerebral necrosis (8 cases) was most often associated with cerebral embolism (80% of the cases).

27. Surgical Experience with Temporary and Permanent A-V Sequential Demand Pacing

JOSH FIELDS,* BAROOTH V. BERKOVITS* and
JACK M. MATLOFF, Los Angeles, California

The hemodynamic advantages of the normal atrioventricular sequence can be maintained in patients with atrial bradycardia-arrhythmias and/or complete heart

*By invitation

block by the temporary or permanent use of atrioventricular sequential demand pacing. This pacing modality is achieved with a two-catheter system utilizing a conventional bipolar ventricular catheter and a new, preformed J-shaped bipolar atrial electrode. Temporary use in 12 postoperative cardiac surgical patients with sinus or nodal bradycardia (with and without A-V block) yielded a 12-22 mm Hg increase in blood pressure over that achieved with ventricular pacing. In each case of escape tachyarrhythmia or ectopic activity, the use of A-V sequential pacing achieved capture and maintenance of a stable rate and rhythm *without* large doses of suppressant drugs.

Permanent A-V sequential demand pacemakers have been implanted for up to three years in 49 patients, 6 following prosthetic valve replacement and 43 with intermittent or constant atrial arrhythmias (17) and bradytachy-arrhythmias (26). Fifteen of these patients also had A-V dissociation. There was dramatic clinical improvement in 44 of these patients, manifest by relief of syncope, control of tachyarrhythmias and relief of congestive heart failure. This latter result was achieved by virtue of increased cardiac outputs. From this experience, it would appear that A-V sequential demand pacing provides the most comprehensive modality of pacing yet available.

28. Experience with Atrial Pacemaker Wires Implanted During Cardiac Operations

NOEL L. MILLS* and JOHN L. OCHSNER, New Orleans, Louisiana

Two-hundred and seventy-five patients had implantation of a pacemaker wire on the right atrium at the time of cardiac operation. The operations involved were repair of congenital heart defects and valvular and coronary bypass procedures. Forty-three patients had arrhythmias in the postoperative period—the most common being atrial fibrillation, ventricular premature beats, and atrial flutter. A unipolar wire was implanted in all except 8 patients. Bipolar wires were used for patients who had sinus bradycardia at the conclusion of the operation and left while being atrially paced.

The atrial wire was used also in the postoperative period as an exploring electrode to diagnose atrial arrhythmias. Connection of the atrial wire to the chest electrode of the standard electrocardiograph obtains such information that proper diagnosis and prompt treatment could be instituted. In addition, the electrode was used as an atrial pacer for conversion of flutter to normal sinus rhythm, for sinus bradycardia, and for evaluation of coronary bypass grafts.*

In 13 patients implantation of the wire was unsuccessful and in 4 the wires were broken on removal. No early or late sequelae were observed. From our experience we advocate the routine use of atrially implanted electrodes for diagnostic and therapeutic use after cardiac operations.

3:30 P.M. Executive Session (Limited to Active and Senior Members)
Regency Ballroom

*By invitation

TUESDAY EVENING, APRIL 17, 1973

**7:00 P.M. President's Reception
 Regency Ballroom**

**8:00 P.M. President's Dinner and Dancing
 Regency Ballroom**

Attendance open to all physicians and their ladies. Tickets must be obtained at the Registration Desk by 5:00 P.M. on Monday, April 16, 1973.

Dinner Dress Preferred

WEDNESDAY MORNING, APRIL 18, 1973

8:30 A.M. Scientific Session
Regency Ballroom

29. Liquid Membrane Oxygenator

HERBERT W. WALLACE,* MARC T. ZUBROW,* HELENE BROOKS,*
WILLIAM J. ASHER,* NORMAN N. LI* and T. PETER STEIN,*
Philadelphia, Pennsylvania

Sponsored by William S. Blakemore

A new concept of blood oxygenation based on the encapsulation of gas bubbles within a thin film of inert fluorochemical avoids a blood-gas interface but allows adequate gas transfer. The encapsulated bubbles are passed countercurrent to the blood flow. Oxygen passes through the liquid membrane into the blood, and CO₂ takes the reverse course. The bubbles emerge from the blood phase and collapse, releasing CO₂. The fluorochemical is reused. *In vitro* experiments demonstrated the method's feasibility (estimated O₂ transfer ≥ 100 cc/min/m²). The compalibility of fluorochemical and blood was evaluated with a device designed to generate a continuous blood-fluorochemical interface. There were no measured untoward effects of fluorochemical upon human blood (24 parameters studied) during 24-hour exposure of over 52 m² of interface at 25°C. *In vivo* all ten dogs survived 4 hours of veno-venous perfusion (200 cc/min). No alterations of 24 blood parameters occurred acutely or during a four- to six-week period of follow-up. The animals were sacrificed, and no gross or microscopic alterations of organs were detected. A prototype oxygenator is now undergoing *in vitro* and *in vivo* evaluation. The initial results have been promising and will be discussed.

30. The Lande'-Edwards Membrane Oxygenator During Heart Surgery: Oxygen Transfer, Microemboli Counts and Bender Gestalt Visual Motor Test Scores

ROBERT G. CARLSON,* ARNOLD J. LANDE',* JAMES BAXTER,*
RUSSELL H. PATTERSON, JR.,* and C. WALTON LILLEHEI,
New York, New York

In 130 patients, 7-103 Kg, the Lande'-Edwards Membrane Oxygenator provided simple, safe, total cardiopulmonary support (3M² membrane/40 kg). Venous blood drained through the membranes into a reservoir by gravity only and was pumped back into the patient.

Maximum oxygen transfer was 52/ml/min/M² of membrane, range 12-52 depending on temperature and body weight. At moderate hypothermia (30°C), venous saturation was 85% and arterial 100%. Partial support was provided during warming in larger patients to reduce the number of oxygenator units needed. At mild hypothermia (33-36°C) maximum oxygen transfer measured was 312 ml/min/6M² membrane (range 35-53).

Decreased morbidity of the membrane compared with the bubble oxygenator was evidenced by decreased microemboli counts, e.g., comparable counts per minute with membrane were 1500 v.s. 18,000 with bubble oxygenator. Use of a

*By invitation

microfilter (Pall) reduced these counts by 90% in both. Postoperative visula motor function (Bender-Gestalt) deteriorated in 9% (membrane) and 40% (bubble) patients. This membrane provided safe total cardiopulmonary support for heart surgery and is recommended for decreased morbidity particularly in complex operations.

31. Intraaortic Balloon Assist for Postcardiotomy Cardiogenic Shock

R. L. BERGER and V. K. SAINI,* Boston, Massachusetts

Intraaortic balloon pump (IABP) support was provided in eleven patients with cardiogenic shock following coronary artery bypass grafts, resection of left ventricle, mitral valve surgery or a combination of these operations. Nine of the eleven could not be weaned from cardiopulmonary bypass (CPB) as evidenced by a systolic pressure of less than 75 mm. of Hg. and a left atrial pressure of greater than 24 mm. of Hg. in spite of maximal volume and pressor therapy. Institution of IABP converted the nonpulsatile flow of CPB into a pulsatile one, raised the arterial and lowered the left atrial pressures and allowed discontinuation of CPB. Eight of the nine patients left the operating room and four became long term survivors. Aortocoronary bypass graft flow was measured with electromagnetic probes in one patient and IABP produced a 60% increase in flow. Two of the eleven patients sustained cardiac arrest on the first and second postoperative days and remained in deep shock following resuscitation. IABP support produced initial improvement but ultimately both died. Extensive documentation of clinical, hemodynamic and metabolic changes were obtained in all cases. This experience indicates that IABP can be instrumental in salvaging postcardiotomy cardiogenic shock patients.

32. Objective Assessment of the Effects of Aorto-Coronary Bypass Operation on Cardiac Function

HOOSHANG BOLOOKI,* LEONARD SOMMER,* STEVEN MALLON,*
ABELARDO VARGAS* and MICHAEL GILL,* Miami, Florida
Sponsored by Gerard A. Kaiser

Controversy exists as to the actual effects of direct myocardial revascularization on cardiac function. In order to evaluate this subject more precisely, we have studied eleven parameters of cardiac function (as a pump and as a muscle) and left ventricular compliance in twenty consecutive patients before and within 4-10 months after successful myocardial revascularization. Eight patients were operated because of acute cardiac ischemia—preinfarction angina (Group A) and twelve had surgery because of chronic intractable angina pectoris (Group B). Postoperatively, all these patients were free of angina and had returned to work; also arteriograms had shown patency of all grafts. After surgery, cardiac index (CI), myocardial contractility (V_{max} , $dp/dt/Kp$), left ventricular end-diastolic pressure (Edp) and compliance (dp/dv) among other parameters, showed insignificant changes in either group. Statistical analysis of data was carried out in a number of ways using various classifications. There was an increase in V_{max} in 4 of 8 (50%) of patients in Group A as opposed to 3 of 12 (25%) of patients in

*By invitation

Group B. Left ventricular compliance was unchanged in Group A, but had decreased by 50% or more in one-half of patients in Group B. CI had increased by 25% or more in 6 of 7 patients (in both groups) who had a control CI below 2.5 L/min/m² ($P < 0.02$). In this group, however, the stroke volume remained unchanged. These results indicate that direct myocardial revascularization produces an excellent palliation of anginal symptoms, but the postoperative improvement in cardiac function is most likely to occur in that group of patients who are suffering from acute cardiac ischemia.

33. Preinfarction Angina Pectoris—A Surgical Emergency

R. R. GOODIN,* T. V. INGLESBY,* A. M. LANSING *
and M. W. WHEAT, JR., Louisville Kentucky

From November, 1971, to September, 1972, 19 patients with preinfarction angina pectoris, 11.3% of patients studied because of coronary atherosclerosis, were studied by cardiac catheterization and coronary arteriography. All patients were candidates for aorto-coronary bypass surgery if technically feasible.

Twelve patients had saphenous vein aorto-coronary bypass graft surgery with one death and good clinical results in the 11 survivors. Seven patients were not operated upon and in five of these patients there have occurred 3 deaths and 2 non-fatal myocardial infarctions during the first three months of follow-up.

From a clinical standpoint, severity and duration of pain, frequency of pain and EKG changes, the two patient groups could not be differentiated. Fifty percent of operated and 57% of non-operated patients had had previous myocardial infarctions. The average number of vessels with over 50% occlusion was 2.17% in operated and 2.85% in non-operated group. Sixty-seven percent of operated and 71% of non-operated patients had abnormal contractions by left ventricular cineangiography. These preliminary results suggest that once the diagnosis of preinfarction angina pectoris is established and appropriate studies carried out, the patient's best interests are served by immediate aorto-coronary artery bypass surgery.

34. Surgical Treatment of Ventricular Irritability

E. D. MUNDTH, M. J. BUCKLEY, R. W. DeSANCTIS,*
W. M. DAGGETT and W. G. AUSTEN, Boston, Massachusetts

Myocardial revascularization and/or resection of a ventricular aneurysm appears to be an effective method of treating persistent and medically refractory ventricular irritability (VI) following acute myocardial infarction (AMI). Nine patients with medically refractory VI, varying from 2 days to 6 weeks post-AMI, have undergone cine coronary arteriography and left ventricular angiography and surgical treatment. Five of the nine patients were less than 2 weeks post-AMI. They were all hemodynamically unstable and 3 were in cardiogenic shock. All 5 had institution of intra-aortic balloon pump assistance (IABPA) with hemodynamic improvement but had persistent VI despite antiarrhythmic drug therapy and appropriate electrical pacing. The other four patients demonstrated intractable VI 2 to 6 weeks post-AMI. IABPA was used in one patient in this group to facilitate management during diagnostic study and induction of anesthesia as well as postoperatively. Seven of nine patients had a demonstrable left ventricular

*By invitation

aneurysm. Aneurysmectomy was carried out in 7 patients and was combined with 1, 2 or 3 vein bypass grafts in 3. Revascularization alone was carried out in 2 patients. Six of nine patients have survived. VI was improved and readily manageable in all 6 survivors. In 4 of the 6 survivors postoperative VI was managed with minimal antiarrhythmic therapy. Left ventricular function has been good or excellent in all 6 survivors.

35. Serial Angiographic Evaluation of Aortocoronary Vein Grafts in Sixty Consecutive Patients, Two Weeks, One Year, and Three Years after Operation

CLAUDE M. GRONDIN,* JEAN-PAUL MARTINEAU,*
CLAUDE MEERE* YVES R. CASTONGUAY,* GILLES LEPAGE*
and PIERRE R. GRONDIN, Montreal, Quebec, Canada

Although early results in aortocoronary vein grafts (ACVG) have been promising, critical appraisal must await long term studies.

Serial angiographic studies were conducted in 60 consecutive patients who underwent ACVG at the Montreal Heart Institute. All patients were studied two weeks, one year, and three years after operation. Occlusion occurred in 7 of the 91 grafts on the initial study and in 8 additional grafts on the second study. On this second study, most grafts displayed diffuse reduction in caliber—average reduction: 30 percent. In 3 grafts, reduction in caliber was greater than 75 percent. Two years later, all 3 grafts were occluded.

Except for these 3 grafts and 2 additional grafts which became occluded, there was no further attrition rate or reduction in caliber—segmental or diffuse—after one year in the remaining 71 grafts. The overall patency rate was therefore 92.3 percent after two weeks, 82.5 percent after one year, and 77 percent after three years. Hence the occlusion rate of ACVG after one year was only 5.5 percent.

These results indicate that the initial enthusiasm for ACVG was not unwarranted.

36. Coronary Bypass Grafting in 376 Consecutively Operated Patients with Three Operative Mortalities

JOHN E. HUTCHINSON, III,* GEORGE E. GREEN,
HAROUTUNE A. MEKHJIAN* and HARVEY G. KEMP,* New York,
New York

In the twenty-one month period from January, 1971, to September, 1972, 376 patients had coronary bypass grafts performed for coronary atherosclerosis. Three hundred and seventy-three patients were discharged from the hospital (alive) and three patients died. The operative mortality in this consecutively operated group was 0.8%. Four late deaths have occurred to date, two of them (being) due to pulmonary emboli, one due to hepatitis, and the fourth to myocardial infarction.

In this series, single grafts were performed in 60 patients, double grafts in 185, triple grafts in 121 patients, and quadruple grafts in ten patients. Included in this consecutive series are twenty-four patients with threatened infarction syndromes and twenty patients with diffuse scarring of the left ventricle.

*By invitation

We feel that the major factors accounting for this low mortality are (1) Increased experience in the performance of small vessel anastomosis. (2) Total avoidance of endarterectomy. (3) The performance of the distal coronary anastomosis with ventricular fibrillation rather than anoxic cardiac arrest. (4) The use of the internal mammary arteries and small veins from the lower legs as the bypass conduits of choice.

The specific details of these factors will be discussed.

WEDNESDAY AFTERNOON, APRIL 18, 1973

2:00 P.M. Scientific Session
Regency Ballroom

37. Pneumothorax Complicating Continuous Ventilatory Support

MICHAEL STEIER,* NATHANIEL CHING,* ENRIQUE BONFILS ROBERTS* and THOMAS F. NEALON JR.,
New York, New York

There has been a 35 per cent increase in the incidence of pneumothorax in our hospital in the last 3 years. The increase has been due to iatrogenic causes related to improvements in the care of critically ill patients—external cardiac massage, central venous pressure monitoring and continuous ventilatory support. The factors associated with continuous ventilatory support seemed appropriate for presentation before the Association.

In the period from January 1, 1968 to December 31, 1971, 61 patients at our institution developed pneumothorax during continuous ventilatory support. The significant factors leading to the occurrence of this complication include:

1. volume-controlled ventilation
2. large tidal volumes
3. high inspiratory pressures
4. positive end expiratory pressure
5. history of chronic obstructive lung disease and previous rib fractures.

These factors will be discussed in detail.

The importance of early diagnosis based on physical findings allowing immediate treatment will be stressed. Eight per cent of patients so diagnosed and treated succumbed as compared to 38 per cent of those in whom diagnosis waited confirmation with a chest x-ray due to the hazard of pressure ventilation in patients with pneumothorax.

38. Electronmicroscopy and Physical Chemistry of Healing in Prosthetic Heart Valves, Skirts, and Struts: Modification by Electrochemically Clean, Physiologic Surfaces

P. N. SAWYER, B. STANCZEWSKI,* N. RAMASAMY,*
G. W. KAMMLOTT,* J. G. STEMPAK* and S. SRINIVASAN,*
Brooklyn, New York

Repeated attempts have been made to decrease the incidence of valve thrombosis, infection, embolism, fibroblastic overgrowth, with orifice occlusion and/or poppet sticking. Sequential studies from this laboratory have revealed various solutions to these problems. The first problem, valve thrombosis, can be prevented by use of "electrochemically clean" oxide free negatively charged non-thrombogenic metallic surfaces. Aluminum, Starr-Edwards stellite 21, titanium and a recently developed metal surface from Howmedica seem to provide this antithrombogenic characteristic along with appropriate uniform net negative surface potentials.

*By invitation

Light electron and scanning electron microscopic studies of the surfaces and composition of healing characteristics of 100 valves implanted in the mitral and tricuspid annuli of calves for periods as long as 765 days have been completed. The scanning electron microscope and transmission electron microscope pictures have proven of real value in developing insight into the pathologic processes found in dysfunctioning valves.

The use of cloth around struts leads to abnormal fibrin deposition in the cloth interstices covering the metallic valve surfaces with invasive fibroblasts followed by abnormal collagen production, sequential platelet deposition and significant repetitive onion layering of fibrin on strut surfaces. This results in increasing strut diameter, poppet sticking, and increased trans-valvular pressure gradients.

Infection, fibroblastic overgrowth on the cuff, and orifice occlusion can be prevented by use of the dacron reinforced autogenous venous skirt which heals as normal autogenous tissue to the valve orifice. The remaining problems are obviated by the use of clean surfaces and appropriately designed valves as shown by flow, electrochemical, and valve surface studies before and following implantation. The characteristic photographs and surface phenomena changes found using the new techniques will be presented along with related interfacial potentials and characteristic light, electron, and scanning electron microscopic histology.

39. Evaluation of Aortic Valve Homograft Failures

ROBERT B. WALLACE, STEPHEN P. LONDE* and
JACK L. TITUS,* Rochester, Minnesota

Between May 1965 and June 1972, 229 patients (163 males and 66 females) had replacement of their aortic valve with an aortic valve homograft. The hospital mortality rate was 4.8% (11 patients). Of the 218 patients dismissed from the hospital, 22 have required reoperation for homograft failure 3 to 73 months after the initial operation, and 16 others have died during the follow-up period.

Valves removed at reoperation or autopsy were studied grossly and morphologically, and factors possibly related to the status of the homograft were evaluated. These factors included valve preparation (92 of the valves used in this series were sterilized with β -propiolactone and 137 were sterilized by irradiation), age and sex of the donor, condition of graft and recipient valve at time of insertion, function of graft at the time of hospital dismissal, and the relationship of graft function to the cause of death. Correlations were done in an attempt to define better those patients in whom the operation might be most appropriate and to delineate those factors that may be important in the long-term results of this procedure.

40. Surgical Management of Acquired Tricuspid Valve Disease

A. CARPENTIER,* A. DELOCHE,* A. HANANIA,* A. PIWNICA,*
Cl. FARGE* and Ch. DUBOST,* Paris, France
Sponsored by Dwight C. McGoon

Acquired tricuspid valve disease (A.T.V.D.) raises two questions which continue to challenge the surgeon. (1) In what circumstances the tricuspid disease

*By invitation

should be corrected? (2) and if so, by what means? A four years experience with the Carpentier tricuspid annuloplasty (C.A.) allows us to bring new answers to these questions.

From May 1967 through September 1972, 375 A.T.V.D. have been treated at Broussais Hospital with correction of severe mitral (271) or mitroaortic (104) diseases.

Kay annuloplasty (K.A) was used in 103 cases with an hospital mortality of 39% (1967-1969). C.A. has been used in 137 cases with a mortality of 9.5% (1969-1972). Starr prosthesis has been used in 135 cases with a mortality of 37% (1967-1972).

In the first 62 C.A., follow up study (1 to 4 years) including cardiac catheterization in 30 patients has shown constant effectiveness of the repair, no recurrent insufficiency, no A.V. block, no thromboembolic complications at the site of the tricuspid valve.

These results led us to a shift in our policy concerning the treatment of A.T.V.D.:

1. All A.T.V.D. clinically detected and operatively confirmed have to be corrected.

2. The C.A. is the only method capable to physiologically correct the disease and prevent the recurrent dilatation of the annulus. It can be used either isolated or combined with a commissurotomy in 90% of the A.T.V.D., as shown by our experience.

41. Closed Mitral Commissurotomy on Cannulated Pump Standby Through A Sternal Split Incision—An Alternative to Routine Open Commissurotomy

VINCENT L. GOTT, ANTONIO REVILLA,* JAMES S. DONAHOO,* EDWARD H. KLOPP,* and ROBERT K. BRAWLEY,*
Baltimore, Maryland

Open mitral commissurotomy offers several advantages over closed commissurotomy, including greater patient safety on pump bypass and easier conversion to prosthetic replacement if necessary. In some patients, however, fibrous obliteration of the commissures makes direct incision somewhat imprecise and the surgeon, in turn, may use a transventricular dilator to accomplish the commissurotomy. Engineering analysis demonstrates that a dilator used this way in the flaccid heart applies equal stress to the leaflets and commissures and not infrequently produces leaflet fracture, whereas in the beating heart, the stresses appear to be concentrated at the commissures because of the teathering mechanism of the chordae. All patients having mitral commissurotomy at this hospital since 1965 have been reviewed. One hundred patients had a standard closed commissurotomy (93% successful commissurotomies, 2% mortality) and 25 patients had open commissurotomy (40% requiring prosthesis, 4% mortality). In an attempt to combine the mechanical advantages of closed commissurotomy with the best features of open commissurotomy, we have more recently carried out all mitral commissurotomies as a "closed" procedure through a median sternotomy on cannulated pump standby with a newly designed "right angle" transventricular dilator. Eighteen patients have had this new procedure. One patient had a valve unsuited for commissurotomy and was easily converted to open cardiotomy for prosthetic replacement. The remaining 17 patients obtained

*By invitation

excellent commissurotomies with no morbidity or mortality. This technique for mitral commissurotomy appears to offer several advantages over other currently used methods.

42. The Open Approach to Mitral Commissurotomy

JAMES O. FINNEGAN,* HORACE MacVAUGH, III, DENNIS C.
GRAY,* CLAUDE R. JOYNER* and JULIAN JOHNSON,
Philadelphia, Pennsylvania

Of the 1005 operations on the mitral valve (380 open) done at the Hospital of the University of Pennsylvania from 1961 to 1971, 592 (317 open) consisted of mitral commissurotomy only.

As the first half of the 10 year period progressed the closed technic was used less and less, being reserved for the more favorable valves, but still constituted 50% for the period. For that 5 year period the mortality was 4% closed and 11% open—combined 7.5%. During the second 5 year period, the open technic was used exclusively, with a single death—less than 1% mortality.

The improved results for mitral commissurotomy in the second 5 year period was no doubt due to better selection of patients for the procedure, the freer replacement of the badly diseased valves, and improvements in postoperative care. These will be described, along with pre and postoperative complications.

The operative and late (5.3%) deaths in the open group have been analyzed. Of the surviving patients all but one have been followed and 73% have been restored to Class I (NYHA).

43. Mitral Valve Replacement with Beall Mitral Valve Prosthesis

N. P. ROSSI, C. KONGTAHWORN* and J. L. EHRENHAFT,
Iowa City, Iowa

To evaluate the function of the Beall valve, 100 consecutive cases of single valve replacements in the mitral position with a Beall prosthesis were analyzed. The procedures were performed from November 1967 to January 1972 allowing a followup of at least one year and up to five years. Eighty percent were performed for rheumatic disease and 20% for mitral incompetence due to nonrheumatic causes. Sixty-five percent of patients were in New York Heart Association Class IV, 26% in Class III and 9% in Class II. We found a 2% incidence of thromboembolism, 2% of paravalvular leak, and 19% of hemolysis. The problems relating to hemolysis were encountered early and were resolved within six months. Twenty-four percent had a previous mitral commissurotomy. Mortality was confined to patients in functional Classes III and IV (17% hospital and 7% late). Of 65 patients who were in functional Class IV, 59 showed improved functional classification, 25 out of 26 patients in functional Class III had changed to Class I or II postoperatively, and six out of nine patients in functional Class II had improved to Class I.

The results of analysis suggested satisfactory clinical improvement after replacement with Beall valve mitral prosthesis. There was a low incidence of thromboembolism but a high initial incidence of hemolysis.

*By invitation

44. Mitral Valve Replacement with Cloth-Covered, Composite Seat Prostheses: The Case for Early Operation

LAWRENCE I. BONCHEK* and ALBERT STARR, Portland,
Oregon

Operative and late complications of prosthetic valves have usually limited mitral valve replacement (MVR) to functional class (FC) III or IV patients refractory to medical management.

146 patients have undergone MVR with cloth-covered, composite seat prostheses (models 6310, 6320). 129 were FC III or IV. Operative mortality was 2% (3/146), and late mortality was 9% (13/146). Only two late deaths were valve related (one leak, one infection). Nine patients had emboli (6%) two with significant residuals, in 2,216 months of patient followup.

110 patients were FC I or II postoperatively. Since this improvement did not correlate with preoperative FC, a new *prognostic* classification was introduced to correlate preoperative *duration* of symptoms and *response* to medical therapy with postoperative result. 75% (15/20) survivors with unsatisfactory functional results postoperatively were in the worst prognostic classification (C2).

The striking reduction in *valve related complications*, and the minimal operative mortality, indicate that MVR with the current prosthesis should be offered early to patients with recent deterioration who respond to medical treatment (prognostic class A1). The poor functional results seen after MVR for neglected mitral disease may thus be avoided.

THE AMERICAN ASSOCIATION FOR THORACIC SURGERY

FUTURE MEETINGS

1974	April 22-24	Las Vegas Hilton Las Vegas, Nevada	•••
1975	April 14-16	Americana Hotel New York, New York	

*By invitation

The American Association for Thoracic Surgery, 1972-1973

Honorary members

- ALLISON, PHILIP Radcliffe Infirmary, Oxford, England
- BARRETT, NORMAN R. Old Palace Place, Richmond Green, Surrey, England
- BELSEY, RONALD Frenchay Hospital, Bristol, England
- BJÖRK, VIKING O. Karolinska Institute, Stockholm, Sweden
- BOEREMA, I. Surgical Clinic, University of Amsterdam, Netherlands
- BROCK, RUSSELL C. Guy's Hospital, London, England
- BROM, A. GERARD University Hospital, Leiden, Holland
- CRAFOORD, CLARENCE Sabbatsberg Sjukhus, Stockholm, Sweden
- d'ABREU, A. L. Queen Elizabeth Hospital, Edgbaston, Birmingham, England
- ELOESSER, LEO APTO Post 39, Tacámbaro, Michoacan, Mexico
- HOLMAN, EMILE 722 Funston Ave., San Francisco, Calif. 94118
- LOGAN, ANDREW Royal Infirmary, Edinburgh 3, Scotland
- OCHSNER, ALTON 1516 Jefferson Highway, New Orleans, La. 70121
- THOMAS, CLEMENT PRICE Court Green, St. Ann's Hills, Midhurst, Sussex, England
- THOMPSON, VERNON Vicarage House, Lloves, Hereford, England
- ZERBINI, E. J. Rua Itapeva, 500-6° Andar São Paulo, Brazil

Active members

- ADAM, MAURICE 3434 Swiss Ave., Dallas, Texas 75204
- ADKINS, PAUL C. 2150 Pennsylvania Ave., N.W., Washington, D.C. 20037
- ADLER, RICHARD H. 100 High St., Buffalo, N.Y. 14203
- ALLEN, PETER 2966 West 45th Ave., Vancouver, B.C., Canada
- ALLEY, RALPH D. Albany Medical Center Hospital, Albany, N.Y. 12208
- ALMOND, CARL H. University of Missouri Medical Center, Columbia, Mo. 65201
- ANDERSEN, MURRAY N. 462 Grider St., Buffalo, N.Y. 14215
- ANDREWS, NEIL C. School of Medicine, University of California, Davis, Calif. 95616
- ANKENEY, JAY L. 2065 Adelbert Rd., Cleveland, Ohio 44106
- ARONSTAM, ELMORE M. 5159 Crown Ave., La Canada, Calif. 91011

If any correction should be made in the spelling of names or in the addresses of members, please notify the Executive Secretary of the Association, Mr. William T. Maloney, Six Beacon Street, Suite 620, Boston, Mass. 02108.

- ASHBAUGH, DAVID G. 1670 Ada St., Boise, Idaho 83702
- ASHMORE, PHILLIP G. 750 West Broadway, Vancouver 9, B.C., Canada
- ATTAR, SAFUHH M. A. University Hospital, Baltimore, Md. 21201
- AUSTEN, W. GERALD Massachusetts General Hospital, Boston, Mass. 02114
- BAFFES, THOMAS G. 4055 Main St., Skokie, Ill. 60076
- BAHNSON, HENRY T. Presbyterian-University Hospital, Pittsburgh, Pa. 15213
- BAIRD, RONALD J. 72 Clarendon Ave., Toronto 7, Ontario, Canada
- BARKER, WALTER L. 2912 N. Commonwealth Ave., Apt. 11C, Chicago, Ill. 60657
- BARNER, HENDRICK B. 1325 South Grand Blvd., St. Louis, Mo. 63104
- BARRETT, RAYMOND J. 28870 Inkster Rd., Southfield, Mich. 48076
- BARSAMIAN, ERNEST M. 1400 Veterans of Foreign Wars Parkway, Boston, Mass. 02132
- BARTLEY, THOMAS D. 2226 N.W. Pettygrove, Portland, Ore. 97210
- BARWINSKY, JAROSLAW 205 Medical Arts Building, 404 Graham Ave., Winnipeg 1, Canada
- BAUE, ARTHUR E. 216 S. Kingshipway, St. Louis, Mo. 63110
- BEALL, ARTHUR C., JR. 1200 M. D. Anderson Blvd., Houston, Texas 77025
- BEATTIE, EDWARD J., JR. 444 East 68th St., New York, N.Y. 10021
- BELL, JOHN W. Forks Mediclinic, P.O. Box 89, Forks, Wash. 98331
- BENDER, HARVEY W., JR. Vanderbilt University Medical Center, Nashville, Tenn. 37203
- BENFIELD, JOHN R. 1000 W. Carson St., Torrance, Calif. 90509
- BENOIT, HECTOR W., JR. 503 Plaza Parkway Bldg., Kansas City, Mo. 64112
- BERT, RALPH, JR. 508 West Sixth Ave., Suite 504, Spokane, Wash. 99204
- BERGER, ROBERT L. 750 Harrison Ave., Boston, Mass. 02118
- BERGMANN, MARTIN 8515 Delmar Blvd., St. Louis, Mo. 63124
- BERNATZ, PHILIP E. Mayo Clinic, Rochester, Minn. 55902
- BERNHARD, WILLIAM F. 300 Longwood Ave., Boston, Mass. 02115
- BESKIN, CHARLES A. 3929 Convention St., Baton Rouge, La. 70806
- BIGELOW, WILFRED G. Toronto General Hospital, Toronto 2, Ontario, Canada
- BLACK, HARRISON 319 Longwood Ave., Boston, Mass. 02115
- BLAIR, EMIL 4200 East Ninth Ave., Denver, Colo. 80220
- BLAKE, HUAL 7765 Devonshire Dr., Knoxville, Tenn. 37919
- BLAKEMORE, WILLIAMS 19th and Lombard St., Philadelphia, Pa. 19146
- BLALOCK, JOHN B. 1516 Jefferson Highway, New Orleans, La. 70121
- BLOODWELL, ROBERT D. 1021 E. Robinson St., Orlando, Fla. 32801
- BLOOMER, WILLIAM E. 3440 Atlantic Ave., Long Beach, Calif. 90807
- BLUMENSTOCK, DAVID A. Mary Imogene Bassett Hospital, Cooperstown, N.Y. 13326
- BLUNDELL, PETER E. Montreal General Hospital, Montreal, Quebec, Canada
- BOSHER, LEWIS H. 1200 E. Broad St., Richmond, Va. 23219
- BOUGAS, JAMES A. 110 Francis St., Boston, Mass. 02115
- BOWMAN, FREDERICK O., JR. 161 Fort Washington Ave., New York, N.Y. 10032
- BOYD, ARTHUR D. Department of Surgery, New York University Medical Center, 550 First Ave., New York, N.Y. 10016
- BOYD, DAVID P. 605 Commonwealth Ave., Boston, Mass. 02215
- BOYD, THOMAS F. 452 Pleasant St., Malden, Mass. 02148
- BRADHAM, R. RANDOLPH Ashley House, Suite 2J,

- BRAUNWALD, NINA S. Department of Surgery, Peter Bent Brigham Hospital, 721 Huntington Ave., Boston, Mass. 02115
- BRINDLEY, G. VALTER, JR. Scott and White Clinic, Temple, Texas 76501
- BROCKMAN, STANLEY K. 29th St. and Ellis Ave., Chicago, Ill. 60616
- BROOKS, JAMES W. 1200 E. Broad St., Richmond, Va. 23219
- BROWN, IVAN W., JR. 1600 Lakeland Hills Blvd., Lakeland, Fla. 33802
- BROWN, LEE B. 926 E. McDowell Rd., Phoenix, Ariz. 85006
- BRUNEAU, JACQUES 3875 St. Urbain, Suite 307, Montreal 131, Quebec, Canada
- BRYANT, LESTER R. University of Kentucky Medical Center, Lexington, Ky. 40506
- BUCKLEY, MORTIMER J. Massachusetts General Hospital, Boston, Mass. 02114
- BURDETTE, WALTER J. M. D. Anderson Hospital and Tumor Institute, Houston, Texas 77025
- BURKE, JOHN F. Massachusetts General Hospital, Boston, Mass. 02114
- CALLAGHAN, JOHN C. Suite 550, 8409 112 St., Edmonton, Alberta, Canada
- CAMISHION, RUDOLPH C. Suite 303, Cooper Parkway, West North Park Dr., Pennsauken, N.J. 08109
- CAMPBELL, DANIEL C., JR. 2665 Clevelanland Ave., Suite 107, Fort Myers, Fla. 33901
- CAMPBELL, GILBERT S. 4301 W. Markham St., Little Rock, Ark. 72201
- CANTRELL, JAMES R. 325 Ninth Ave., Seattle, Wash. 98104
- CAREY, JOSEPH S. 2200 Santa Monica Blvd., Santa Monica, Calif. 90404
- CARTER, MAX G. 670 George St., New Haven, Conn. 06511
- CARTER, PAUL R. 1421 Alpine Dr., West Covina, Calif. 91791
- CASTANEDA, ALDO R. University of Minnesota Hospitals, Minneapolis, Minn. 55414
- CENTER, SOL 637 DuPont Bldg., Miami, Fla. 33131
- CHESNEY, JOHN G. 7450 Red Rd., S. Miami, Fla. 33136
- CLARK, RICHARDE. 4960 Audubon Ave., St. Louis, Mo. 63110
- CLAUSS, ROY H. 1249 Fifth Ave., New York, N.Y. 10029
- CLEVELAND, RICHARD J. Harbor General Hospital, 1000 W. Carson Street, Torrance, Calif. 90509
- CLOWES, GEORGE H. A., JR. 818 Harrison Ave., Boston, Mass. 02118
- COHEN, MORLEY 295 Dromore Ave., Winnipeg, Manitoba, Canada
- COLE, FRANCIS H. 176 South Bellevue, Memphis, Tenn. 38106
- COLLINS, HAROLD A. Department of Surgery, Vanderbilt University Hospital, Nashville, Tenn. 37203
- COLLINS, JOHN J., JR. Peter Bent Brigham Hospital, Boston, Mass. 02115
- CONNAR, RICHARD G. One David Blvd., Tampa, Fla. 33606
- CONNOLLY, JOHN E. University of California at Irvine, Irvine, Calif. 92664
- CONRAD, PETER W. 2304 Westmoreland St., Falls Church, Va. 22046
- COOK, WILLIAM A. 1300 Morris Park Ave., New York, N.Y. 10461
- COOLEY, DENTON A. 6621 Fannin St., Houston, Texas 77025
- CORDELL, A. ROBERT Bowman Gray School of Medicine, Winston-Salem, N.C. 27103
- COUVES, CECIL M. 11-101 Clinical Sciences Bldg., University of Alberta, Edmonton, Alberta, Canada
- COWLEY, R. ADAMS University Hospital, Baltimore, Md. 21201
- CRAWFORD, E. STANLEY 1200 Moursund Ave., Houston, Texas 77025
- CROSS, FREDERICK S. 11311 Shaker Blvd., Cleveland, Ohio 44104
- CULINER, MORRIS M. 2233 Post St., San Francisco, Calif. 94115

- DAGGETT, WILLARD M. Massachusetts General Hospital, Department of Surgery, Fruit Street, Boston, Mass. 02114
- DAICOFF, GEORGE R. University of Florida College of Medicine, Gainesville, Fla. 32603
- DALE, W. ANDREW 2000 Church St., Nashville, Tenn. 37203
- DALTON, MARTIN L., JR. 3801 21st St. Lubbock, Texas 79410
- DAMMANN, JOHN F. University of Virginia Medical Center, Charlottesville, Va. 22901
- DANIELSON, GORDON K., JR. Mayo Clinic, Rochester, Minn. 55901
- DAUGHTRY, DeWITT C. 1550 N.W. 10th Ave., Miami, Fla. 33136
- DAVILA, JULIO C. Henry Ford Hospital, 2799 W. Grand Blvd., Detroit, Mich. 48202
- DAVIS, MILTON V. 3434 Swiss Ave., Suite 405, Dallas, Texas 75204
- DEATON, W. RALPH, JR. 1027 Professional Village, Greensboro, N.C. 27401
- DEMOS, NICHOLAS J. 100 Clifton Place, Jersey City, N.J. 07304
- DeMUTH, WILLIAM E., JR. Milton S. Hershey Medical Center, Hershey, Pa. 17033
- DeNIORD, RICHARD N. 1911 Thomson Dr., Lynchburg, Va. 24501
- DERRICK, JOHN R. University of Texas Medical Branch, Galveston, Texas 77551
- DESFORGES, GERARD 452 Pleasant St., Malden, Mass. 02148
- DETERLING, RALPH A., JR. 171 Harrison Ave., Boston, Mass. 02111
- DeWALL, RICHARD A. 247 Northview Rd., Dayton, Ohio 45419
- DeWEESE, JAMES A. 260 Crittenden Blvd., Rochester, N.Y. 14642
- DILLARD, DAVID H. 12712 39th N.E., Seattle, Wash. 98155
- DILLON, MARCUS L., JR. Department of Surgery, Veterans Administration Hospital, Lexington, Ky. 40507
- DIVELEY, WALTER L. 121 Twenty-First Ave., North, Nashville, Tenn. 37203
- DOBELL, ANTHONY R. C. 2300 Tupper St., Montreal 108 Quebec, Canada
- DOMM, SHELDON E. 1918 W. Clinch Ave., Knoxville, Tenn. 37916
- DOOLEY, BYRON N. Oak Hills Medical Center, Pasteur Drive, San Antonio, Texas 78229
- DRAKE, EMERSON H. 19 Bramhall St., Portland, Me. 04102
- DRAPANAS, THEODORE 1430 Tulane Ave., New Orleans, La. 70112
- EASTRIDGE, CHARLES E. Veterans Administration Hospital, Memphis, Tenn. 38115
- EBERT, PAUL A. The New York Hospital, Rm. F-739, New York, N.Y. 10021
- ECKER, ROGER R. 5323 Harry Hines Blvd., Dallas, Texas 75235
- EDMUNDS, L. HENRY JR. University of California Medical Center, San Francisco, Calif. 94112
- EDWARDS W. STERLING University of New Mexico School of Medicine, Albuquerque, N.M. 87106
- EFFLER, DONALD B. Euclid and East 93rd St., Cleveland, Ohio 44106
- EHRENHAFT, JOHANN L. University Hospitals, Iowa City, Iowa 52240
- EISEMAN, BEN University of Colorado Medical Center, 4200 E. Ninth Ave., Denver, Colo. 80220
- ELLIS, F. HENRY, JR. 605 Commonwealth Ave., Boston, Mass. 02115
- ELLISON, ROBERT G. Medical College of Georgia, Augusta, Ga. 30902
- EMERSON, GEORGE L. 11 Rochester St., Scottsville, N.Y. 14546
- FABER, L. PENFIELD 1753 W. Congress Parkway, Chicago, Ill. 60612
- FAVALORO, RENE G. Buenos Aires, Argentina, Clinica Guemes, Cordoba 3933

- FERGUSON, THOMAS B. 4989 Barnes Hospital Plaza,
St. Louis, Mo. 63110
- FINDLAY, CHARLES W., JR. 161 Fort Washington Ave.,
New York, N.Y. 10032
- FINEBERG, CHARLES 829 Spruce St., Philadelphia, Pa. 19107
- FISHMAN, NOEL H. Department of Surgery, Rm. 488-m, University
of California Medical Center, San Francisco, Calif. 94122
- FITZPATRICK, HUGH F. St. Luke's Hospital, New York, N.Y. 10025
- FLEMMMA, ROBERT J. 9800 W. Bluemound Rd., Milwaukee, Wis. 53226
- FLYNN, PIERCE J. 399 E. Highland Ave., Suite 506,
San Bernardino, Calif. 92410
- FONKALSRUD, ERIC W. UCLA Medical Center, Los Angeles, Calif. 90024
- FORD, JOSEPH M. 1056 Fifth Ave., New York, N.Y. 10028
- FORD, WILLIAM B. 412 N. Whitfield St., Pittsburgh, Pa. 15213
- FOSBURG, RICHARD G. Box 255, Naval Hospital, San Diego, Calif. 92134
- FOSTER, JOHN H. Vanderbilt University Hospital, Nashville, Tenn. 37203
- FOX, ROBERT T. 2136 Robin Crest Lane, Glenview, Ill. 60025
- FRANK, HOWARD A. 330 Brookline Ave., Boston, Mass. 02215
- FRATER, ROBERT W. M. 17 Gladwin Pl., Bronxville, N.Y. 10708
- FRENCH, SANFORD W., III 307 East Buena Vista, Barstow, Calif. 92311
- FROBESE, ALFRED S. 1425 Scrope Rd., Rydal, Pa. 19046
- GAENSLER, EDWARD A. 229 Dudley Rd., Newton Center, Mass. 02159
- GAGNON, EDOUARD D. 30 Est. Blvd. St. Joseph, Suite 1002, Montreal,
Quebec, Canada
- GARAMELLA, JOSEPH J. 1629 Medical Arts Bldg.,
Minneapolis, Minn. 55402
- GARDNER, RICHARD E. 414 Geary Medical Bldg., Suite 412,
San Francisco, Calif. 94118
- GARRETT, H. EDWARD 910 Madison Ave., Suite 823,
Memphis, Tenn. 38103
- GARZON, ANTONIO A. 450 Clarkson Ave., Brooklyn, N.Y. 11203
- GERARD, FRANKLYN P. 144 S. Harrison St., Suite 809,
E. Orange, N.J. 07018
- GERST, PAUL H. 622 W. 168th St., New York, N.Y. 10032
- GIANNELLI, STANLEY, JR. 153 W. 11th St., New York, N.Y. 10011
- GILBERT, JOSEPH W., JR. Emory University Branch Post Office, P. O. Box
15063, Atlanta, Ga. 30333
- GLASS, BERTRAM A. 3600 Prytania St., New Orleans, La. 70115
- GLENN, WILLIAM W. L. 333 Cedar St., New Haven, Conn. 06510
- GOBBEL, WALTER G., JR. Veterans Administration Hospital,
Nashville, Tenn. 37203
- GOLDMAN, BERNARD S. 223 Strathallen Wood, Toronto,
Ontario, Canada 305
- GONZALEZ, LUIS L. Eden and Bethesda Avenues, Cincinnati, Ohio 45219
- GOTT, VINCENT L. Johns Hopkins Hospital, Baltimore, Md. 21205
- GRAVEL, JOFFRE-ANDRE 170 Grande-Allee West, Quebec 6,
Quebec, Canada
- GREEN, GEORGE E. St. Luke's Hospital, Amsterdam Ave. at 114th St.,
New York, N.Y. 10025
- GREENBERG, JACK J. 4300 Alton Rd., Miami Beach, Fla. 33140
- GREENFIELD, LAZAR J. 921 N.E. 13th St., Oklahoma City, Okla. 73104
- GREER, ALLEN E. 1211 N. Shartel, Oklahoma City, Okla. 73103
- GRILLO, HERMES C. Massachusetts General Hospital, Boston, Mass. 02114
- GRIMES, ORVILLE F. University of California Hospital,
San Francisco, Calif. 94122

- GRONDIN, PIERRE 5000 E. Belanger, Montreal 410, Quebec, Canada
- GROVES, LAURENCE K. Cleveland Clinic, Cleveland, Ohio 44106
- GWATHMEY, OWEN 5700 Old Richmond Ave., Richmond, Va. 23226
- HAIRSTON, PETER 80 Barre Street, Medical College Hospital,
Charleston, S.C. 29401
- HALL, DAVID P. 1000 E. Third St., Chattanooga, Tenn. 37403
- HALLER, J. ALEX, JR. Johns Hopkins Hospital, Baltimore, Md. 21205
- HALLMAN, GRADY L., JR. 6621 Fannin St., Houston, Texas 77025
- HANLON, C. ROLLINS 55 E. Erie St., Chicago, Ill. 60611
- HARDY, JAMES D. University of Mississippi Medical Center,
Jackson, Miss. 39216
- HARDY, KENNETH L. 3115 Webster St., Oakland, Calif. 94609
- HARRISON, ALBERT W. 3155 Stagg Drive, Beaumont, Texas 77701
- HARRISON, ROBERT W. 1810 Wealthy St., S.E., Grand Rapids,
Mich. 49605
- HATCHER, CHARLES R., JR. Emory University Clinic,
Atlanta, Ga. 30322
- HAUPT, GEORGE J. 306 Lankenau Medical Bldg., Philadelphia, Pa. 19151
- HEANEY, JOHN P. Medical Professional Bldg., San Antonio, Texas 78212
- HEIMBECKER, RAYMOND O. Toronto General Hospital, Toronto 2,
Ontario, Canada
- HERRERA, RODOLFO 6a. Avenida 8-71, Zone 10, Guatemala City,
Guatemala
- HEWITT, ROBERT LEE 1430 Tulane Avenue, New Orleans, La. 70112
- HEWLETT, THOMAS H. 17821 South Pioneer Blvd., Artesia, Calif. 90701
- HIGGINSON, JOHN F. 2320 Bath St., Suite 213,
Santa Barbara, Calif. 93105
- HILL, LUCIUS D. 1118 Ninth Ave., Seattle, Wash. 98101
- HIROSE, TEURO 5830 Tyndall Ave., Bronx, N.Y. 10471
- HOLDER, THOMAS M. 510 Medical Plaza Bldg., 4320 Wornall Rd.,
Kansas City, Mo. 64111
- HOLLAND, ROBERT H. 3216 Beverly Drive, Dallas, Texas 75205
- HOLSWADE, GEORGE R. 517 E. 71st St., New York, N.Y. 10021
- HOOD, R. MAURICE 3100 Red River, Austin, Texas 78705
- HOOD, RICHARD H., JR. Drawer E, Cotulla, Texas 78014
- HOPEMAN, ALAN R. Associate Professor of Surgery, Section of Thoracic
Surgery, N311 University Medical Center, Columbia, Mo. 65201
- HOWARD, HECTOR S., JR. 910 Madison Ave., Memphis, Tenn. 38103
- HUDSPETH, ALLEN S. Bowman Gray School of Medicine,
Winston-Salem, N.C. 27103
- HUFNAGEL, CHARLES A. 3800 Reservoir Rd., N.W.,
Washington, D.C. 20007
- HUGHES, RICHARD K. 2293 Dallin St., Salt Lake City, Utah 84109
- HUMPHREY, EDWARD W. Veterans Administration Hospital,
Minneapolis, Minn. 55417
- HUNTER, JAMES A. 1725 W. Harrison St., Chicago, Ill. 60612
- HURLEY, EDWARD J. University of California, Davis, Calif. 95616
- HUTCHIN, PETER 214 Avenida Cortez, La Jolla, Calif. 92037
- JAHNKE, EDWARD J., JR. 1596 San Leandro Lane,
Santa Barbara, Calif. 93103
- JAMPLIS, ROBERT W. Palo Alto Clinic, Palo Alto, Calif. 94301
- JAVID, HUSHANG 1725 W. Harrison St., Chicago, Ill. 60612
- JENSIK, ROBERT J. 1725 W. Harrison St., Chicago, Ill. 60612

- JOHNS, THOMAS N. P. 6305 Towana Rd., Richmond, Va. 23226
 JOHNSON, FRANK E. 1012 Metropolitan Arts Bldg., 825 So. 8th Street,
 Minneapolis, Minn. 55404
 JOHNSON, W. DUDLEY 9800 W. Bluemound Rd., Milwaukee, Wis. 53226
 JOHNSTON, FRANK R. Bowman Gray School of Medicine,
 Winston-Salem, N.C. 27103
 JONES, THOMAS W. 715 Minor Ave., Seattle, Wash. 98104
 JOYNT, GEORGE H. C. 25 Leonard Ave., Suite 102, Toronto 2b,
 Ontario, Canada
 JUDE, JAMES R. 220 N.E. 17th St., Miami, Fla. 33132
 JULIAN, ORMAND C. 1716 Sandpiper, Palm Desert, Calif. 92260
 KAHN, DONALD R. 1300 University Ave., Madison, Wisc. 53706
 KAISER, GEORGE C. 1325 S. Grand Blvd., St. Louis, Mo. 63104
 KAISER, GERARD A. Department of Surgery, University of Miami School
 of Medicine, P.O. Box 875, Biscayne Annex, Miami, Fla. 33152
 KARLSON, KARL E. 110 Lockwood St., Providence, R.I. 02903
 KAUSEL, HARVEY W. Albany Medical Center Hospital,
 Albany, N.Y. 12208
 KAY, JEROME HAROLD 123 S. Alvarado St., Los Angeles, Calif. 90057
 KEE, JOHN L., JR. 3707 Gaston Ave., Dallas, Texas 75246
 KEMLER, R. LEONARD 21 Woodland St., Hartford, Conn. 06105
 KENNEDY, JOHN H. Texas Medical Center, Houston, Texas 77025
 KERTH, WILLIAM J. Pacific Medical Center, San Francisco, Calif. 94115
 KESHISHIAN, JOHN M. 2520 L St., N.W., Washington, D.C. 20037
 KEY, JAMES A. Toronto General Hospital, Toronto, Ontario, Canada
 KILMAN, JAMES W. 410 West Tenth Ave., Columbus, Ohio 43210
 KILLEN, DUNCAN A. 510 Medical Plaza Bldg., 4320 Warnall Rd.,
 Kansas City, Mo. 64111
 KING, HAROLD 1100 West Michigan St., Indianapolis, Ind. 46207
 KING, ROBERT D. 1100 West Michigan St., Indianapolis, Ind. 46207
 KIRKLIN, JOHN W. University of Alabama Medical Center,
 Birmingham, Ala. 35233
 KIRSCHNER, PAUL A. 2 East 92nd St., New York, N.Y. 10028
 KITTLE, C. FREDERICK 5454 S. Shore, Chicago, Ill. 60615
 KOVARIK, JOSEPH L. 1633 Fillmore St., Denver, Colo. 80206
 LAFORET, EUGENE G. 2000 Washington St., Newton Lower Falls,
 Mass. 02162
 LANGSTON, HIRAM T. 952 Pine Tree Lane, Winnetka, Ill. 60093
 LAWRENCE, G. HUGH 1118 Ninth Ave., Seattle, Wash. 98101
 LAWRENCE, MONTAGUE S. University Hospitals, Iowa City, Iowa 52240
 LEE, WILLIAM H., JR. 80 Barre St., Charleston, S.C. 29401
 LEES, WILLIAM M. 6518 N. Nokomis Ave., Lincolnwood, Ill. 60646
 LEFEMINE, ARMANDA A. 60 Highland Circle, Wayland, Mass. 01778
 LEININGER, BERNARD J. 55 E. Washington St., Chicago, Ill. 60602
 LEMMON, WILLIAM M. 220 North 15th St., Philadelphia, Pa. 19102
 LEPAGE, GILLES 445 Lockhart Ave., Montreal 16, Quebec, Canada
 LEPLEY, DERWARD, JR. 9800 W. Bluemound Rd., Milwaukee,
 Wis. 53226
 LEVITSKY, SIDNEY University of Illinois College of Medicine,
 P.O. Box 6998, Chicago, Ill. 60680
 LIDDLE, HAROLD V. 535 East First South, Salt Lake City, Utah 84102
 LILLEHEI, RICHARD C. University Hospitals, P.O. 490,
 Minneapolis, Minn. 55455
 LINDESMITH, GEORGE G. 1136 W. 6th St., Los Angeles, Calif. 90017

- LINDSEY, EDWARD S. 1430 Tulane Ave., New Orleans, La. 70112
 LITTLEFIELD, JAMES B. Memorial University Faculty of Medicine,
 St. John's, Newfoundland, Canada
 LITWAK, ROBERT S. 5th Ave. at 100th St., New York, N.Y. 10029
 LOGAN, WILLIAM D., JR. 272 Boulevard, N.E., Atlanta, Ga. 30312
 LONG, DAVID M., JR. University of Illinois College of Medicine,
 P.O. Box 6988, Chicago, Ill. 60680
 LOWER, RICHARD R. 1200 East Broad St., Richmond, Va. 23219
 LUCIDO, JOSEPH L. 634 N. Grand Blvd., St. Louis, Mo. 63103
 LYNN, R. BEVERLEY R. R. No. 1, Westbrook, Ontario, Canada
 MacKENZIE, JAMES W. Rutgers Medical School, CMDNJ, P.O. Box 2100,
 New Brunswick, N.J. 08903
 MACKLER, S. ALLEN 104 South Michigan Ave., Chicago, Ill. 60603
 MacLEAN, LLOYD D. Royal Victoria Hospital, Montreal 2,
 Quebec, Canada
 MADOFF, IRVING M. 1180 Beacon St., Brookline, Mass. 02146
 MAGOVERN, GEORGE J. Suite 265—One Allegheny Sq., Allegheny Center,
 Pittsburgh, Pa. 15212
 MALETTE, WILLIAM G. University of Kentucky Medical Center,
 Lexington, Ky. 40506
 MALM, JAMES R. 161 Fort Washington Ave., New York, N.Y. 10032
 MALONEY, JAMES V., JR. UCLA Medical Center, Los Angeles,
 Calif. 90024
 MANDELBAUM, ISIDORE 1100 West Michigan St., Indianapolis,
 Ind. 46207
 MANNIX, EDGAR P., JR. 2850 Sixth Ave., San Diego, Calif. 92103
 MARK, JAMES B. D. Department of Surgery, Stanford University School
 of Medicine, Stanford, Calif. 94305
 MATLOFF, JACK M. Cedars-Sinai Medical Center, P.O. Box 54265,
 Los Angeles, Calif. 90054
 MAY, IVAN A. 3115 Webster St., Oakland, Calif. 94609
 MAYER, JOHN H., JR. 503 Plaza Parkway Bldg., Kansas City, Mo. 64112
 McBURNEY, ROBERT P. Suite 524, 910 Madison Ave., Memphis,
 Tenn. 38103
 McCLENATHAN, JAMES E. The Children's Hospital, Washington,
 D.C. 20009
 McGOON, DWIGHT C. Mayo Clinic, Rochester, Minn. 55902
 McLAUGHLIN, JOSEPH S. 22 South Greene St., Baltimore, Md. 21201
 McNAMARA, JOSEPH JUDSON 396 Dune Circle, Kailua, Hawaii 96734
 McVAUGH, HORACE 1. S. Ravdin Institute, 3400 Spruce Street,
 Philadelphia, Pa. 19104
 MECKSTROTH, CHARLES V. Ohio State University Hospital,
 Columbus, Ohio 43210
 MEREDITH, JESSE H. Bowman Gray School of Medicine, Winston-Salem,
 N.C. 27103
 MEYER, BERT W. 1136 West Sixth St., Los Angeles, Calif. 90017
 MILLER, DON R. University of Kansas Medical Center,
 Kansas City, Kans. 66103
 MILLER, FLETCHER A. Creighton-St. Joseph Hospital,
 Omaha, Nebr. 68102
 MILLER, GEORGE E. 214 Sixth Ave., West, Calgary, Alberta, Canada
 MILLS, WALDO O. 1001 Broadway, Suite 216, Seattle, Wash. 98122
 MINOR, GEORGE R. B. P. 50 Tunis, Belvedere, Tunisia
 MITCHEL, BEN F., JR. 3434 Swiss Ave., Suite 404, Dallas, Texas 75204
 MOORE, THOMAS C. 1000 W. Carson St., Torrance, Calif. 90509

- SCHRAMMEL, ROBERT J. 4440 Magnolia St., New Orleans, La. 70115
 SCHUSTER, SAMUEL R. 300 Longwood Ave., Boston, Mass. 02115
 SCHWARTZ, SEYMOUR I. 260 Crittenden Blvd., Rochester, N.Y. 14620
 SCOTT, HENRY J. 3350 Cote des Neiges, Suite 540, Montreal 25,
 Quebec, Canada
 SCOTT, STEWART M. One Northwood Road, Asheville, N.C. 28803
 SEALY, WILL C. Duke University Hospital, Durham, N.C. 27706
 SENNING, ÅKE Surgical University Clinic A, Kantonsspital, 8006
 Zurich, Switzerland
 SEYBOLD, WILLIAM D. 6624 Fannin St., Houston, Texas 77025
 SHIELDS, THOMAS W. 333 E. Huron St., Chicago, Ill. 60611
 SHUMWAY, NORMAN E. Stanford Medical Center, Palo Alto, Calif. 94302
 SILVER, DONALD Duke University Medical Center, Durham, N.C. 27706
 SIRAK, HOWARD D. The Children's Hospital, 17th St. at Livingston
 Park, Columbus, Ohio 43203
 SKINNER, DAVID B. 601 N. Broadway, Baltimore, Md. 21205
 SLOAN, HERBERT University Hospital, Ann Arbor, Mich. 48104
 SMOLEFF, EDWARD A. 5301 F St., Sacramento, Calif. 95819
 SMYTH, NICHOLAS P. D. 110 Irving St., N.W., Washington, D.C. 20010
 SOROFF, HARRY S. 818 Harrison Ave., Boston, Mass. 02118
 SPEAR, HAROLD C. 909 Interama Blvd., N. Miami Beach, Fla. 33162
 SPENCER, FRANK C. 550 First Ave., New York, N.Y. 10016
 STANFORD, WILLIAM Wilford Hall, USAF Medical Center (SGHST),
 Lackland Air Force Base, Texas 78236
 STANSEL, HORACE C., JR. 333 Cedar St., New Haven, Conn. 06510
 STARKEY, GEORGE W. B. 110 Francis St., Boston, Mass. 02215
 STARR, ALBERT 3181 S.W. Sam Jackson Park Rd., Portland, Ore. 97201
 STATE, DAVID Albert Einstein College of Medicine, New York, N.Y. 10061
 STEICHEN, FELICIEN M. 3459 Fifth Ave., Pittsburgh, Pa. 15213
 STEMMER, EDWARD A. Veterans Administration Hospital, Long Beach,
 Calif. 90801
 STEPHENSON, SAM E., JR. 2000 Jefferson St., Jacksonville, Fla. 32209
 STERN, HAROLD 2 Church St., So., New Haven, Conn. 06519
 STERNS, LAURENCE P. 8409 112 St., Edmonton, Alberta, Canada
 STILES, QUENTIN R. 1136 West 6th Street, Los Angeles, Calif. 90017
 STRANAHAN, ALLAN Albany Medical Center Hospital,
 Albany, N.Y. 12208
 SUGG, WINFRED L. 5323 Harry Hines Blvd., Dallas, Texas 75235
 SULLIVAN, HERBERT J. Medical Arts Bldg., Hamilton, Ontario, Canada
 SYMBAS, PANAGIOTIS N. 69 Butler St., S.E., Atlanta, Ga. 30303
 TABER, RODMAN E. 500 Saddle Lane, Grosse Pointe Woods, Mich. 48236
 TAKARO, TIMOTHY Veterans Administration Hospital, Oteen, N.C. 28801
 TAYLOR, FREDERICK H. 1012 Kings Dr., Charlotte, N.C. 28207
 TAYLOR, WARREN J. 452 Pleasant St., Malden, Mass. 02148
 TEMPLETON, JOHN Y., III 829 Spruce St., Philadelphia, Pa. 19107
 THAL, ALAN P. University of Kansas Medical Center,
 Kansas City, Kans. 66103
 THOMAS, GEORGE I. 715 Minor Ave., Seattle, Wash. 98104
 THOMAS, GORDON W. Int. Grenfell Association, St. Anthony,
 Newfoundland, Canada
 THOMAS, PAUL A., JR. Lancaster and City Line Aves.,
 Philadelphia, Pa. 19151
 THOMSON, NORMAN B., JR. St. Francis Hosp., Port Washington Blvd.
 Roslyn, N.Y. 11576
 TICE, DAVID A. 550 First Ave., New York, N.Y. 10016
 TIMMES, JOSEPH J. New Jersey College of Medicine,
 Jersey City, N.J. 07304

- TIMMIS, HILARY H. William Beaumont Hospital, 3601 West 13 Mile Rd.,
 Royal Oak, Mich. 48072
 TOCKER, ALFRED M. Medcenter Bldg., 932 North Topeka,
 Wichita, Kans. 67214
 TRIMBLE, ALAN S. Toronto General Hospital, Toronto 2, Ontario, Canada
 TRINKLE, J. KENT Department of Surgery, University of Texas Medical
 School at San Antonio, 7703 Floyd Curl Dr., San Antonio, Texas 78229
 TRUMMER, MAX J. Mercy Hosp. Med. Ctr. 4077 Fifth Ave.
 San Diego, Calif. 92103
 TRUSLER, GEORGE A. 123 Edward St., Suite 1225, Toronto, 101,
 Ontario, Canada
 TSUJI, HAROLD K. 2220 Lynn Road, Suite 204, Thousand Oaks,
 Calif. 91360
 URSCHER, HAROLD C., JR. 3810 Swiss Ave., Dallas Texas 75204
 VASKO, JOHN S. 410 W. 10th Avenue, Columbus, Ohio 43210
 VEITH, FRANK J. 111 East 210th St., New York, N.Y. 10467
 WADDELL, WILLIAM R. 4200 East Ninth Ave., Denver, Colo. 80220
 WALDHAUSEN, JOHN A. Milton S. Hershey Medical Center,
 Hershey, Pa. 17033
 WALKER, GEORGE R. 289 Cedar St., P.O. Box 970, Sudsbury,
 Ontario, Canada
 WALKER, JAMES H. 1323 Quarrier St., East, Charleston, W. Va. 25301
 WALLACE, ROBERT B. 200 1st St., S.W., Rochester, Minn. 55901
 WARDEN, HERBERT E. West Virginia University Medical Center,
 Morgantown, W. Va. 26506
 WATKINS, ELTON, JR. 605 Commonwealth Ave., Boston, Mass. 02215
 WEBB, WATTS R. 750 E. Adams St., Syracuse, N.Y. 13210
 WEINBERG, MILTON, JR. 1725 W. Harrison St., Suite 448,
 Chicago, Ill. 60612
 WEISEL, WILSON 2266 North Prospect Ave., Milwaukee, Wis. 53202
 WELDON, CLARENCE S. 4960 Audubon Ave., St. Louis, Mo. 63110
 WESOLOWSKI, SIGMUND A. Mercy Hospital, Rockville Centre,
 N.Y. 11570
 WHEAT, MYRON W., JR. Department of Surgery, Health Sciences Center,
 P.O. Box 1055, Louisville, Ky. 40201
 WICHERN, WALTER A., JR. 428 West 59th St., New York, N.Y. 10019
 WILCOX, BENSON R. University of North Carolina School of Medicine,
 Chapel Hill, N.C. 27514
 WILDER, ROBERT J. 200 W. Cold Spring Lane, Baltimore, Md. 21210
 WILKINS, EARLE W., JR. Massachusetts General Hospital,
 Boston, Mass. 02114
 WILLIAMS, G. RAINEY 800 Northeast 13th St., P.O. Box 26901,
 Oklahoma City, Okla. 73190
 WILLMAN, V. L. 1325 South Grand Blvd., St. Louis, Mo. 63104
 WILSON, HUGH E., III 6011 Harry Hines Blvd., Dallas, Texas 75235
 WILSON, JOHN L. Stanford Medical Center, Palo Alto, Calif. 94304
 WILSON, ROBERT F. Department of Surgery, W.S.U. School of
 Medicine, 540 E. Canfield St., Detroit, Mich. 48201
 WITMER, ROBERT H. 126 East Chestnut St., Lancaster, Pa. 17602
 WOLCOTT, MARK W. 500 Foothill Blvd., Salt Lake City, Utah 84113
 WOLFF, WILLIAM I. 10 Nathan D. Perlman Place, New York, N.Y. 10003
 YEH, THOMAS J. Memorial Medical Center, P.O. Box 6688, Station C,
 Savannah, Ga. 31405

YOUNG, W. GLENN, JR. Box 3396, Duke University Medical Center,
Durham, N.C. 27706
YOUNG, WILLIAM P. 1300 University Ave., Madison, Wis. 53706
ZAROFF, LAWRENCE I. 80 Pelham Rd., Rochester, N.Y. 14610

Associate members

ADAMS, JESSE E., JR. 1000 E. 3rd St., Chattanooga, Tenn. 37403
ADELMAN, ARTHUR . . . 601 E. 63rd St., Suite 503, Kansas City, Mo. 64110

BAISCH, BRUCE F. 644 E. Regent St., Inglewood, Calif. 90301
BRYANT, J. RAY 1169 Eastern Parkway, Louisville, Ky. 40217
BURBANK, BENJAMIN 244 Henry St., Brooklyn, N.Y. 11201

CAHAN, WILLIAM G. 444 East 68th St., New York, N.Y. 10021
CHANDLER, JOHN H. 616 West Forest Ave., Jackson, Tenn. 38301
CHODOFF, RICHARD J. 1905 Spruce St., Philadelphia, Pa. 19103
COOKE, FRANCIS N. 25 S.E. Second Ave., Miami, Fla. 33131
CRACOVANER, ARTHUR J. 103 East 78th St., New York, N.Y. 10021
CRASTNOPOL, PHILIP 8 N. Circle Dr., Great Neck, L.I., N.Y. 11020
CRECCA, ANTHONY D. 376 Roseville Ave., Newark, N.J. 07107
CRUTCHER, RICHARD R. 2101 Nicholasville Rd., Lexington, Ky. 40503

De BORD, ROBERT A. 414 St. Mark Court, Peoria, Ill. 61603
DECKER, ALFRED M., JR. 8 Church St., Saranac Lake, N.Y. 12983
DeMATTEIS, ALBERT . . . 2700-62 Avenue South, St. Petersburg, Fla. 33711

FELTON, WARREN L., II 5700 N.W. Grand Blvd., Oklahoma City,
Okla. 73103

FINNERTY, JAMES . . . Brookhaven Medical Arts Bldg., Patchogue, N.Y. 11772
FRIEDLANDER, RALPH Grand Concourse and Mt. Eden Parkway,
Bronx, N.Y. 10457

FRIESEN, STANLEY R. University of Kansas Medical Center,
Kansas City, Kans. 66103

FULLER, JOSIAH 205 West 2nd St., Duluth, Minn. 55802

GENTSCH, THOMAS O. 1150 N.W. 14th St., Miami, Fla. 33136
GERBASI, FRANCIS S. 81 Lochmoor Blvd., Grosse Pointe Shores,
Mich. 48236

HAUSMANN, PAUL F. 2309 West State St., Milwaukee, Wis. 53233
HEAD, LOUIS R. 55 East Washington St., Chicago, Ill. 60602
HENLY, WALTER S. 900 Medical Center Professional Bldg.,
Houston, Texas 77025

HERING, ALEXANDER C., Capt., MC, USN USS Sanctuary (AH 17)
c/o F.P.O., San Francisco, Ca. 96601

HERR, RODNEY H. 123 East Idaho St., Boise, Idaho 83702

INGRAM, IVAN N. 1439 Brixton Rd., Pasadena, Calif. 91105
IOVINE, VINCENT M. 2520 L St., N.W., Washington, D.C. 20037

JARETZKI, ALFRED, III 161 Fort Washington Ave.,
New York, N.Y. 10032

JENSEN, NATHAN K. 1629 Medical Arts Bldg.,
Minneapolis, Minn. 55402
JOHNSON, CLIVE R. 1519 Pennsylvania, Fort Worth, Texas 76104
JUDD, ARCHIBALD R. 304 N. Fourth St., Hamburg, Pa. 19526

KAUNITZ, VICTOR H. 3878 Delaware Ave., Tonawanda, N.Y. 14223
KRAEFT, NELSON H. 143 Miccosukee Rd., Tallahassee, Fla. 32303
KUNDERMAN, PHILIP J. . . . 185 Livingston Ave., New Brunswick, N.J. 08902
KUNSTLER, WALTER E. 1538 Sherbrooke St., West, Montreal 109,
Quebec, Canada

LASLEY, CHARLES H. 1200 South Druid Rd., Clearwater, Fla. 33516
LEIBOVITZ, MARTIN 300 Utica Square Medical Center,
Tulsa, Okla. 74114

LEVOWITZ, BERNARD S. 555 Prospect Place, Brooklyn, N.Y. 11238
LEWIS, J. Eugene, Jr. 1401 So. Grand, St. Louis, Mo. 63104
LINBERG, EUGENE J. Suite 14 Landmark Bldg., 328 5th Ave. S.,
Naples, Fla. 33940

LUI, ALFRED H. F. Wayne County General Hospital, Eloise, Mich. 48132

MAHAFFEY, DANIEL E. . . . 366 Medical Towers South, Louisville, Ky. 40202
MAIN, F. BEACHLEY Allegheny General Hospital, Pittsburgh, Pa. 15212
MANGIARDI, JOSEPH L. 222 Front St., Mineola, N.Y. 11501
McCORD, COLIN W. 421 W. 113th St., New York, N.Y. 10025
McKEOWN, JOHN J., JR. . . . 935 Cedar Grove Road, Wynnewood, Pa. 19096
McPHAIL, JASPER L. 214 Baptist Medical Arts Bldg.,
Little Rock, Ark. 72202

MENDELSSOHN, EDWIN 1351 West Tabor Rd., Philadelphia, Pa. 19141
MILLER, ARTHUR C. 453 W. Freeway, Roseburg, Ore. 97470
MILLER, CAROL C. 304 Humphrey St., Swampscott, Mass. 01907
MILLER, DONALD B. Mary Fletcher Hospital, Burlington, Vt. 05401

NARODICK, BENJAMIN G. 2040 W. Wisconsin Ave.,
Milwaukee, Wis. 53233

NEERKEN, ADRIAN J. 404 Bronson Medical Center,
Kalamazoo, Mich. 49004

NETTERVILLE, RUSH E. 514 E. Woodrow Wilson Drive,
Jackson, Miss. 39216

NEWMAN, ROBERT W. University of Tennessee, Physicians Office
Building, 1928 Alcoa Highway, Suite 318, Knoxville, Tenn. 37920

OCHSNER, ALTON, JR. 3525 N. Causeway Blvd., Suite 601,
Metairie, La. 70002

O'NEILL, JAMES F. 1425 Woodland Rd., Rydal, Pa. 19046
OVERSTREET, JOHN WM. 2210 Maroneal St., Houston, Texas 77025

PEMBERTON, ALBERT H. 2040 West Wisconsin Ave.,
Milwaukee, Wis. 53203

PERRY, JOHN F., JR. # 2 Red Fox Rd., North Oaks,
St. Paul, Minn. 55110

PIERUCCI, LOUIS, JR. Suite 300, Cooper Parkway West, North Park
Drive, Pennsauken, N.J. 08109

PINKHAM, ROLAND D. . . . 801 Broadway, Suite 901, Seattle, Wash. 98122
PRATT, LAWRENCE A. 219 N. Austin St., Paris, Ill. 61944

- RAMS, JAMES J. The Western Pennsylvania Hospital,
4800 Friendship Ave., Pittsburgh, Pa. 15224
- ROBBINS, S. GWIN 20 South Dudley St., Memphis, Tenn. 38103
- ROSS, RALEIGH R. 2 Medical Arts Square, Austin, Texas 78705
- RYAN, BERNARD J. 375 East Main St., Bay Shore, N.Y. 11706
- RYAN, THOMAS C. Veterans Administration Hospital, Garner's
Ferry Road, Columbia, S.C. 29201
- SANES, GILMORE M. 301 Buckingham Rd., Pittsburgh, Pa. 15215
- SELMAN, MORRIS W. 2302 Meadowwood Drive, Toledo, Ohio 43602
- SHERMAN, PAUL H. 2425 Lake Sue Drive, Orlando, Fla. 32803
- SIDERY, HARRY 1815 N. Capitol Ave., Indianapolis, Ind. 46202
- SILVER, ARTHUR W. 612 W. Duarte Rd., Suite 603,
Arcadia, Calif. 91006
- SNYDER, HOWARD E. 103½ E. Ninth Ave., Winfield, Kans. 67156
- STAYMAN, JOSEPH W. 8815 Germantown Ave., Philadelphia, Pa. 19118
- STENSTROM, JOHN D. 220-1105 Pandora Ave., Victoria,
British Columbia, Canada
- THROWER, WENDELL B. Furnace Brook Medical Bldg., Suite 32,
1261 Furnace Brook Parkway, Quincy, Mass. 02169
- TILLOU, DONALD J. Hillcrest Road, R.D. 1, Elmira, N.Y. 14903
- TRICERRI, FERNANDO E. La Ferratiere, 1297 Founex, Switzerland
- VAN FLEIT, WILLIAM E. 401 Jefferson Medical Arts Bldg.,
South Bend, Ind. 46617
- WATKINS, DAVID H. 6039 North Waterbury Rd.,
Des Moines, Iowa 50312
- ZUHDI, M. NAZIH 1121 North Shartel, Oklahoma City,
Okla. 73103

Senior members

- ABBOTT, OSLER Emory University Clinic, Atlanta, Ga. 30322
- ADAMS, HERBERT D. 605 Commonwealth Ave., Boston, Mass. 02215
- ADAMS, WILLIAM E. 55 East Erie St., Chicago, Ill. 60611
- ALLBRITTEN, FRANK F., JR. University of Kansas Medical Center,
Kansas City, Kan. 66103
- AMBERSON, J. B. 16 Sherwood Drive, Hillsdale, N.J. 07642
- AUERBACH, OSCAR Veterans Administration Hospital,
East Orange, N.J. 07019
- AUFSES, ARTHUR H. 301 E. 66th St., New York, N.Y. 10021
- BADGER, THEODORE L. 264 Beacon St., Boston, Mass. 02116
- BAILEY, CHARLES P. 34 East 67th St., New York, N.Y. 10021
- BARKLEY, HOWARD T. 609 Hermann Professional Bldg.,
Houston, Texas 77025
- BARONOFESKY, IVAN D. 550 Washington Street, Suite 611,
San Diego, Calif. 92103
- BATTERSBY, JAMES S. 1040 W. Michigan St., Indianapolis, Ind. 46202
- BEECHER, HENRY K. 10 Shattuck St., Boston, Mass. 02115
- BENEDICT, EDWARD B. 24 Essex Road, Chestnut Hill, Mass. 02167
- BENSON, CLIFFORD D. 1515 David Whitney Bldg., Detroit, Mich. 48226

- BERRY, FRANK B. 169 East 69th St., New York, N.Y. 10021
- BETTS, REEVE H. Veterans Administration Hospital,
Oteen, N.C. 28805
- BISGARD, J. DEWEY 422 Doctors Bldg., Omaha, Neb. 68131
- BLADES, BRIAN Veterans Administration Hospital, Suite 112C,
50 Irving St., N.W., Washington, D.C. 20422
- BLOCH, ROBERT G. Montefiore Hospital, New York, N.Y. 10067
- BLOOMBERG, ALLAN E. 100 E. Gunhill Rd., Bronx, N.Y. 10028
- BRANTIGAN, OTTO C. 104 West Madison St., Baltimore, Md. 21201
- BREWER, LYMAN A., III 658 S. Bonnie Brae St., Los Angeles,
Calif. 90057
- BROWN, ROBERT K. 1727 Gilpin St., Denver, Colo. 80218
- BROWNRIGG, GARRETT M. 47 Queens Rd., St. Johns, Newfoundland
- BUCKINGHAM, WILLIAM W. 310 W. 49th St., Apt. 405,
Kansas City, Mo. 64112
- BUGDEN, WALTER F. 1200 E. Genesee St., Syracuse, N.Y. 13210
- BURFORD, THOMAS H. 4989 Barnes Hospital Plaza,
St. Louis, Mo. 63110
- BURNETT, W. EMORY 47 E. Righters Mill Rd., Narberth, Pa. 19072
- CARLSON, HERBERT A. 21 Seventh Pl., Long Beach, Calif. 90802
- CARLSON, ROBERT I. N. 38th Pl., Phoenix, Ariz. 85012
- CARR, DUANE 20 S. Dudley St., Memphis, Tenn. 38103
- CARTER, B. NOLAN 5005 Willow Hills Lane, Cincinnati, Ohio 45243
- CHAMBERS, JOHN S., JR. 2850 Sixth Ave., San Diego, Calif. 92103
- CLAGETT, O. THERON Mayo Clinic, Rochester, Minn. 55902
- CLATWORTHY, H. WILLIAM, JR. 695 Bryden Road, Columbus,
Ohio 43205
- CLERF, LOUIS H. 5575 8th Ave., North, St. Petersburg, Fla. 33702
- COHEN, ROY B. Stanford Hospital, Palo Alto, Calif. 94302
- COLE, DEAN B. Professional Bldg., Richmond, Va. 23219
- CONDON, WILLIAM B. 2045 Franklin St., Denver, Colo. 80218
- CONKLIN, WILLIAM S. 511 Southwest Tenth ave., Portland, Ore. 97205
- COTTON, BERT H. 111 Congress St., Pasadena, Calif. 91105
- COURNAND, ANDRE 630 W. 168th St., New York, N.Y. 10032
- CRANDELL, WALTER B. Veterans Administration Hospital,
White River Junction, Vt. 05001
- CURRERI, ANTHONY R. 1300 University Ave., Madison, Wis. 53706
- CUTLER, PRESTON R. 535 E. 1st South St., Salt Lake City,
Utah 84102
- DAILEY, JAMES E. 3214 Reba Drive, Houston, Texas 77019
- DANIEL, ROLLIN A. 2000 Hayes St., Nashville, Tenn. 37203
- DANIELS, ALBERT C. Box 6535 Carmel, Calif. 93921
- DAVIDSON, LOUIS R. 1025 Fifth Ave., New York, N.Y. 10028
- DAY, J. CLAUDE 3790 Woodward Ave., Detroit Mich. 48201
- De BAKEY, MICHAEL E. 1200 Moursund Ave., Houston, Texas 77025
- DeCAMP, PAUL T. 1514 Jefferson Highway, New Orleans, La. 70121
- DELARUE, NORMAN C. 25 Donlea Drive, Toronto 17,
Ontario, Canada
- DENNIS, CLARENCE 8208 Hamilton Spring Ct.,
Bethesda, Md. 20034
- DODRILL, FOREST DEWEY 641 David Whitney Bldg.,
Detroit, Mich. 48226
- DORNER, RALPH A. 710 Equitable Bldg., Des Moines, Iowa 50309
- DORSEY, JOHN M. 2650 Ridge Ave., Evanston, Ill. 60201

- DOUGLASS, RICHMOND 32 Vassar View Rd., Poughkeepsie, N.Y. 12603
- DOVELL, CHAUNCEY E. (Address unknown)
- DRASH, EVERETT C. University of Virginia Hospital, Charlottesville, Va. 22901
- DUGAN, DAVID J. 3115 Webster Street, Oakland, Calif. 94609
- EVANS, BYRON H. 3291 N. Hilliard Lane, Fresno, Calif. 93703
- FALOR, WILLIAM H. 208 Medical Arts Bldg., Akron, Ohio 44304
- FAULKNER, WILLIAM B., JR. 20 San Rafael Way, San Francisco, Calif. 94127
- FELL, EGBERT H. P.O. Box 227, Kealahakua, Kona, Hawaii
- FISCHER, WALTER W. 170 E. 78th St., New York, N.Y. 10021
- FLICK, JOHN B. 819 Black Rock Rd., Gladwyne, Pa. 19035
- GEARY, PAUL 1117 Waterway Lane, Delray Beach, Fla. 33444
- GEBAUER, PAUL Leahi Hospital, Honolulu, Hawaii 96816
- GERBODE, FRANK Pacific Medical Center, San Francisco, Calif. 94115
- GIBBON JOHN H., JR. 2103 N. Providence Rd., Media, Pa. 19063
- GLENN, FRANK 525 E. 68th St., New York, N.Y. 10021
- GOLDMAN, ALFRED c/o P.O. Box 2025, Palm Springs, Calif. 92262
- GROSS, ROBERT E. 300 Longwood Ave., Boston, Mass. 02115
- GROW, JOHN B. 2045 Franklin St., Room 910, Denver, Colo. 80205
- HARKEN, DWIGHT E. 67 Bay State Road, Boston, Mass. 02215
- HARPER, FREDERICK R. 2045 Franklin St., Denver, Colo. 80218
- HARRINGTON, STUART W. Mayo Clinic, Rochester, Minn. 55902
- HARRISON, ELLIOTT 1260 W. 38th St., Vancouver 13, B.C., Canada
- HART, DERYL Duke University Medical Center, Durham, N.C. 27706
- HARTER, JOHN S. 1169 Eastern Parkway, Louisville, Ky. 40217
- HEAD, JEROME R. 55 E. Washington St., Chicago, Ill. 60602
- HELMSWORTH, JAMES A. Cincinnati General Hospital, Cincinnati, Ohio 45229
- HEROY, WILLIAM W. 29 E. Gate Road, Huntington, N.Y. 11743
- HOLINGER, PAUL H. 700 N. Michigan Ave., Chicago, Ill. 60611
- HOLMAN, CRANSTON W. 862 Fifth Ave., New York, N.Y. 10021
- HOPKINS, WILLIAM A. 1293 Peachtree St., N.E., Atlanta, Ga. 30309
- HUDSON, THEODORE R. 251 E Chicago Ave., Chicago, Ill. 60611
- HUDSON, W. A. Hudsonakers, Jasper, Ark. 72641
- HUGHES, FELIX A., JR. 263 N. Rose Rd., Memphis, Tenn. 38117
- HUMPHREYS, GEORGE H., II 180 Fort Washington Ave., New York, N.Y. 10032
- HURWITZ, ALFRED 4300 Alton Rd., Miami Beach, Fla. 33140
- JARVIS, FRED J. 819 Boylston Ave., Seattle, Wash. 98104
- JOHNSON, ELGIE K. 1077 Northern Blvd. Roslyn, N.Y. 11576
- JOHNSON, HOLLIS E. 2122 W. End Ave., Nashville, Tenn. 37205
- JOHNSON, JULIAN 31 Righter's Mill Rd., Gladwyne, Pa. 19035
- JOHNSTON, J. HARVEY, JR. 710 N. State St., Jackson, Miss. 39201
- JONES, JOHN C. 1136 W. Sixth St., Los Angeles, Calif. 90017
- KAY, EARLE B. 2475 E. 22nd St., Cleveland, Ohio 44115
- KEELEY, JOHN L. P.O. Box 1336, Hines, Ill. 60141

- KELLEY, WINFIELD O. Uncas-on-Thames, Norwich, Conn. 06361
- KERGIN, FREDERICK G. 61 Ardwood Gate, Toronto 178, Ontario, Canada
- KESSLER, CHARLES R. 801 Princeton Ave., Suite 327, Birmingham, Ala. 35211
- KING, RICHARD 340 Boulevard, N.E., Atlanta, Ga.
- KLASSEN, KARL P. Ohio State University, Columbus, Ohio 43215
- KLEPSE, ROY G. 1835 Eye St., N.W., Washington, D.C. 20006
- KLOPSTOCK, ROBERT Veterans Administration Hospital, Brooklyn, N.Y. 11209
- KNOEPP, LOUIS F. Veterans Administration Hospital, Alexandria, La. 71301
- LAIRD, ROBERT R.R. # 2, Woodbridge, Ontario, Canada
- LAM, CONRAD R. Henry Ford Hospital, Detroit, Mich. 48202
- LAMBERT, ADRIAN 768 Park Ave., New York, N.Y. 10021
- LEAHY, LEON J. 176 Bryant St., Buffalo, N.Y. 14222
- LEEDS, SANFORD E. 2211 Post St., San Francisco, Calif. 94115
- LESTER, CHARLES W. 320 E. 72nd St., New York, N.Y. 10021
- LEVEN, N. LOGAN 1464 Lowry Medical Arts Bldg., St. Paul, Minn. 55102
- LEWIS F. JOHN Department of Surgery, Northwestern University Medical School, Chicago, Ill. 60611
- LILLEHEI, C. WALTON 435 E. 70th St., New York, N.Y. 10021
- LINDSKOG, GUSTAF E. 38 Griffing Ford Rd., Branford, Conn. 06405
- LOCKWOOD, A. L. 300 Bloor St., E., Toronto, Ontario, Canada
- LONGMIRE, WILLIAM P., JR. UCLA Medical Center, Los Angeles, Calif. 90024
- LYNCH, JOSEPH P 2000 Washington Street, Newton Lower Falls, Mass. 02162
- MacMANUS, JOSEPH E. 50 High St., Buffalo, N.Y. 14203
- MAHONEY, EARLE B. 260 Crittenden Blvd., Rochester, N.Y. 14620
- MAIER, HERBERT C. 3 E. 71st St., New York, N.Y. 10021
- MAURER, ELMER P. R. 250 Wm. Howard Taft Rd., Cincinnati, Ohio 45219
- MAUTZ, F. R. 13241 Ravenna Rd., Chardon, Ohio 44024
- McDONALD, JOHN R. Harper Hospital, Detroit, Mich. 48201
- McINTOSH, CLARENCE A. 900 Sherbrooke St., West, Montreal, Quebec, Canada
- MEADE, RICHARD H. 750 San Jose Drive, S.E., Grand Rapids, Mich. 49506
- MELICK, DERMONT W. University of Arizona College of Medicine, Tucson, Ariz. 85721
- MELTZER, HERBERT 14127-98th Ave., Edmonton, 51, Alberta, Canada
- MENDELSON, HARVEY J. 2065 Adelbert Rd., Cleveland, Ohio 44106
- MERENDINO, K. ALVIN University of Washington, School of Medicine, Seattle, Wash. 98105
- MERKEL, CARL G. 8 Church St., Saranac Lake, N.Y. 12983
- MEYER, HERBERT WILLY Box 507, Rancho Santa Fe, Calif. 92067
- MICHELSON, ELLIOTT 200 W. Cold Spring Lane, Baltimore, Md. 21210
- MISCALL, LAURENCE 11 E. 68th St., New York, N.Y. 10021
- MOERSCH, RICHARD N. 399 E. Highland Ave., Suite 506, San Bernardino, Calif. 92404
- MOORE, RICHMOND L. 3320 Woodridge Pl., Lynchburg, Va., 24503
- MYERS, J. ARTHUR 1316 Mayo Memorial Bldg., Minneapolis, Minn. 55455

- NIXON, JAMES W. 1121 Nix Professional Bldg., San Antonio,
Texas 78205
- OATWAY, WILLIAM H., JR. 146 Monarch Bay, Laguna Niguel,
Calif. 92677
- OLSEN, ARTHUR M. Mayo Clinic, Rochester, Minn. 55902
- OVERHOLT, RICHARD H. 135 Francis St., Boston, Mass. 02215
- PAPPER, EMANUEL M. P.O. Box 875, Biscayne Annex,
Miami, Fla. 33152
- PARKER, EDWARD F. 158 Rutledge Ave., Charleston, S.C. 29408
- PHILLIPS, FRANCIS J. 2023 Leussac Drive, Anchorage, Alaska 99503
- PICKHARDT, OTTO C. 25 E. 77th St., New York, N.Y. 10028
- POOL, JOHN L. The Norwalk Hospital, 24 Stephens St.,
Norwalk, Conn. 06856
- POPPE, J. KARL 2525 N.W. Lovejoy St., Portland, Ore. 97210
- PROCTOR, OSCAR S. 1101 Garraty Rd., San Antonio, Texas 78209
- RAVITCH, MARK M. 3459 Fifth Ave., Pittsburgh, Pa. 15213
- READ, C. THOMAS Lakeview Medical Arts Center, 13200 N. 103rd Ave.,
Sun City, Ariz. 85351
- RIENHOFF, WILLIAM F., JR., 911 Poplar Hill Rd., Baltimore, Md. 21210
- RIGGINS, H. McLEOD 1031 Fifth Ave., New York, N.Y. 10028
- RIGLER, LEO G. Los Angeles Center for Health Sciences,
Los Angeles, Calif. 90024
- RIPSTEIN, CHARLES B. 15 Birch St., Great Neck, L.I., N.Y. 11020
- ROBERTSON, ROSS 6067 Blink Bonnie Rd., W. Vancouver, B.C., Canada
- ROGERS, W. L. 414 Geary Medical Bldg., Suite 412,
San Francisco, Calif. 94118
- ROSEMOND, GEORGE P. 3401 N. Broad St., Philadelphia, Pa. 19140
- RUMEL, WILLIAM R. 535 E. 1st S. St., Salt Lake City, Utah 84102
- SALYER, JOHN M. 1125 E. 17th St., Santa Ana, Calif. 92701
- SAMSON, PAUL C. 15 LaSalle Ave., Piedmont, Calif. 94611
- SAROT, IRVING A. 197 E. 85th St., New York, N.Y. 10028
- SCOTT, HENRY W., JR. Vanderbilt University Hospital,
Nashville, Tenn. 37203
- SEILER, HAWLEY H. 517 Bayshore Blvd., Tampa, Fla. 33606
- SELEY, GABRIEL P. 799 Park Ave., New York, N.Y. 10021
- SHAW, ROBERT R. Nangrahar Medical School, Jalalabad, Afghanistan
- SHUMACKER, HARRIS B., JR. Indiana University Medical Center,
Indianapolis, Ind. 46207
- SIMEONE, FIORINDO A. 164 Summit Ave., Providence, R.I. 02906
- SKINNER, EDWARD F. 20 S. Dudley St., Memphis, Tenn. 38103
- SKINNER, GEORGE F. 36 Coburg St., St. John, New Brunswick,
Canada
- SMITH, DAVID T. Duke University Medical Center,
Durham, N.C. 27706
- SNYDER, JOHN M. 800 Ostrum St., Bethlehem, Pa. 18015
- SOMMER, GEORGE N. J., JR. 120 W. State St., Trenton, N.J. 08608
- SOUTTER, LAMAR 577 Bridge St., Dedham, Mass. 02026
- STEPHENS, H. BRODIE 1105 Greenwich St., San Francisco,
Calif. 94109
- STRIEDER, JOHN W. 2000 Washington St., Newton Lower Falls,
Mass. 02162
- STRODE, JOSEPH E. 888 So. King St., Honolulu, Hawaii 96813

- STRUG, LAWRENCE H. 2435 Octavia St., New Orleans, La. 70115
- SWAN, HENRY, II 6700 W. Lakeridge Rd., Denver, Colo. 80227
- SWENSON, ORVAR Children's Memorial Hospital,
Chicago, Ill. 60614
- THOMPSON, SAMUEL A. 850 Park Ave., New York, N.Y. 10021
- TOUROFF, ARTHUR S. W. 333 E. 79th St., New York, N.Y. 10021
- TYSON, M. DAWSON Hitchcock Clinic, Hanover, N.H. 03755
- VAN ALLEN, CHESTER M. State Hospital, Bikaner, Rajputana, India
- VARCO, RICHARD L. University of Minnesota Medical Center,
Minneapolis, Minn. 55455
- VINEBERG, ARTHUR M. 3170 de Lavigne Road, Montreal 218,
Quebec, Canada
- VORWALD, ARTHUR J. 1741 Deckner Ave., Green Bay, Wis. 54302
- WALKUP, HARRY E. R.F.D. 1, Worton, Md. 21678
- WANGENSTEEN, OWEN H. University of Minnesota Medical Center,
Minneapolis, Minn. 55414
- WATERMAN, DAVID H. 1918 W. Clinch Ave., Knoxville, Tenn. 37916
- WATSON, WILLIAM L. 340 E. 72nd St., New York, N.Y. 10021
- WEINBERG, JOSEPH A. 111 Marquez Place, Pacific Palisades,
Calif. 90272
- WILLAUER, GEORGE 6129 Greene St., Philadelphia, Pa. 19144
- WILLIAMS, MARK H. 63 Front St., Binghamton, N.Y. 13905
- WILSON, JULIUS L. 924 Canyon Rd., Santa Fe, N. Mex. 87501
- WILSON, NORMAN J. Parramore Hospital, Crown Point, Ind. 46307
- WIPER, THOMAS B. 40 Bayview Ave., Belvedere, Calif. 94920
- WOODS, FRANCIS M. 135 Francis St., Boston, Mass. 02115
- WRIGHT, GEORGE W. 11311 Shaker Blvd., Cleveland, Ohio 44104
- WYLIE, ROBERT H. 161 Fort Washington Ave., New York, N.Y. 10032

THE AMERICAN ASSOCIATION FOR THORACIC SURGERY

Charter Members

June 7, 1917

E. Wyllis Andrews	Arthur A. Law
John Auer	William Lerche
Edward R. Baldwin	Howard Lilienthal
Walter M. Boothby	William H. Luckett
William Branower	Morris Manges
Harlow Brooks	Walton Martin
Lawrason Brown	Rudolph Matas
Kenneth Bulkley	E. S. McSweeney
Alexis Carrel	Samuel J. Melter
Norman B. Carson	Willy Meyer (Founder)
J. Frank Corbett	James Alexander Miller
Armistead C. Crump	Robert T. Miller
Charles N. Dowd	Fred J. Murphy
Kennon Dunham	Leo S. Peterson
Edmond Melchior Eberts	Eugene H. Pool
Max Einhorn	Walther I. Rathbun
Herman Fischer	Martin Rehling
Albert H. Garvin	B. Merrill Ricketts
Nathan W. Green	Samuel Robinson
John R. Hartwell	Charles I. Scudder
George J. Heuer	William H. Stewart
Chevalier Jackson	Franz Torek
H. H. Janeway	Martin W. Ware
James H. Kenyon	Abraham O. Wilensky
Adrian V. S. Lambert	Sidney Yankauer

Meetings of the American Association for Thoracic Surgery

1918 - Chicago	President, Samuel J. Meltzer
1919 - Atlantic City	President, Willy Meyer
1920 - New Orleans	President, Willy Meyer
1921 - Boston	President, Rudolph Matas
1922 - Washington	President, Samuel Robinson
1923 - Chicago	President, Howard Lilienthal
1924 - Rochester, Minn.	President, Carl A. Hedblom
1925 - Washington	President, Nathan W. Green
1926 - Montreal	President, Edward W. Archibald
1927 - New York	President, Franz Torek
1928 - Washington	President, Evarts A. Graham
1929 - St. Louis	President, John L. Yates
1930 - Philadelphia	President, Wyman Whittemore
1931 - San Francisco	President, Ethan Flagg Butler
1932 - Ann Arbor	President, Frederick T. Lord
1933 - Washington	President, George P. Muller
1934 - Boston	President, George J. Heuer
1935 - New York	President, John Alexander
1936 - Rochester, Minn.	President, Carl Eggers
1937 - Saranac Lake	President, Leo Eloesser
1938 - Atlanta	President, Stuart W. Harrington
1939 - Los Angeles	President, Harold Brunn
1940 - Cleveland	President, Adrian V. S. Lambert
1941 - Toronto	President, Fraser B. Gurd
1944 - Chicago	President, Frank S. Dolley
1946 - Detroit	President, Claude S. Beck
1947 - St. Louis	President, I. A. Bigger
1948 - Quebec	President, Alton Ochsner
1949 - New Orleans	President, Edward D. Churchill
1950 - Denver	President, Edward J. O'Brien
1951 - Atlantic City	President, Alfred Blalock
1952 - Dallas	President, Frank B. Berry
1953 - San Francisco	President, Robert M. Janes
1954 - Montreal	President, Emile Holman
1955 - Atlantic City	President, Edward S. Welles
1956 - Miami Beach	President, Richard H. Meade
1957 - Chicago	President, Cameron Haight
1958 - Boston	President, Brian Blades
1959 - Los Angeles	President, Michael E. De Bakey
1960 - Miami Beach	President, William E. Adams
1961 - Philadelphia	President, John H. Gibbon, Jr.
1962 - St. Louis	President, Richard H. Sweet (Deceased 1-11-62)
1963 - Houston	President, O. Theron Clagett
1964 - Montreal	President, Julian Johnson
1965 - New Orleans	President, Robert E. Gross
1966 - Vancouver, B.C.	President, John C. Jones
1967 - New York	President, Herbert C. Maier
1968 - Pittsburgh	President, Frederick G. Kergin
1969 - San Francisco	President, Paul C. Samson
1970 - Washington, D.C.	President, Edward M. Kent
1971 - Atlanta	President, Hiram T. Langston
1972 - Los Angeles	President, Thomas H. Burford
	President, John W. Strieder

INDEX OF AUTHORS

	Page
Adkins, Paul C., Washington, D.C.	22
*Anderson, Kathryn D., Washington, D.C.	6
*Asher, William J., Philadelphia, Pa.	42
Austen, W.G., Boston, Mass.	46
*Baxter, James, New York, N.Y.	42
*Benzing, George, III, Cincinnati, Ohio	10
Berger, R.L., Boston, Mass.	44
*Berkovits, Barouh V., Los Angeles, Calif.	36
Bernhard, William F., Boston, Mass.	14
*Bolooki, Hooshang, Miami, Fla.	44
*Bonchek, Lawrence I., Portland, Oreg.	60
Bowman, Frederick O., Jr., New York, N.Y.	8
*Brawley, Robert K., Baltimore, Md.	56
*Brooks, Helene, Philadelphia, Pa.	42
*Buckberg, Gerald D., Los Angeles, Calif.	30
Buckley, M.J., Boston, Mass.	46
*Burdick, James F., Bethesda, Md.	22
*Carlson, Robert G., New York, N.Y.	42
*Carpentier, A., Paris, France	54
*Castonguay, Yves R., Montreal, Quebec	48
*Charrette, E.J.P., Kingston, Ontario	28
*Ching, Nathaniel, New York, N.Y.	52
*Christiansen, Christine, Bethesda, Md.	22
*Collins, John, Chicago, Ill.	16
Connolly, John E., Irvine, Calif.	32
*Craenen, Josepha., Columbus, Ohio	12
Crawford, E. Stanley, Houston, Texas	26
Daggett, W.M., Boston, Mass.	46
*de Leval, M., San Francisco, Calif.	34
*Deloche, A., Paris, France	54
*DeSanctis, R.W., Boston, Mass.	46
*DiOrio, David A., Atlanta, Ga.	18
*Donahoo, James S., Baltimore, Md.	56
*DuBost, Ch., Paris, France	54
*Edie, Richard N., New York, N.Y.	8
Ehrenhaft, J.L., Iowa City, Iowa	58
*Ellis, Kent, New York, N.Y.	8
Faber, L. Penfield, Chicago, Ill.	22
*Falk, Emily A., New York, N.Y.	26
*Farge, Cl., Paris, France	54
*Fellows, Kenneth E., Boston, Mass.	14
*Fennessy, J. J., Chicago, Ill.	20
*Fields, Josh, Los Angeles, Calif.	36
*Finnegan, James O., Philadelphia, Pa.	58
*Futagawa, Shunji, Tokyo, Japan	24
*Fyler, Donald C., Boston, Mass.	14
*Gee, J.B.L., New Haven, Conn.	16
Gerbode, Frank, San Francisco, Calif.	34

*By invitation

	Page
*Gersony, Welton, New York, N.Y.	8
*Gill, Michael, Miami, Fla.	44
*Gingell, Robert, Baltimore, Md.	8
Glenn, W.W.L., New Haven, Conn.	16
*Goodin, R.R., Louisville, Ky.	46
Gott, Vincent L., Baltimore, Md.	56
*Gray, Dennis C., Philadelphia, Pa.	58
Green, George E., New York, N.Y.	48
*Griep, Randall B., Stanford, Calif.	30
*Grondin, Claude M., Montreal, Quebec	48
Grondin, Pierre, Montreal, Quebec	48
Gross, Robert E., Boston, Mass.	14
Haller, J. Alex, Baltimore, Md.	8
*Hanania, A., Paris, France	54
Hatcher, Charles R., Jr., Atlanta, Ga.	18
Head, Louis R., Chicago, Ill.	16
Helmsworth, James A., Cincinnati, Ohio	10
*Henderson, R.D., Toronto, Ontario	24
*Hill, J.D., San Francisco, Calif.	34
Hill, Lucius D., Seattle, Wash.	20
*Ho, Charles, Baltimore, Md.	8
*Hogan, J., New Haven, Conn.	16
*Holcomb, W.G., New Haven, Conn.	10
*Hosier, Don M., Columbus, Ohio	12
*Hottenrott, Christof E., Los Angeles, Calif.	30
*Hutchinson, John E., III, New York, N.Y.	48
*Indeglia, R., Providence, R.I.	34
*Inglesby, T.V., Louisville, Ky.	46
*Isom, O. Wayne, New York, N.Y.	26
*Iyengar, S.R.K., Kingston, Ontario	28
Jensik, Robert J., Chicago, Ill.	22
Johnson, Julian, Philadelphia, Pa.	58
*Jolly, Philip C., Seattle, Wash.	20
*Joseph, William L., Washington, D.C.	22
*Joyner, Claude R., Philadelphia, Pa.	58
*Kakos, Gerard S., Columbus, Ohio	12
*Kammloft, G.W., Brooklyn, N.Y.	52
*Kaplan, Samuel, Cincinnati, Ohio	10
*Kemp, Harvey G., New York, N.Y.	48
Kilman, James W., Columbus, Ohio	12
*Kim, C.S., New Haven, Conn.	16
Kittle, C.F., Chicago, Ill.	20
*Klopp, Edward H., Baltimore, Md.	56
*Kongtahworn, C., Iowa City, Iowa	58
*Kurkji, Henry J., Los Angeles, Calif.	30
*Kutin, Neil D., New York, N.Y.	26
*Lambert, E., Buffalo, N.Y.	10
*Landé, Arnold J., New York, N.Y.	42
*Lansing, A.M., Louisville, Ky.	46
*Lawless, Peter, Seattle, Wash.	20

*By invitation

	Page
*Lepage, Gilles, Montreal, Quebec	48
*Li, Norman N., Philadelphia, Pa.	42
Lillehei, C. Walton, New York, N.Y.	42
*Londe, Stephen P., Rochester, Minn.	54
Lynn, R.B., Kingston, Ontario	28
*Macur, M.F., San Francisco, Calif.	34
MacVaugh, Horace, III, Philadelphia, Pa.	58
*Mallon, Steven, Miami, Fla.	44
Malm, James R., New York, N.Y.	8
Maloney, James V., Los Angeles, Calif.	30
*Martineau, Jean Paul, Montreal, Quebec	48
*Matano, I., New Haven, Conn.	16
Matloff, Jack M., Los Angeles, Calif.	36
*McCart, Peter, Irvine, Calif.	32
*Meere, Claude, Montreal, Quebec	48
*Mekhjian, Haroutune A., New York, N.Y.	48
*Mielke, C.H., San Francisco, Calif.	34
*Milloy, Frank J., Chicago, Ill.	22
*Mills, Noel L., New Orleans, La.	38
*Monson, David O., Chicago, Ill.	22
*Motoyama, E.K., New Haven, Conn.	16
Mundth, E.D., Boston, Mass.	46
Nealon, Thomas F., Jr., New York, N.Y.	52
Ochsner, John L., New Orleans, La.	38
*Olszowka, J., Buffalo, N.Y.	10
*Palmer, Arthur, Chicago, Ill.	16
*Patterson, Russell H., Jr., New York, N.Y.	42
Pearson, F.G., Toronto, Ontario	24
*Piwnica, A., Paris, France	54
*Poirier, R.A., New Haven, Conn.	16
*Ramasamy, N., Brooklyn, N.Y.	52
Randolph, Judson G., Washington, D.C.	6
*Revilla, Antonio, Baltimore, Md.	56
*Roberts, Enrique Bonfils, New York, N.Y.	52
Rossi, N.P., Iowa City, Iowa	58
*Saini, V.K., Boston, Mass.	44
Sawyer, P.N., Brooklyn, N.Y.	52
*Schreiber, J. Tracy, Cincinnati, Ohio	10
*Schwartz, David C., Cincinnati, Ohio	10
*Shaker, I.J., Baltimore, Md.	8
Shumway, Norman E., Stanford, Calif.	30
Simeone, F.A., Providence, R.I.	34
*Sommer, Leonard, Miami, Fla.	44
Spencer, Frank C., New York, N.Y.	26
*Srinivasan, S., Brooklyn, N.Y.	52
*Stanczewski, B., Brooklyn, N.Y.	52
*Stanton, William E., Irvine, Calif.	32
Starr, Albert, Portland, Oreg.	60
*Steier, Michael, New York, N.Y.	52
*Stein, T. Peter, Philadelphia, Pa.	42

*By invitation

	Page
Stemmer, Edward A., Irvine, Calif.	32
*Stempak, J.G., Brooklyn, N.Y.	52
*Stinson, Edward B., Stanford, Calif.	30
*Subramanian, S., Buffalo, N.Y.	10
*Sugiura, Mitsuo, Tokyo, Japan	24
Symbas, P.N., Atlanta, Ga.	18
*Tamura, H., Providence, R.I.	34
*Thibault, William, Irvine, Calif.	32
*Thomas, Arthur N., San Francisco, Calif.	18
*Titus, Jack L., Rochester, Minn.	54
Trusler, G.A., Toronto, Ontario	12
*Tyras, D.H., Atlanta, Ga.	18
*Vargas, Abelardo, Miami, Fla.	44
*Venugopal, P., Buffalo, N.Y.	10
*Vlad, P., Buffalo, N.Y.	10
*Wagner, H., Buffalo, N.Y.	10
*Wallace, Herbert W., Philadelphia, Pa.	42
Wallace, Robert B., Rochester, Minn.	54
*Ware, R.E., Atlanta, Ga.	18
*Wells, Samuel A., Durham, N.C.	22
*West, Thomas, Seattle, Wash.	20
Wheat, M.W., Jr., Louisville, Ky.	46
*Williams, Thomas E., Jr., Columbus, Ohio	12
*Witoszka, M., Providence, R.I.	34
*Wolfe, Walter G., Washington, D.C.	22
*Zubrow, Marc T., Philadelphia, Pa.	42

*By invitation