GRANT NUMBER DAMD17-94-J-4116

TITLE: Biology of Breast Cancer: A Predoctoral Training Program

PRINCIPAL INVESTIGATOR: Nita J. Maihle, Ph.D.

CONTRACTING ORGANIZATION: Mayo Clinic and Foundation
Rochester, Minnesota 55905

REPORT DATE: August 1998

TYPE OF REPORT: Final

PREPARED FOR: Commander
U.S. Army Medical Research and Materiel Command
Fort Detrick, Frederick, Maryland 21702-5012

DISTRIBUTION STATEMENT: Approved for public release;
distribution unlimited

The views, opinions and/or findings contained in this report are
those of the author(s) and should not be construed as an official
Department of the Army position, policy or decision unless so
designated by other documentation.

DTIC QUALITY INSPECTED 4

19990811 133
Implementation of a new predoctoral program in the "Biology of Breast Cancer" has facilitated the training of investigators committed to future careers in the study of breast cancer. The scope of this program has been limited to the training of predoctoral (i.e., Ph.D. and M.D., Ph.D.) candidates. USAMRDC support for this program has resulted in the development of a truly outstanding, multidisciplinary, didactic curriculum in tumor biology, which includes a strong emphasis in breast cancer. To date, 16 trainees have matriculated into this new training program. Two trainees have successfully completed this training program and have left the Mayo Clinic to continue their training/careers in breast cancer research. All of the remaining trainees are conducting breast cancer relevant thesis research and continue to make excellent progress in their studies. This final report includes the product of our last task (i.e., Task 6) of our original statement of work which is a formal written evaluation, and also includes the comments of our two external reviewers (see Appendix). The longer term continuation and success of this new training program has been assured by the recent (8/98) award of a new predoctoral training grant from the NCI (CA 75926).
FOREWORD

Opinions, interpretations, conclusions and recommendations are those of the author and are not necessarily endorsed by the U.S. Army.

Where copyrighted material is quoted, permission has been obtained to use such material.

Where material from documents designated for limited distribution is quoted, permission has been obtained to use the material.

Citations of commercial organizations and trade names in this report do not constitute an official Department of Army endorsement or approval of the products or services of these organizations.

In conducting research using animals, the investigator(s) adhered to the "Guide for the Care and Use of Laboratory Animals," prepared by the Committee on Care and Use of Laboratory Animals of the Institute of Laboratory Resources, national Research Council (NIH Publication No. 86-23, Revised 1985).

For the protection of human subjects, the investigator(s) adhered to policies of applicable Federal Law 45 CFR 46.

In conducting research utilizing recombinant DNA technology, the investigator(s) adhered to current guidelines promulgated by the National Institutes of Health.

In the conduct of research utilizing recombinant DNA, the investigator(s) adhered to the NIH Guidelines for Research Involving Recombinant DNA Molecules.

In the conduct of research involving hazardous organisms, the investigator(s) adhered to the CDC-NIH Guide for Biosafety in Microbiological and Biomedical Laboratories.

[Signature] 8/15/98

FI - Signature  Date
# Annual Report for DAMD 17-94-J-4116

**August, 1998**

**Biology of Breast Cancer: A Predoctoral Training Program**

## Table of Contents

<table>
<thead>
<tr>
<th>Item</th>
<th>Page</th>
</tr>
</thead>
<tbody>
<tr>
<td>Front Cover</td>
<td>1</td>
</tr>
<tr>
<td>Standard Form (SF) 298</td>
<td>2</td>
</tr>
<tr>
<td>Foreword</td>
<td>3</td>
</tr>
<tr>
<td>Table of Contents</td>
<td>4</td>
</tr>
<tr>
<td>Introduction</td>
<td>5</td>
</tr>
<tr>
<td>Body</td>
<td>6-15</td>
</tr>
<tr>
<td>Conclusions</td>
<td>15</td>
</tr>
<tr>
<td>References</td>
<td>15</td>
</tr>
<tr>
<td>Appendices (Listing)</td>
<td>16</td>
</tr>
<tr>
<td>A. Table I: Academic Credentials of Trainees</td>
<td>17</td>
</tr>
<tr>
<td>B. Table II: BBC Trainees, Their Mentors, and Thesis Titles</td>
<td>18</td>
</tr>
<tr>
<td>C. Table III: Publications of BBC Trainees</td>
<td>19-24</td>
</tr>
<tr>
<td>D. Table IV: Meetings/Extramural Courses Attended by BBC Trainees</td>
<td>25</td>
</tr>
<tr>
<td>E. BBC Faculty and Their Research Interests</td>
<td>26-28</td>
</tr>
<tr>
<td>F. Selected Course Syllabuses and Lecture Outlines</td>
<td>29-43</td>
</tr>
<tr>
<td>G. ‘97 - ‘98 Journal Club Titles</td>
<td>44-46</td>
</tr>
<tr>
<td>H. Internet Home Page for the Biology of Breast Cancer/Tumor Biology Program</td>
<td>47-48</td>
</tr>
<tr>
<td>I. Recruitment Advertisement (Example)</td>
<td>49</td>
</tr>
<tr>
<td>K. BBC Training Program Extramural Support</td>
<td>51</td>
</tr>
<tr>
<td>L. Written Self Study</td>
<td>52</td>
</tr>
<tr>
<td>M. Dr. Harold Moses’ Critique (External Reviewer)</td>
<td>53</td>
</tr>
<tr>
<td>N. Dr. Mina Bissell’s Critique (External Reviewer)</td>
<td>54</td>
</tr>
<tr>
<td>O. AACE Abstract</td>
<td>55</td>
</tr>
<tr>
<td>P. List of Personnel</td>
<td>56</td>
</tr>
</tbody>
</table>
INTRODUCTION

This annual report covers activities for DAMD 17-94-J-4116 entitled the "Biology of Breast Cancer: A Predoctoral Training Program" for the period September, 1997 through August, 1998, and also provides a cumulative summary of the activities of this new training program since its inception. In this report we present documentation for the successful completion of the past academic year, and the initiation of the 1998-1999 academic year.

The six tasks that were presented in the original application are listed, below:

- **Task 1:** Organize Biology of Breast Cancer Predoctoral Training Program Faculty and Curriculum.
- **Task 2:** Establish New Courses in Specialized Aspects of Tumor Biology, Emphasizing the Cell and Molecular Biology of Breast Cancer.
- **Task 3:** Establish Appropriate Mechanisms for Student Recruitment Into This New Training Program.
- **Task 4:** Implement the New Biology of Breast Cancer Predoctoral Training Program Curriculum.
- **Task 5:** Assess Student Progress.
- **Task 6:** Assess Biology of Breast Cancer Predoctoral Training Program Effectiveness and Formalize Assessment in a Written Report.

As noted in the written review of our last progress report (February 12, 1997), we had successfully completed all of these tasks except Task 6 at the time of our last report. Enclosed as a component of this final report (Appendix L) is the formal written self evaluation that was submitted to members of our External Advisory Committee (Drs. Bissell, King and Moses) in June, 1998. The comments of two of these external reviewers (Drs. Bissell and Moses) are also included in the Appendix (Appendices M, N).
Goals

The goals of the Biology of Breast Cancer (BBC) and Tumor Biology Training Program are fourfold:

- First, to provide trainees with a solid and uniquely multidisciplinary knowledge base in the biology of cancer using breast cancer as the paradigm.
- Second, to guide the development of each individual trainee so that they achieve their fullest academic and research potential.
- Third, to aid trainees in the establishment of their professional network of peers and colleagues in the field of breast cancer research.
- Fourth, to stimulate new working alliances between students, fellows, and staff participating in breast cancer research, education, and clinical endeavors at the Mayo Clinic and within the Mayo Cancer Center.

Overview

The “Biology of Breast Cancer” is a multidisciplinary predoctoral training program in the biology of cancer, with a specific emphasis on breast cancer. The focus of the program is to provide an educational environment that stimulates excellence in scientific thought and training while simultaneously providing exposure to all of the major fields of study relevant to tumor biology. While a defining feature of this program is its research focus and integral link with clinical aspects of breast cancer, a general foundation in tumor biology is both important and essential in achieving this goal. The curriculum for this program is outlined in the course syllabus material, provided below, and the thesis research topics of the students matriculating in the program. This information clearly details and substantiates the major breast cancer research focus of this new training program. Research and training are broadly focused on gene regulation, cell cycle control, cancer genetics, oncogene and tumor suppressor action, tumor immunology, signal transduction, antitumor pharmacology, with a particular emphasis on breast cancer, but also includes investigators with research programs in ovarian, uterine, lung, G.I., brain, and prostate cancers. Students participate in laboratory-based research, as well as in a formal tumor biology curriculum that integrates current concepts in cell growth control with the natural history of human tumors.

The Biology of Breast Cancer Training Program has been supported by extramural training grants since its inception. Currently, these training grants include an award from the US Army Medical Research and Materiel Command in the “Biology of Breast Cancer” (DAMD17-94-J-4116), and a T32 Training Grant from the National Cancer Institute in “Tumor Biology” (CA75926). In addition, the program operates under the generous support of the Mayo Foundation through the Mayo Graduate School and the Mayo Cancer Center. Individual trainees also have been successful in competition for individual research awards (see Appendix K).
Program Structure

Administrative Structure

The BBC training program is administrated through the Mayo Graduate School, and is closely allied with the Mayo Cancer Center. Day-to-day program administration operates largely through the Director (Dr. Salisbury) and Co-director (Dr. Maihle) of the Biology of Breast Cancer and Tumor Biology Training grants. Long range planning and administration operates through the Tumor Biology Education Committee (Drs. Salisbury, Maihle, Federspiel, Jelinek, and Tindall, and a trainee, Mr. J. Baines). The Education Committee meets each academic quarter to discuss student recruitment, student progress, and coordination within the Tumor Training Program curriculum. In addition, the directors of the three cancer-related pre- and postdoctoral training grants (Drs. Salisbury, Maihle, Getz, and David) and the Director of the Mayo Cancer Center (Dr. Prendergast) interact to coordinate ongoing programs and activities related to cancer research and education at Mayo in general. The Biology of Breast Cancer Training faculty also meet quarterly, in addition to frequent interactions through participation in program courses, journal clubs, research workshops, and a biweekly social hour called the “Tumor Biology Tea.”

Qualifications of the Program Faculty

The BBC training faculty (see Appendix E for listing) consists of approximately 50 full and associate members drawn from each of the basic science departments, as well as clinical faculty who participate in scholastic activities of the program but who do not have active research laboratories. The level of individual faculty participation varies each year for specific courses, topics and journal clubs. Nonetheless, a growing and enthusiastic cadre of participating faculty has emerged. In future years the program may elect to restructure its faculty based on degree of faculty participation, given the mounting enthusiasm for this program, as well as recent and ongoing recruitment of new staff in the area of cancer biology. At this time, however, faculty privileges will remain as they are in order to promote both the inclusively and multidisciplinary nature of this new training program. While most individual faculty are associated with traditional discipline-based basic science and clinical departments (such as Oncology, Molecular Biology, Experimental Pathology, Pharmacology, etc.) the administrative structure of the Program is that of an interdisciplinary programmatic unit. This programmatic structure reflects the interdisciplinary nature of the major research and academic efforts of the associated faculty.

- Full members of the training faculty have an established track record of accomplishment in biomedical research as demonstrated by significant publications of high scientific merit, excellence, and innovation. Overall, the faculty have consistent records of extramural funding in support of their individual research programs.
- The collective interests of this training faculty are quite broad, but all show direct breast cancer relevance. These interests include: cell signaling, cancer genetics, gene regulation, tumor immunology, oncogene and tumor suppressor action, cell cycle regulation, tumor virology, gene therapy, hormonal regulation, and molecular cytology. Most training faculty are also members of the NCI-designated Mayo Cancer Center.
- Faculty drawn from both clinical and basic science departments contribute to the Training Program through participation in a variety of relevant educational activities and as clinical instructors. Faculty from outside the program may serve as advisory members of qualifying examination and thesis committees, however, they may not serve as research mentors for students in the Biology of Breast Cancer Training Program.
Program Curriculum

Introduction

The program curriculum and thesis research is a predoctoral training program leading to the Ph.D. degree in Biomedical Sciences. Each year, 3 to 5 students are accepted into the program for an appointment term of 4 to 5 years. The program strives to maintain a steady state level of 15 to 20 students. Trainee stipends initially are supported through institutional funds. Students qualify for support from extramural training grants (third through fifth year), following successful completion of the qualifying examination. The training program curriculum includes didactic course work, journal clubs, research seminars and workshops, and tutorial and special clinical activities. Students who matriculate into the program must meet the general course requirements of the Mayo Graduate School in which a minimum of 15 credits are required from the Graduate School Core Curriculum. Students must also complete 20 credits from the didactic Tumor Biology Program curriculum and the remainder of their credit requirements can be completed through elective courses offered by the Tumor Biology Program, the Mayo Graduate School or by special topics courses given at other institutions and sanctioned by the Mayo Graduate School. For students in the Biology of Breast Cancer Program these electives must include a course entitled “Biology of Breast Cancer” (see Appendix F for outlines of selected courses). A student's program of core courses is individually developed by the Education Committee in consultation with the student and his/her advisor.

Summary of the Curriculum:

Graduate School Core Offerings (15 Credits, minimum)

- Genome Biology (3)
- Immunology (3)
- Principles of Cell and Tissue Design (3)
- Biochemistry (3)
- Genetics (1)
- Pharmacology (2)
- Developmental Biology & Statistics (1)
- Biology of Disease (1)
- Ethics (1)

Required Biology of Breast Cancer Course Offerings (20 Credits)

- Tumor Biology I: Introduction to Tissue and Tumor Biology (3)
- Tumor Biology II: Origins of Human Cancer (3)
- Tumor Biology III: Growth Factors, Oncogenes, and Tumor Suppressors (3)
- Biology of Breast Cancer (1)
- Current Topics in Tumor Biology: Journal Club (1) x 3
- Research Seminars in Tumor Biology and Tumor Biology Interest Group (1)
- AACR Course in Histopathology of Cancer, Keystone CO. (1)
- Laboratory Rotations in Tumor Biology (3 required rotations, 2 credits each = 6 total)
- Research in Tumor Biology (Thesis Research (0))

Recommended Electives

- Quantitative Biology I-III, Neuroscience (1), Integrated Physiology (5)
- Tumor Immunology (1), Business of Science and the Science of Business (1)
- Cytogenetics (2)
Biology of Breast Cancer Track and Core Curriculum Schedule

<table>
<thead>
<tr>
<th>Year I or II</th>
<th>M</th>
<th>T</th>
<th>W</th>
<th>Th</th>
<th>F</th>
</tr>
</thead>
<tbody>
<tr>
<td>Fall Quarter</td>
<td>9:00-10:00</td>
<td>Biochemistry</td>
<td>Immunology</td>
<td>Biochemistry</td>
<td>Immunology</td>
</tr>
<tr>
<td>10:00-11:00</td>
<td>Genome Biology</td>
<td></td>
<td>Genome Biology</td>
<td></td>
<td>Genome Biology</td>
</tr>
<tr>
<td>11:00-12:30</td>
<td>Tumor Biology I</td>
<td>Tumor Biology</td>
<td>Tumor Biology Journal Club</td>
<td>Tumor Biology I</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Year 1 or II</th>
<th>M</th>
<th>T</th>
<th>W</th>
<th>Th</th>
<th>F</th>
</tr>
</thead>
<tbody>
<tr>
<td>Winter Quarter</td>
<td>9:00-10:00</td>
<td>Cell Biology</td>
<td>Genetics</td>
<td>Cell Biology</td>
<td>Cell Biology</td>
</tr>
<tr>
<td>11:00-12:30</td>
<td>Tumor Biology II</td>
<td>Tumor Biology</td>
<td>Tumor Biology Journal Club</td>
<td>Tumor Biology II</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Year 1 or II</th>
<th>M</th>
<th>T</th>
<th>W</th>
<th>Th</th>
<th>F</th>
</tr>
</thead>
<tbody>
<tr>
<td>Spring Quarter</td>
<td>9:00-10:00</td>
<td>Pharmacology</td>
<td>Pharmacology</td>
<td>Pathobiology</td>
<td></td>
</tr>
<tr>
<td>10:00-11:00</td>
<td>Development &amp; BioStatistics</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>11:00-12:30</td>
<td>Tumor Biology III</td>
<td>Tumor Biology</td>
<td>Tumor Biology Journal Club</td>
<td>Tumor Biology III</td>
<td></td>
</tr>
</tbody>
</table>

Additional advanced elective courses may be chosen in any area from the Mayo Graduate School Bulletin to fulfill the overall degree requirement of 35 credits. In addition, all students in the training program are required to take formal classes in Radiation Safety, Animal Care and Use, and also participate in an NIH Grant Writing Workshop. These required courses are not administrated by the Mayo Graduate School, and, therefore, are not offered for Graduate School credit.
Program Implementation

Innovative Strategies for Teaching

Tumor Biology track courses establish a solid foundation in the biology of cancer. In practice, the biology of breast cancer is emphasized as an illustrative paradigm throughout the program curriculum. Issues of fundamental importance to understanding breast cancer are also relevant to other cancers, and the converse is also true. Likewise, there are many instances where the illustrating example of a relevant point may involve biological systems as diverse as yeast and frogs (not breast cancer research, but directly relevant to improving our understanding of breast cancer). Thereby, the tumor biology curriculum is grounded in basic cancer biology with the aim of leading students to a thorough understanding of all aspects at the forefront of breast cancer research. In order to accomplish this, our curriculum has been strategically developed to provide a strong foundation in the study of cancer using breast cancer as the model wherever possible. For example, to illustrate the significant emphasis on and integration of breast cancer in the overall curriculum, topics in breast cancer are featured in three 3 credit hour courses in the curriculum (see below), as well as in approximately one out of every three journal club presentations. Moreover, all trainees are required to register for a didactic course in “The Biology of Breast Cancer” (TBIO 8305, page 134 in the Mayo Graduate School Bulletin). The three major courses were organized and are taught using breast cancer as the principle paradigm for instruction and content: “Introduction to Tissue and Tumor Biology” (Tumor Biology I, TBIO 5000), “Origins of Human Cancer” (Tumor Biology II, TBIO 8000), and “Growth Factors, Oncogenes and Tumor Suppressors” (Tumor Biology III, TBIO 8005) (see course outlines in Appendix F).

Tumor Biology Track courses share several distinguishing features and innovative strategies for teaching which enhance learning and student/faculty participation.

- **Course Structure:** Tumor Biology I, II, and III are given as an integrated series during the first three consecutive quarters following matriculation. Individually, these courses meet three times per week (1 1/2 hour per session) with an overview and historical review of a selected topic(s) for the current week presented in a didactic lecture format during the first class session. This is followed by a student presentation of a current or historically relevant research paper(s) in the area of the week’s topic during the second class session using the journal club format. Finally, a round table small group problem set discussion format is used to focus on questions and problems relevant to the week’s topic during the third class session. The research paper presentation and problem set discussions are carefully organized in order to thoroughly integrate research topics relevant to the theme of the week. The effectiveness of these active learning/teaching strategies is apparent to all participants in these courses. The active involvement of students in the learning and discovery process, in information processing, and in the application of information to problems requires that students are accountable for learned information on an immediate and ongoing basis. Problem centered learning also puts learning into context and facilitates learning transfer. These sessions allow students to organize and categorize information into meaningful units and to ‘discover’ novel relationships and extract and assimilate important points in an interactive and participatory manner.
• **Balance of Historic and Current Scientific Perspectives:** Given the rapid pace of progress in the biological sciences and the exponential rate of growth of relevant literature, the general philosophy that is promoted within the program is to teach less better. The objective here is to lay a strong foundation in cancer biology with the clear understanding that what is particularly relevant and important today, may not be so tomorrow. Therefore, emphasis is placed on developing effective learning skills, and the application of these skills to both historic paradigms, as well as critical review and evaluation of issues at the forefront of modern cancer biology.

• **Commitment, Accountability, and Responsibility:** Integral to the Tumor Biology Program teaching philosophy is Peer Performance Assessment and the Team Learning Model. These strategies create a climate in which all students are encouraged to grow. This results in a classroom environment where students from diverse backgrounds (including clinical fellows) feel welcome to fully participate in discussions and problem solving. In this way, desired student performances are tied directly to the efforts of the students themselves, to the involvement of students in the teaching-learning process, to the opportunities to make choices, and to the degree to which they interact with their peers and instructors. Emphasis is placed on organization and presentation skills, accountability tracking, and peer assessment and feedback. Our experience with this learning model is that trainees rapidly gain a level of professional expectation of their peers (and themselves) that both promotes and enhances the general level of academic and scholastic effort among trainees.

• **Continuing Education:** Senior students who have completed the formal didactic course requirements, and postdoctoral fellows who are supported by other cancer-related training grants actively participate in the journal club sessions that are an integral component of the Tumor Biology curriculum. In this way, senior trainees contribute to the critical mass of the class and also enhance the sophistication and the multidisciplinary nature of the discussions. Through this mechanism, more advanced trainees revisit current topics in cancer biology throughout their advanced training years. In this manner, advanced trainees have the opportunity to reinforce key concepts, to help new trainees better understand these concepts, and also to remain current with the rapid pace of development in the dynamic field of cancer biology. Likewise, Program faculty (and their laboratory personnel) actively participate in these regular weekly journal club sessions.

• **Integration of the Clinical Perspective:** The training program is designed to give the trainee a broad and well-rounded understanding of cancer from the basic science,
population science, and clinical perspectives. Integration of the clinical activities of the Mayo Cancer Center into the training program is achieved in several ways:

◊ First, clinical fellows and residents from a broad spectrum of cancer relevant programs (e.g., oncology, hem/onc, orthopedic research, gynecologic oncology, pediatric oncology, etc.) formally enroll and participate in training program courses and journal clubs. Active participation by clinical residents and fellows adds considerably to the multidisciplinary perspective of the student body, and to stimulating class discussions.

◊ Second, clinical staff give didactic lectures in areas of their specialty in Biology of Breast Cancer Training Program courses. For example in Tumor Biology II and III, individual lectures are given by clinical staff practicing in Surgery, Medical Oncology, Radiation Oncology, and Surgical Pathology.

◊ Third, program trainees are required to attend Mayo Cancer Center Research Workshops, Mayo Cancer Center Grand Rounds, and appropriate departmental, Research Society, Oncology Society and Hematology Society lectures, and receive course credit for participating in these activities (TBio 5101: “Research Seminars in Tumor Biology”). Students are made aware of relevant speakers through Email, direct mailings, and weekly announcements during the weekly journal club.

◊ The program curriculum is integrated with clinical practice wherever possible through special course related activities. For example, small group tours of the Surgical Pathology Suite (in TBio I) allow students to observe gross dissection of surgical specimens (including breast tumors), rapid freezing and cryomicrotomy, microscopic examination (via video monitors) and diagnosis by staff pathologists, and reporting to operating room surgeons. Additionally, Tumor Biology Program trainees are required to attend clinical rounds with a Mayo staff member (any division or department). Direct exposure to clinical activities such as these are useful for students to understand, based on first-hand observation, the intensity, dedication, and skill involved in the clinical care and treatment of cancer patients. These activities also promote the involvement of our clinical faculty in Biology of Breast Cancer Training Program activities.

◊ Finally, Tumor Biology trainees may have one or more clinical staff advisors participate as members of their Thesis Advisory Committee. Sometimes this involvement is fairly technical, e.g., participation by a Mayo Cancer Center biostatistician in study design and analysis. In other instances, however, clinical advisors may be directly involved in helping the trainee define a clinically relevant question, and/or assist them with tumor specimen acquisition and/or data analysis.

Additional Academic Activities

Seminars by Students, Faculty and Invited Speakers: Extensive institutional resources support seminars by nationally and internationally recognized scientists and clinicians on the Mayo Rochester campus. Approximately 350-400 speakers come to the Rochester campus each year. Trainees are, therefore, exposed to diverse biomedical research opportunities, and institutionally and departmentally-based research seminars throughout the year. In December of 1997, students of the Tumor Biology Program hosted Dr. Judah Folkman (Harvard University) who presented the Annual Findling Lecture of the Mayo Graduate School. The 1998-1999 academic year will feature a series of visiting speakers who will focus on the broad topic of Epigenetics and Cancer. During each of their research years, Tumor Biology Trainees also present research seminars and research posters in multidisciplinary research workshops and retreats (e.g., the Mayo Graduate School Annual Research Symposium, the Joint Mayo Cancer Center/Laboratory Medicine Retreat). More recently, we have formalized the Tumor Biology Interest Group (TBIG) for Mayo Graduate
School course credit. This monthly research workshop provides the opportunity for all Tumor Biology trainees (pre- and postdoctoral) to regularly present their research plans, proposals, and results in a constructively critical internal forum.

**Attendance at National Research Meetings:** All students are supported to attend at least one national scientific meeting each year even if they are not presenting an abstract. If they are presenting their work, attendance at additional meetings is encouraged and supported by the research mentor’s laboratory. Mentors take an active role in introducing students to the professional culture and ‘networking’ critical to success in any biomedical research career through this mechanism. In recent years, Biology of Breast Cancer Training Program trainees have attended and presented at the following national meetings: AACR, ASCB, FASEB, Annual Oncogene Meetings, Annual Human Cancer Meeting, Cold Spring Harbor Cancer Genetics Meetings, Salk Tyrosine Phosphorylation Meetings, and various Gordon and Keystone Conferences (see Table IV, Appendix D).

**Research Training**

**Selection of Thesis Laboratory, Mentor, and Thesis Committee Members:** Trainees typically matriculate in June through August and are required to complete three laboratory-based (minimum 8 weeks each) rotations during their first year. Any laboratory-based Mayo Graduate School faculty member may serve as a mentor for these research rotations. Selection of the thesis mentor follows completion of successful laboratory-based rotations by mutual consent of the student and mentor with the sanction of the Training Program Education Committee and Mayo Graduate School Education Committee. The qualifying examination consists of a written thesis proposal, an oral presentation (TBIG forum), and its defense before a thesis committee consisting of a minimum of 4 faculty (including the thesis advisor), and when appropriate, an extramural committee member from outside the institution. Typically clinical or extramural committee members’ research specialties are related to the general area of the student’s thesis topic. Following successful completion of the qualifying examination, research progress is assessed through regular Thesis Advisory Committee meetings (minimum of one Thesis Advisory Committee meeting per year). While Thesis Advisory Committee members are available for advice, technical assistance, and consultation throughout the year, these meetings provide a formal opportunity for input by the Thesis Advisory Committee on progress and experimental aspects of the thesis project. The chair of the Thesis Advisory Committee formally reports the outcome of each committee meeting in writing to the Training Program Education Committee and to the Mayo Graduate School.

**Thesis Research:** The Biology of Breast Cancer Training Program places strong emphasis on thesis research. All laboratory-based faculty have demonstrated records of research training at both the predoctoral and postdoctoral levels. The specific details of an individual student’s research training plan are developed following the selection of a thesis advisor and in consultation with the Thesis Advisory Committee. The thesis research project must be hypothesis driven and experimental in nature and must, in addition, have a direct relevance to the biology of cancer. For a listing of BBC trainees and their thesis titles, see Appendix B.

**Ph.D. Thesis:** The thesis is the most important document that the Ph.D. candidate prepares during the course of graduate study, and is a record of the scientific accomplishments that justify the awarding of the Ph.D. degree. The thesis is archival. Consequently, the Mayo Graduate School has developed standards for its format and style, and our trainees adhere to these guidelines. The thesis examination consists of a formal thesis research seminar open to all members of the Mayo community followed by a meeting with the Thesis Examining Committee.
during which the scientific merit and accomplishments of the candidate are evaluated. Successful completion of a research thesis typically also results in two or more research manuscripts submitted for publication in peer-reviewed journals of high scientific standards.

Student Recruitment, Progress and Track Evaluation

Trainee Candidates: Students recruited into the Biology of Breast Cancer Training Program are selected on the basis of outstanding academic credentials, a stated desire to study and conduct research in the area of breast cancer biology, and an assessment of individual research potential by the training faculty (see Appendix A for table of BBC academic credentials). Many of the applicants to the Mayo Graduate School have had research experience within the Mayo system through summer undergraduate research internships. Typically, candidates for admission to Mayo’s graduate programs apply directly to the Graduate School where their academic credentials, letters of recommendation, and personal statements are placed on record. Applicants are selected for on-site interview by the Mayo Graduate School Admissions Committee. The interview process involves faculty and student assessment of each applicant’s research and academic interests. The Biology of Breast Cancer Training Program also has placed special emphasis on recruitment and training of under-represented minorities. Currently the predoctoral class consists of a total of 16 students, 3 of whom are minorities (19%, including one Native American, and two Hispanic students). A sample student recruitment advertisement is included in the Appendix of this report (Appendix I).

Trainee Evaluation

Trainee evaluation takes place at several levels and is assessed by comparison of established and objective data relating recruitment credentials, program completion, academic performance, placement, and ultimately career achievement.

- Academic performance of trainees, including coursework evaluation and consideration of reports from the trainee’s Qualifying Exam and Thesis Advisory Committees.
- Successful completion of the degree program.
- Success in gaining competitive pre- and postdoctoral fellowships and/or extramural funding.
- Ultimately, the appointment of these trainees to independent research positions with evidence of ongoing research activities relevant to cancer biology.

Curriculum and Program Evaluation

Curriculum and Program evaluation includes the areas listed below, as well as additional areas defined by the External Review Committee:

- Ability to recruit and retain outstanding Ph.D. candidates.
- Course content and appropriateness to the biology of breast cancer.
- Thoroughness of didactic and formal training in the biology of cancer.
- Effectiveness of teaching and examining methods and procedures.
- Vitality and effectiveness of student/faculty interactions in the academic components of the program.
- Evidence of faculty mentorship and establishment of intramural and extramural professional networks.
- Scope and role of individual faculty participation in the Biology of Breast Cancer Training Program.
• Integration of the clinical perspective and understanding of physician and patient concerns in the diagnosis and treatment of cancer.

• Overall effectiveness of the Program Director, Co-Director, Education Committee, and of the Graduate School in administrating the Biology of Breast Cancer Training Program.

In addition to these standards to be used for self-evaluation, the Biology of Breast Cancer Training Program has successfully completed two rigorous external reviews, each of which has resulted in an extramural training award (i.e., the US Army Medical Research and Materiel Command award for predoctoral training in the “Biology of Breast Cancer” (DAMD17-94-J-4116), and a T32 Training Grant from the National Cancer Institute for predoctoral training in “Tumor Biology” (CA75926). One aspect of the USAMRMC predoctoral award is a requirement for formalization of an External Advisory Committee and periodic external review by this Committee. This formal written self evaluation and two external reviewers’ comments (i.e., Dr. M. Bissell and Dr. H. Moses) are included as a component of the Appendix of this final progress report (Appendices L-N).

In addition, our experiences with the development and implementation of this new training program will be presented (poster format) at the 1998 American Association for Cancer Education Meeting to be held in Portland, Oregon (see Appendix O for abstract).

Syllabus Outlines for Selected Tumor Biology Track Courses

Mayo Tumor Biology track courses cover a series of topics of historic relevance and primary importance to cancer biology. In addition, each year course organization and content have evolved according to current trends and in order to incorporate breaking forefront issues in this dynamic field. Course syllabus outlines for 1997-1998, indicating topic, format, and faculty are included in the Appendix (Appendix F).

CONCLUSIONS

In the period 1994-1998 funding from the USAMRMC Breast Cancer Research Program was used to begin the implementation of a new multidisciplinary training program in the “Biology of Breast Cancer” at the Mayo Clinic in Rochester, Minnesota. The initiation of this new training program has resulted in the development of a new didactic curriculum in Tumor Biology with a special emphasis on breast cancer, a new journal club, and new working alliances among Mayo Clinic faculty interested in breast cancer research. To date, a total of sixteen students have matriculated into this training program, two of whom have successfully completed their training and have left the Mayo Clinic to continue their research training/careers in the field of breast cancer research. This new training program has recently completed formal internal and external reviews, and a copy of these reviews is included as a component of this final report. While the leaders of this program have identified future goals and objects to continue to enhance the quality of this new training program the overall consensus from these reviews is that this new program has been quite successful to date, and importantly, provides a truly outstanding foundation for the training of future generations of investigators dedicated to the field of breast cancer research. On behalf of all the participants in this new training program we express our sincere gratitude to the USAMRMC Breast Cancer Research Program for providing us with this opportunity.

REFERENCES - None.
# Appendix

<table>
<thead>
<tr>
<th>Exhibit</th>
<th>Contents</th>
<th>Page</th>
</tr>
</thead>
<tbody>
<tr>
<td>A.</td>
<td>Table I: Academic Credentials of Trainees</td>
<td>17</td>
</tr>
<tr>
<td>B.</td>
<td>Table II: BBC Trainees, Their Mentors, and Thesis Titles</td>
<td>18</td>
</tr>
<tr>
<td>C.</td>
<td>Table III: Publications of BBC Trainees</td>
<td>19-24</td>
</tr>
<tr>
<td>D.</td>
<td>Table IV: Meetings/Extramural Courses Attended by BBC Trainees</td>
<td>25</td>
</tr>
<tr>
<td>E.</td>
<td>BBC Faculty and Their Research Interests</td>
<td>26-28</td>
</tr>
<tr>
<td>F.</td>
<td>Selected Course Syllabuses and Lecture Outlines</td>
<td>29-43</td>
</tr>
<tr>
<td>G.</td>
<td>'97 - '98 Journal Club Titles</td>
<td>44-46</td>
</tr>
<tr>
<td>H.</td>
<td>Internet Home Page for the Biology of Breast Cancer/</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Tumor Biology Program</td>
<td>47-48</td>
</tr>
<tr>
<td>I.</td>
<td>Recruitment Advertisement (Example)</td>
<td>49</td>
</tr>
<tr>
<td>K.</td>
<td>Summary of BBC Training Program Extramural Support</td>
<td>51</td>
</tr>
<tr>
<td>L.</td>
<td>Written Self Study/Evaluation</td>
<td>52</td>
</tr>
<tr>
<td>M.</td>
<td>Dr. Harold Moses' Critique (External Reviewer)</td>
<td>53</td>
</tr>
<tr>
<td>N.</td>
<td>Dr. Mina Bissell's Critique (External Reviewer)</td>
<td>54</td>
</tr>
<tr>
<td>O.</td>
<td>American Association for Cancer Education Meeting Abstract on Mayo's</td>
<td>55</td>
</tr>
<tr>
<td></td>
<td>Biology of Breast Cancer Program</td>
<td></td>
</tr>
<tr>
<td>P.</td>
<td>List of Personnel</td>
<td>56</td>
</tr>
<tr>
<td>Name</td>
<td>Undergraduate School</td>
<td>GPA</td>
</tr>
<tr>
<td>-----------------</td>
<td>-----------------------------------------------------------</td>
<td>-----</td>
</tr>
<tr>
<td>Adelsman, Margaret</td>
<td>Bemidji State University, Bemidji, MN</td>
<td>3.85/4.0</td>
</tr>
<tr>
<td>Adriance, Melissa*</td>
<td>California State University, Long Beach, CA</td>
<td>2.7/4.0</td>
</tr>
<tr>
<td>Baines, Jonathan*</td>
<td>University of Arizona, Tucson, AZ</td>
<td>3.39/4.0</td>
</tr>
<tr>
<td>Canales, Nohelia*</td>
<td>Mount St. Mary’s College, Los Angeles, CA</td>
<td>3.34/4.0</td>
</tr>
<tr>
<td>Calhoun, Eric</td>
<td>Truman State University, Kirksville, MO</td>
<td>3.65/4.0</td>
</tr>
<tr>
<td>Eley, Gregory</td>
<td>University of Georgia, Athens, GA</td>
<td>3.01/4.0</td>
</tr>
<tr>
<td>Faupel, Jessica</td>
<td>Gettysburg College, Gettysburg, PA</td>
<td>3.3/4.3</td>
</tr>
<tr>
<td>Holmen, Sheri</td>
<td>Western Michigan University, Kalamazoo, MI</td>
<td>3.56/4.0</td>
</tr>
<tr>
<td>Johnson, Julie</td>
<td>University of Wisconsin, Madison, WI</td>
<td>3.38/4.0</td>
</tr>
<tr>
<td>Lomberk, Gwen</td>
<td>Boston College, Chestnut Hill, MA</td>
<td>3.3/4.0</td>
</tr>
<tr>
<td>Myers, Shannon</td>
<td>Southwestern University, Georgetown, TX</td>
<td>3.8/4.0</td>
</tr>
<tr>
<td>Ritland, Steve</td>
<td>University of Wisconsin, Eau Claire, WI</td>
<td>3.54/4.0</td>
</tr>
<tr>
<td>Rogers, Michael</td>
<td>Brigham Young University, Provo, UT</td>
<td>3.66/4.0</td>
</tr>
<tr>
<td>Schehl, Colleen</td>
<td>University of Dayton, Dayton, OH (Bach.) Oklahoma State University, Stillwater, OK (M.S.)</td>
<td>3.3/4.0</td>
</tr>
<tr>
<td>Walters, Denise</td>
<td>McMaster University, Hamilton, Ontario, Canada</td>
<td>10.6/12</td>
</tr>
<tr>
<td>Xu, Kun</td>
<td>Beijing Medical University, Beijing, P.R. China</td>
<td>3.6</td>
</tr>
</tbody>
</table>

*Members of underrepresented minority groups.
# Table II: Trainees, Mentors, Thesis Titles

<table>
<thead>
<tr>
<th>Student</th>
<th>Mentor</th>
<th>Proposed Thesis Title</th>
</tr>
</thead>
<tbody>
<tr>
<td>Adelsman, M. A</td>
<td>Dr. N. J. Maihle</td>
<td>&quot;Ligand-Independent Dimerization of c-erbB1 Oncogenic v-erbB Products&quot;</td>
</tr>
<tr>
<td>Baines, J. E.</td>
<td>Dr. D. H. Persing</td>
<td>&quot;Novel Immunotherapeutic Approaches to Cervical Cancer&quot;</td>
</tr>
<tr>
<td>Calhoun, E. S.</td>
<td>Dr. D. F. Jelinek</td>
<td>Thesis project not determined</td>
</tr>
<tr>
<td>Canales, N. D.</td>
<td>Dr. S. J. Gendler</td>
<td>Thesis project not determined</td>
</tr>
<tr>
<td>Eley, G. D.</td>
<td>Dr. C. D. James</td>
<td>&quot;Characterization of the Epidermal Growth Factor Receptor Amplicon in Human Glioblastoma&quot;</td>
</tr>
<tr>
<td>Holmen, S. L.</td>
<td>Dr. M. J. Federspiel</td>
<td>&quot;Viral Receptors Engineered to Inhibit Viral Entry&quot;</td>
</tr>
<tr>
<td>Johnson, J. L.</td>
<td>Dr. N. J. Maihle</td>
<td>&quot;Mechanisms of c-erbB1 Oncogenic Signaling&quot;</td>
</tr>
<tr>
<td>Lomberk, G.</td>
<td>Dr. D. I. Smith</td>
<td>Thesis project not determined</td>
</tr>
<tr>
<td>Ritland, S. R.</td>
<td>Dr. S. J. Gendler</td>
<td>&quot;Genetic Linkage for Tumor Modifier Loci in the MMTV-neu Transgenic Mouse Mammary Tumor Model</td>
</tr>
<tr>
<td>Rogers, M. S.</td>
<td>Dr. E. E. Strehler</td>
<td>&quot;Studies on the Structure and Function of Human CLP in Breast Cancer&quot;</td>
</tr>
<tr>
<td>Schehl, C. M.</td>
<td>Dr. F. Couch</td>
<td>&quot;The Role of BRCA1/2 Mutations in Breast Cancer&quot;</td>
</tr>
<tr>
<td>Xu, K.</td>
<td>Dr. F. Prendergast</td>
<td>&quot;Molecular Studies of Farnesyl Transferase Inhibitors&quot;</td>
</tr>
<tr>
<td>Adriance, M.</td>
<td>Rotations</td>
<td>N.A.</td>
</tr>
<tr>
<td>Faupel, J.</td>
<td>Rotations</td>
<td>N.A.</td>
</tr>
<tr>
<td>Myers, S.</td>
<td>Rotations</td>
<td>N.A.</td>
</tr>
<tr>
<td>Walters, D</td>
<td>Rotations</td>
<td>N.A.</td>
</tr>
</tbody>
</table>
TABLE III. PUBLICATIONS OF TRAINEES

Adelsman, Margaret A.

**Papers**


Adriance, Melissa

**Abstracts**


Canales, Nohelia

**Abstracts**

Bower A, Serdoncelllo C, Canales N: The specificity of atrial natriuretic peptide on the release control of melanocyte stimulating hormone from the pituitary.

Canales N, Radice G: Histological analysis of mammary glands in P-cadherin deficient mice.

Bower A, Canales N, Becker K, Ocampo M: Atrial natriuretic peptide release inhibition by dopamine and melanocyte stimulating hormone.

Eley, Gregory

**Papers**


**Abstracts**

Holmen, Sheri L.

Papers


Abstracts


Johnson, Julie L.

Papers


Abstracts


Lomberk, Gwen

Papers


Ritland, Steve R.

Papers


Abstracts


Ritland S, Gendler S: Cancer chemoprevention studies using piroxicam in the ApcMin mouse. 5th International Conference on Eicosanoids and Other Bioactive Lipids in Cancer, Inflammation, and Related Diseases, La Jolla, California, September 1997.

Ritland S, Gendler S: Evaluation of 5-aminosalicylic acid for intestinal tumor chemoprevention in the ApcMin mouse. 5th International Conference on Eicosanoids and Other Bioactive Lipids in Cancer, Inflammation, and Related Diseases, La Jolla, California, September 1997.


Rogers, Michael S.

Papers


Abstracts


Schehl, Colleen M.

Papers


Abstracts

### TABLE IV. MEETINGS/COURSES ATTENDED BY TRAINEES

**Adelsman, Margaret**


**Canales, Noelia**


**Gregory Eley**


**Julie Johnson**

Experimental Pathology and Laboratory Medicine Symposia, 1997.

American Association for Cancer Research Annual Meeting, New Orleans, LA, March 28-April 1, 1998


**Lomberk, Gwen**


**Rogers, Michael**


American Association for Cancer Research Annual Meeting, New Orleans, LA, March 28-April 1, 1998

**Colleen Schehl**

THE FACULTY AND THEIR RESEARCH
Tumor Biology Program

Robert T. Abraham, Associate Professor; Ph.D., Pittsburgh, 1981. Signal transduction; cell-cycle regulation; leukemogenesis.
Margot P. Cleary, Visiting Scientist; Ph.D., Columbia, 1976. Breast cancer; obesity; nutrition.
Fergus J. Couch, Assistant Professor; Ph.D., University College Cork, Ireland, 1992. Identification and characterization of genes involved in familial and sporadic breast and ovarian cancer development. Functional analysis of the BRCA2 breast and ovarian cancer predisposition gene.
Chella S. David, Professor; Ph.D., Iowa State, 1966. Immunogenetic aspects of immune response, with emphasis on the major histocompatibility complex class II a genes and T-cell receptor gene.
Gordon W. Dewald, Professor; Ph.D., North Dakota, 1972. Cytogenetics and molecular cytogenetics of congenital disorders and hematologic malignancies.
Richard L. Ehman, Professor; M.D., Saskatchewan, 1979. Magnetic resonance imaging.
Charles Erlichman, Professor; M.D., Toronto (Canada), 1974. Pharmacology of drugs used in cancer therapy.
Mark J. Federspiel, Assistant Professor; Ph.D., Michigan State, 1987. Retroviral vectors; antiviral strategies; molecular medicine.
Lorraine A. Fitzpatrick, Professor; M.D., Chicago, 1980. Prostate cancer metastatic to bone; skeletal calcification; steroid regulation of metastatic disease.
*Sandra J. Gendler, Associate Professor; Ph.D., USC, 1984. Tumor cell biology; mucins in cancer and cystic fibrosis.
Michael J. Getz, Professor; Ph.D., Texas at Houston, 1972. Molecular biology of peptide growth factors; biology of tissue factor in tumorigenesis.
C. David James, Associate Professor; Ph.D., Wright State, 1986. Cancer genetics; cell cycle regulation.
Diane F. Jelinek, Assistant Professor; Ph.D., Texas Southwestern Medical Center, 1985. Cytokine-mediated signaling and gene expression in normal and malignant human B lymphocytes.
Larry M. Karnitz, Assistant Professor; Ph.D., Iowa, 1989. Signaling mechanisms of oncoproteins and hematopoietic growth factors; molecular radiobiology.
Scott H. Kaufmann, Associate Professor; M.D./Ph.D., Johns Hopkins, 1981. Pharmacology of topoisomerase-directed antineoplastic agents; apoptosis; resistance to anticancer drugs.
Paul J. Leibson, Associate Professor; Ph.D., 1981, M.D., 1979, Chicago. Tumor immunology; lymphocyte activation; antiviral immunity.
Vanda A. Lennon, Professor; M.B.B.S., Sydney (Australia), 1966; Ph.D., Melbourne (Australia), 1973. Immunobiology of autoimmunity and cancer; ionic channel protein antigens in human neoplasms of lung, ovary, and breast (carcinomas), and thymic epithelium (thymoma).
Edward B. Leof, Associate Professor; Ph.D., North Carolina, 1982. Regulation of cellular proliferation; genetics of pneumocystis carinii.
Ricardo V. Lloyd, Professor; M.D./Ph.D., Wisconsin, Madison, 1975. Endocrine tumor biology, especially pituitary and thyroid.
John A. Lust, Assistant Professor; M.D./Ph.D., Boston University, 1983. Role of IL-6 and IL-6R in pathogenesis of multiple myeloma; detection of minimal residual disease in myeloma transplant patients by PCR.

L. James Maher, Associate Professor; Ph.D., Wisconsin, 1988. Nucleic acid biochemistry; triple helix DNA.

Nita J. Mahle, Associate Professor; Ph.D., Yeshiva (Einstein), 1983. Molecular basis of cancer; human breast, ovarian, and prostate carcinomas; gliomas.

David J. McKeen, Professor; Ph.D., Johns Hopkins, 1972. Signaling and gene transcription events in T helper lymphocytes; MHC class II protein transport.

Michael J. McManus, Assistant Professor; M.D., Georgetown, 1983. Molecular pediatric oncology; growth factor receptors; tyrosine kinase signal transduction pathways.

Mark A. McNiven, Associate Professor; Ph.D., Maryland, 1987. Cytoskeletal dynamics in mammalian cells; molecular basis of cellular migration during metastasis; vesicle-based transport in epithelial cells.

L. Joseph Melton, III, Professor; M.D., LSU, 1969. Chronic disease epidemiology.

Heidi Nelson, Associate Professor; M.D., Washington (Seattle), 1981. Colorectal cancer; immunotherapy.


Dennis J. O’Kane, Assistant Professor; Ph.D., SUNY at Stony Brook, 1979. Telomerase activity as a diagnostic marker for cancer; translational research on new tumor markers.

David H. Persing, Associate Professor; M.D./Ph.D., California, San Francisco, 1988. Precore promoter mutations in hepatic tumors; immunogenetic determinants of chronic papillomavirus infections and cervical cancer; association of chronic infections with lymphoproliferation.

Mark R. Pittelkow, Professor; M.D., Mayo, 1979. EGF-related growth factor/receptor function: epidermal keratinocyte and melanocyte regulation of growth and differentiation.

Karl C. Podratz, Professor; M.D./Ph.D., St. Louis, 1974. Molecular prognostic determinants in gynecologic malignancies.

Gregory A. Poland, Professor; M.D., Southern Illinois. Expertise in vaccine development and evaluation, adjuvant development and evaluation; vaccine antigen processing and HLA presentation; and vaccine immunogenetics.

Franklyn G. Prendergast, Professor; M.B.B.S., West Indies, 1968; Ph.D., Minnesota, 1977. Fluorescence spectroscopy; protein structure and dynamics; biochemistry and bioluminescence.

Corey Raffel, Associate Professor; M.D./Ph.D., California, San Diego, 1980. Pediatric neuro- oncology; gene therapy and cancer.

Jeffrey L. Salisbury, Professor; Ph.D., Ohio State, 1978. Cell cycle control; centrosomes; mitotic spindle poles; breast cancer.

David I. Smith, Professor; Ph.D., Wisconsin, 1978. Chromosomal fragile sites; molecular genetics of cancer development.

Thomas C. Spelsberg, Professor; Ph.D., West Virginia, 1967. Steroid action on early (c-myc) gene transcription, steroids and TGF-β action on bone cell functions, and early gene expression.

Emanuel E. Strehler, Associate Professor; Ph.D., ETH Zurich (Switzerland), 1981. Intracellular Ca^{2+} homeostasis and signaling; molecular mechanisms of disease.

Stephen N. Thibodeau, Professor; Ph.D., Washington (Seattle), 1979. Cancer genetics; colon and prostate cancer.


Raul Urrutia, Assistant Professor; M.D., Cordoba (Argentina), 1987. Cell differentiation.

Peter J. Wettstein, Professor; Ph.D., North Carolina at Chapel Hill, 1977. Role of minor histocompatibility antigens in allograft rejection.
Lester E. Wold, Professor; M.D., Chicago, 1977. Immunocytochemistry; bone tumors and tumor-like conditions; breast diseases.
Charles Y-F. Young, Assistant Professor; Ph.D., Brigham Young, 1984. Calpain inhibitor-induced apoptosis in human prostate adenocarcinoma cells.

*Scottsdale campus.
# Tumor Biology I:

**Introduction to Tissue and Tumor Biology (TBIO 5000)**

2:30 - 4:00 p.m. Tue. Thur. Fall Quarter 1998

[50% participation, 25% term paper, 25% lab/problem sets]

<table>
<thead>
<tr>
<th>Date</th>
<th>Topic</th>
<th>Instructor</th>
</tr>
</thead>
<tbody>
<tr>
<td>September 29</td>
<td>Principles of Cell and Tissue Design</td>
<td>Salisbury</td>
</tr>
<tr>
<td>September 30</td>
<td>TBJC: Fundamentals of the Cell Cycle</td>
<td></td>
</tr>
<tr>
<td>October 1</td>
<td>Laboratory - Light and Electron Microscopy</td>
<td></td>
</tr>
<tr>
<td>October 6</td>
<td>Stem Cells, Differentiation, and Cancer</td>
<td>Maihle</td>
</tr>
<tr>
<td>October 7</td>
<td>TBJC: Genomic Instability/Aneuploidy</td>
<td></td>
</tr>
<tr>
<td>October 8</td>
<td>Guest Lecture - Epigenetics and Genetics</td>
<td></td>
</tr>
<tr>
<td>October 13</td>
<td>Properties of Transformed Cells <em>in vitro</em></td>
<td>Maihle</td>
</tr>
<tr>
<td>October 14</td>
<td>TBJC: Temin - Hayflick</td>
<td></td>
</tr>
<tr>
<td>October 15</td>
<td>Senescence and Immortalization</td>
<td></td>
</tr>
<tr>
<td>October 20</td>
<td>Properties of Transformed Cells <em>in vivo</em></td>
<td>Maihle</td>
</tr>
<tr>
<td>October 21</td>
<td>TBJC: Transgenics and Knockouts</td>
<td></td>
</tr>
<tr>
<td>October 22</td>
<td>Xenografts</td>
<td></td>
</tr>
<tr>
<td>October 27</td>
<td>Tissue Biology - Epithelia</td>
<td>Salisbury</td>
</tr>
<tr>
<td>October 28</td>
<td>TBJC: Cell Polarity</td>
<td></td>
</tr>
<tr>
<td>October 29</td>
<td>Laboratory - Epithelia [TERM PAPER OUTLINE DUE by 4:00 p.m.]</td>
<td></td>
</tr>
<tr>
<td>November 3</td>
<td>Tissue Biology - Connective Tissue</td>
<td>Salisbury</td>
</tr>
<tr>
<td>November 4</td>
<td>TBJC: Invasion and Metastasis</td>
<td></td>
</tr>
<tr>
<td>November 5</td>
<td>Laboratory - Connective Tissue</td>
<td></td>
</tr>
<tr>
<td>November 10</td>
<td>Endothelial Cells, Vascular Tissue and Lymphatics</td>
<td>Salisbury</td>
</tr>
<tr>
<td>November 11</td>
<td>TBJC: Angiogenesis</td>
<td></td>
</tr>
<tr>
<td>November 12</td>
<td>Laboratory - Vascular Tissue</td>
<td></td>
</tr>
<tr>
<td>November 17</td>
<td>Pathobiology of Cancer</td>
<td>Salisbury</td>
</tr>
<tr>
<td>November 18</td>
<td>TBJC: Epithelial / Mesenchymal Interactions</td>
<td></td>
</tr>
<tr>
<td>November 19</td>
<td>Surgical Pathology Tours</td>
<td></td>
</tr>
<tr>
<td>November 24</td>
<td>INDEPENDENT STUDY</td>
<td></td>
</tr>
<tr>
<td>November 25</td>
<td>INDEPENDENT STUDY</td>
<td></td>
</tr>
<tr>
<td>November 26</td>
<td>Thanksgiving</td>
<td></td>
</tr>
<tr>
<td>December 1</td>
<td>Normal Breast / Breast Cancer</td>
<td>Salisbury</td>
</tr>
<tr>
<td>December 2</td>
<td>TBJC: Breast Tumor Staging and Grade</td>
<td></td>
</tr>
<tr>
<td>December 3</td>
<td>Laboratory - Breast Pathology</td>
<td></td>
</tr>
<tr>
<td>December 8</td>
<td>Normal Intestine / Colon Cancer</td>
<td>TBA</td>
</tr>
<tr>
<td>December 9</td>
<td>TBJC: HNPCC - MIN Mice</td>
<td></td>
</tr>
<tr>
<td>December 10</td>
<td>Laboratory - GI Tumors</td>
<td></td>
</tr>
<tr>
<td>December 17</td>
<td>[TERM PAPER EVALUATIONS DUE by 4:00 p.m.]</td>
<td></td>
</tr>
</tbody>
</table>
Tumor Biology II:
Origins of Human Cancer (TBio 8000)
2:30 - 4:00 p.m. Tue. Thur. Winter Quarter 1998
[50% participation/50% term paper]

January 7  Origins of Human Cancer: An Overview  Maihle
January 8  Tumor Biology Journal Club:
January 9  Problem Set (Maihle)
January 14 Origins of Human Cancer: Etiology and Genetics  Smith
January 15 Tumor Biology Journal Club:
January 16 Problem Set
January 21 Origins of Human Cancer: Progression and Metastasis  Gendler
January 22 Tumor Biology and Journal Club: Angiogenesis
January 23 Problem Set (Gendler and Maihle)
January 28 Origins of Human Cancer: Epidemiology and Prevention  Yang
January 29 Tumor Biology Journal Club: Intestinal Polyposis and COX-2
January 30 Tumor Immunology: An Overview  Mitchell
February 4  Problem Set (Epidemiology and Prevention)
February 5  Tumor Biology Journal Club (Tumor Immunology)  Jelinek
February 6  Problem Set (Tumor Immunology)
February 11 Paraneoplastic Syndromes (in Breast and Ovarian Cancer)  Lennon
February 12 Tumor Biology Journal Club: Paraneoplastic Autoimmunity
February 13 Problem Set
February 25 Introduction to Clinical Research  O’Fallon
February 26 Tumor Biology Journal Club: Phase I Trial of Dolastatin-10
February 27 Problem Set
March 4  Introduction to Chemotherapy  Ames
March 5  Tumor Biology Journal Club: Inhibitors of Farnesyl Transferase
March 6  Problem Set
March 11 Tumor Imaging: An Overview  Robb
March 12 Experimental Tumor Imaging  Ehman
March 13 Problem Set (Maihle)
March 18 Introduction to Surgical Oncology  Nelson
March 19 Tumor Biology Journal Club: Surgical Procedures in Colon Cancer
March 25 Introduction to Radiation Therapy  Bonner
March 26 Breast Cancer Patient Vignettes
March 27 Experimental Gene Therapy  Raffel

Biology of Breast Cancer Training Program 30
Tumor Biology III
Growth Factors, Oncogenes, and Tumor Suppressors (TBIO 8005)
2:30 - 4:00 p.m. Tue. Thur. Winter Quarter 1998
[50% participation/50% term paper]

April 7  Cell Cycle and Cell Growth Control  Salisbury
April 8  Tumor Biology Journal Club
April 9  Regulation of Immediate Early Gene Expression
         (AACR Meeting 3/28 - 4/1)  Getz
April 14  Growth Factors/GF Receptors  Maihle
April 15  Tumor Biology Journal Club
April 16  Student Discussion Problem Set
April 21  Intracellular Mediators: Kinases and Phosphatases  Maihle
April 22  Tumor Biology Journal Club
April 23  Student Discussion Problem Set
April 28  Intracellular Mediators: G Proteins  Karnitz
April 29  Tumor Biology Journal Club
April 30  Student Discussion Problem Set
May 5  Oncogenes and Viral Oncogenes  Maihle
May 6  Tumor Biology Journal Club
May 8  Student Discussion Problem Set
May 12-14  INDEPENDENT STUDY
May 19  Introduction to Tumor Suppressors  James
May 20  Tumor Biology Journal Club
May 21  Student Discussion Problem Set
May 26  Cancer Genetics  Jenkins/Lloyd
May 27  Tumor Biology Journal Club
May 28  Student Discussion Problem Set
June 2  P53 - Guardian of the Genome  James
June 3  Tumor Biology Journal Club
June 4  Student Problem Set
June 9  Retinoblastoma and Rb  Smith
June 10  Tumor Biology Journal Club
June 11  Student Discussion Problem Set
June 16  To Die Or Not To Die - Apoptosis  TBA
June 17  Tumor Biology Journal Club
June 18  Student Discussion Problem Set
Biology of Breast Cancer

(TBio1 5200)
Guggenheim 1093
1:30-2:30 p.m. Fridays

(50% participation/50% final exam)

- This course is aimed at integrating basic concepts in developmental, cellular and molecular biology of the breast together with current information on the etiology, diagnosis and treatment of breast cancer.
- The faculty include members form diverse basic science and medical disciplines including cell and molecular biology, pathology, oncology and surgery.

   April 11  Breast Cancer: The Magnitude of the Problem                     Ingle
   April 18  Development, Anatomy and Histology and Cell Biology of the Breast  Salisbury
   April 25  Histopathology of the Breast                                  Wold
   May 2     Experimental Models of Breast Cancer                          Gendler
   May 9     Oncogenes, Growth Factors, and Breast Cancer                  Leof
   May 16    Tumor Suppressors and Breast Cancer                          Mr. Ritland
   May 23    Radiation Therapy for Breast Cancer                          Peterson
   May 30    Surgical Treatment of Breast Cancer                          Donohue
   June 6    Breast Cancer Diagnosis and Imaging                          Johnson
   June 13   Systemic Therapy for Breast Cancer                           Ingle
   June 20   Experimental Therapies for Breast Cancer                     Maihle
   June 27   Final Examinations Due (by 5:30 p.m.)
Principles in Pancreatic Cancer

(TBiol 5200)
Guggenheim 1093
3:00-4:00 p.m. Fridays

- This course is aimed at integrating basic concepts in developmental, cellular and molecular biology of the pancreas together with current information on the etiology, diagnosis and treatment of pancreatic cancer.
- The faculty include members from diverse basic science and medical disciplines including cell and molecular biology, pathology, clinical gastroenterology, oncology and surgery.

October 25  Pancreatic Cancer: What is the Problem?
             Dr. Eugene P. DiMagno, Gastroenterology Research Unit, SMH

November 1  Development, Cell Biology and Histology of the Pancreas
             Dr. Raul Urrutia, Gastroenterology Research Unit, SMH

November 8  Histopathology of Pancreatic Cancer
             Dr. Lawrence J. Burgart, Surgical Pathology

November 15 Experimental Models of Pancreatic Cancer
              Dr. Raul Urrutia, Gastroenterology Research Unit, SMH

November 22 Cellular and Molecular Mechanisms of Pancreatic Cancer
              Dr. Raul Urrutia, Gastroenterology Research Unit, SMH

December 6  Current and Future Non-Surgical Treatments of Pancreatic Cancer
              Dr. Richard M. Goldberg, Medical Oncology

December 13 Surgical Treatment of Pancreatic Cancer
              Dr. Michael Sarr, Gastroenterology Research Unit and Surgery, SMH

For more information contact
Dr. Raul Urrutia (4-7500)
Business of Science, Science of Business

TBio 5300
K. E. Bennet, M.B.A. and N. J. Maihle, Ph.D.

Summer Quarter (even years)

1) Introduction (August 2) Orientation and Objectives [KEB/NJM]
2) Administrative Structures in Support of Research (August 5) [KEB]
   Not-for-Profit & For Profit
3) Overview of Research Accounting (August 7) [KEB]
   Research Budgets
   Direct versus Indirect costs
4) Sources of Financial Support for Research (August 9) [NJM]
   Intramural & Extramural Support
   Federal & Private
5) Sources of Financial Support for Research (August 12) [KEB]
   Corporate
   Strategic Alliance, Joint Development
   Licensing/Venture Capital
6) Introduction to Intellectual Property (August 14) [KEB]
   Definition of Intellectual Property
   Protection of Intellectual Property
   Patents & Trade Secrets
   Ownership of Intellectual Property
7) Introduction of Cases (August 16) [KEB]
8) Commercialization of Research Discoveries (August 21) [KEB]
   Licensing
   Market Value of Invention
9) Independent Study on Cases (August 19)
10) Laws and Policies Governing Conduct of Research (August 23)
    Institutional [NJM]
    State, Federal, and International [KEB]
11) Case Presentations "Levamisole" (August 26) [KEB/NJM]
12) Case Presentations - "University of Florida" (August 28) [KEB/NJM]
13) Course Wrap-Up (August 30) [KEB/NJM]
Origins of Human Cancer: Normal Breast & Breast Cancer
1/13/98 TBIO II 8000

Tumor Biology II (8000)
Origins of Human Cancer

Normal Breast Development and Histology, and Breast Cancer
January 13, 1998
Jeffrey L. Salisbury, Ph.D.

Normal Breast
- compound, branched, alveolar gland
- 15-25 irregular lobes
- terminal duct lobular unit (lobules: functional secretory unit in lactation)
  - ducts: lactiferous ducts, lactiferous sinu (ampulla), extralobular ducts, lobular ducts, alveolar ducts (terminal ductules)
- alveoli
  - lining epithelium
  - surrounding myoepithelial cell layer (oxytocin responsive)
- adipose and connective tissue

Mammary Gland Development
- four developmental stages
  - embryonic
  - adolescent
  - lactating
  - involution

Embryonic Stage
- 'primitive milk streak', mammary ridge or milk line
  - thickened epithelial layer derived from ectoderm
- mammary ridge or milk line
  - raised epidermal tissue 4-6 cell layers both sides of ventral midline, other portions invovlute
  - mammary bud (disc, globular, cone = bud)
    - centers of cellular migration and shape changes (mouse 5-6 per side, human only 1 per side)
  - 15-25 secondary buds = mammary cord
    - rapid proliferation = cord opens at skin (nipple)
    - proliferation at opposite end = branching ducts
- development ceases until puberty

Lecture Notes page 1

J.L. Salisbury, Ph.D.
salisbury@mayo.edu
Origins of Human Cancer: Normal Breast & Breast Cancer

Breast Development in the Male
- identical to female until day 13-15 gestation
- mesenchyme condenses around center of mammary bud, and cells of the cord die
  - small segment of the cord detaches from superficial epidermis
    - mammary rudiment (no further development)
- + testosterone ⇒ cell death, bud degeneration
- - testosterone ⇒ male mammary development

Mammary System of the Mouse and Milk Line in Human Fetus and Corresponding Location in Adult

Mammary rudiment in male mouse fetus. The rudiment has separated from the epidermis.

3H-Dihydrotestosterone Receptor Radioautography
(A) Median section through a mammary rudiment. Epidermis at top. Steroid-binding mesenchymal cells are found only around the gland bud and its stalk.
(B) Oblique section through an epithelial gland bud. There is a well-defined envelope of receptor-positive mesenchymal cells around the epithelium, indicating the short range of inductive influence.

Epithelial-Mesenchyme Interactions in the Mouse:
- mesenchyme induces epithelial mammary edge to proliferate
- young bud induces fibroelastic 'mammary' mesenchyme (wandering & smooth muscle)
- day 14 male, testosterone
  - mesenchyme to destroy epithelial anlage
- day 16-17 female, 1' sprout invades fat pad; further growth and branching

Lecture Notes page 2

J.L. Salisbury, Ph.D.
salisbury@mayo.edu
Origins of Human Cancer: Normal Breast & Breast Cancer
1/13/98 TBIO II 8000

Hormonal Regulation of Growth & Function
- puberty
  - estrogen (ovary) ⇒ growth of duct system
  - progesterone (ovary) ⇒ alveoli development
- pregnancy
  - estrogen (ovary & placenta) ⇒ growth of duct system
  - progesterone (ovary & placenta) ⇒ alveoli development
  - prolactin (pars distalis of hypophysis) ⇒ full glandular devel.
- birth
  - estrogen →, progesterone →, prolactin ⇒ lactation
- regression
  - nursing, prolactin → lactation, epithelium degenerates, connective tissue →

Role of Growth Factors
- TGFβ family
  - generally inhibitory for breast epithelial growth
    - TGFβ1 inhibits ductal growth by DNA S in end buds
    - not in stromal cells or proliferation of lobuloalveolar structures
    - TGFβ 2 & 3 ⇒ disappearance of proliferating mammary stem cell layer, involution of ductal end buds, & cessation of glandular growth
    - TGFβ synthesis and secretion of milk proteins and limit their accumulation during pregnancy
  - TGFβ over expression in transgenic mice
    ⇒ general mammary hyperplasia
    ⇒ reduced mammary ductal branching
    ⇒ failure of lobuloalveolar development

Role of Growth Factors
- TGFα and EGF (autocrine & paracrine)
  - mitogens for normal and malignant epithelial cells
  - over expression in transgenic mice
    ⇒ mammary hyperplasia
    ⇒ increased incidence of carcinoma
    ⇒ shorter latency following chemical carcinogenesis
- KGF keratinocyte growth factor (FGF family)
  - secreted by stromal cells
    ⇒ stimulate epithelial cell proliferation
    ⇒ ductal & acinar cell growth

in situ hybridization for KGFR and KGF in mammary rudiments

Nipple and Areola
- accessory areolar glands of Montgomery
  intermediate between sweat glands and true mammary glands
- sebaceous glands (usually lacking hair follicles)
- sweat glands
- smooth muscle (inner, intermediate, outer layers)
- connective tissue (circular, elliptic course)
- richly innervated (Meissner's corpuscles, Merkel's discs, Krause's end bulbs, Pacinian corpuscles)

Whole mount of cleared mouse mammary fat pad week 4 to 6 illustrating terminal end buds. Hematoxylin stain.

Lecture Notes page 3

J.L. Salisbury, Ph.D.
salisbury@mayo.edu
Origins of Human Cancer: Normal Breast & Breast Cancer
1/13/98 TBIO II 8000

Inactive Breast

- parenchyma (sparsely consists mainly of duct elements)
  - cuboidal epithelial cells and myoepithelial cells
- stroma (abundant)
  - loose irregular cellular connective tissue immediately surrounding ducts
  - dense irregular connective tissue beyond the area of the lobule
  - fat

SEM of an Acinus of the Mammary Gland

Branching myoepithelial cells occupy the grooves between the bases of the secretory cells.

From Nagas, T. Cell Tissues Rep 200:1, 1983

Lactating Mammary Gland

- parenchyma (abundant and consists mainly of alveolar elements)
  - cuboidal epithelial cells
  - myoepithelial cells
  - product within lumen of alveoli and ducts
- stroma (reduced)
  - small amount of connective tissue between alveoli and septa between between lobules
- lymphocytes often present

Acinus of a Lactating Mammary Gland

Identify:
A) lipid droplets
B) casein granules
C) secretory cells
D) myoepithelial cells
E) capillary

Secretion in Breast Alveolar Cells

- merocrine (exocytic)
  - milk proteins casein, lactalbumin
- apocrine
  - loss of portion of cell membrane and some cytoplasm
  - lipid droplets
- diffusion
  - water, ions
- transcytosis
  - IgA

Breast Cancer

- Incidence:
  - 180,000 new cases per year
  - 1:8 women in US and Canada (lifetime risk)
  - rare before age 20, rarely diagnosed in < age 25
  - past age 25 incidence rises steadily to peak around menopause
- Deaths
  - 45,000 deaths annually (US)
  - second most common cause of cancer death

Lecture Notes page 4

J.L. Salisbury, Ph.D.
salisbury@mayo.edu
Origins of Human Cancer: Normal Breast & Breast Cancer
1/13/98 TBIO II 8000

Risk Factors
- High (relative risk > 4.0)
  - age
  - personal history BC
  - family history
  - proliferative disease w/ atypia
- Moderate (relative risk 2-4)
  - first degree relative w/ BC
  - personal history ovarian, or endometrial cancer
  - age 1st full term pregnancy ≥30
  - multiparous
  - obesity, postmenopausal
  - upper socioeconomic class
- Low (relative risk < 1-2)
  - menarche before age 12
  - menopause after age 55
  - Caucasian race
  - moderate alcohol intake
  - long duration estrogen replacement therapy (>15 years)

Invasive Carcinomas of Breast Cancer

<table>
<thead>
<tr>
<th>Histological Type</th>
<th>Frequency</th>
<th>5-year survival</th>
</tr>
</thead>
<tbody>
<tr>
<td>Infiltrating ductal carcinoma</td>
<td>43.6%</td>
<td>79</td>
</tr>
<tr>
<td>Infiltrating lobular carcinoma</td>
<td>5.9%</td>
<td>84</td>
</tr>
<tr>
<td>Infiltrating ductal &amp; lobular</td>
<td>1.6%</td>
<td>95</td>
</tr>
<tr>
<td>Medullary carcinoma</td>
<td>2.8%</td>
<td>87</td>
</tr>
<tr>
<td>Mucinous (colloid) carcinoma</td>
<td>2.1%</td>
<td>95</td>
</tr>
<tr>
<td>Comedocarcinoma</td>
<td>1.4%</td>
<td>87</td>
</tr>
<tr>
<td>Paget's Disease</td>
<td>1.0%</td>
<td>79</td>
</tr>
<tr>
<td>Papillary Carcinoma</td>
<td>0.8%</td>
<td>96</td>
</tr>
<tr>
<td>Tubular carcinoma</td>
<td>0.6%</td>
<td>96</td>
</tr>
<tr>
<td>Adenocarcinoma, NOS</td>
<td>7.5%</td>
<td>65</td>
</tr>
<tr>
<td>Carcinoma, NOS</td>
<td>3.5%</td>
<td>62</td>
</tr>
</tbody>
</table>

Noninvasive Carcinomas of Breast Cancer

<table>
<thead>
<tr>
<th>Histological Type</th>
<th>Frequency</th>
<th>5-year survival</th>
</tr>
</thead>
<tbody>
<tr>
<td>Intraductal carcinoma (DCIS)</td>
<td>3.6%</td>
<td>&gt;99</td>
</tr>
<tr>
<td>Lobular carcinoma in situ (LCIS)</td>
<td>1.6%</td>
<td>&gt;99</td>
</tr>
<tr>
<td>Intralobular &amp; LCIS</td>
<td>0.2%</td>
<td>&gt;99</td>
</tr>
<tr>
<td>Papillary carcinoma</td>
<td>0.4%</td>
<td>&gt;99</td>
</tr>
<tr>
<td>Comedocarcinoma</td>
<td>0.3%</td>
<td>&gt;99</td>
</tr>
</tbody>
</table>

Hormone Receptor Status
- cancers which display estrogen receptors have a better prognosis (ER+)
  - differentiated
  - respond to hormonal manipulation
  - tamoxifen (anti-estrogen) treatment ~75% ER+ patients respond
- significance of progesterone receptor (PR) status less well understood
  - generally ER+ are also PR+
  - cancers that are PR+ and ER- have a worse prognosis

Additional Markers
- Cathepsin D (acidic lysosomal protease)
  - correlation with metastasis
- C-erb B-2 (C-neu)
  - correlation with high nuclear grade and aneuploidy

Flow Cytometry

Lecture Notes page 5

J.L. Salisbury, Ph.D.
salisbury@mayo.edu
Origins of Human Cancer: Normal Breast & Breast Cancer
1/13/98 TBIO II 8000

Flow Cytometry

- diploid normal cell population
- aneuploid tumor cells in general have a poorer prognosis
- Flow Cytometry is a rapid method for determining the degree of aneuploidy in a tumor sample

Aneuploidy and low S phase

Diagnostic Procedures

- self-examination
  - regular basis to follow normal breast architecture
  - 1 cm tumor (detectable) 5-10 years
- mammography
  - most sensitive and specific
  - location of cancers

(Histology) Grading (modified Scarff-Bloom-Richardson)

<table>
<thead>
<tr>
<th>Tubule Formation (% tubules composed of tubular structures)</th>
<th>Score</th>
</tr>
</thead>
<tbody>
<tr>
<td>&gt; 75</td>
<td>1</td>
</tr>
<tr>
<td>10-75%</td>
<td>2</td>
</tr>
<tr>
<td>&lt; 10%</td>
<td>3</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Nuclear Ploymorphism</th>
<th>Score</th>
</tr>
</thead>
<tbody>
<tr>
<td>small, uniform shape and staining</td>
<td>1</td>
</tr>
<tr>
<td>moderate increase in size and variation</td>
<td>2</td>
</tr>
<tr>
<td>marked variation</td>
<td>3</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Mitotic Count (per 10 high power fields)</th>
<th>Score</th>
</tr>
</thead>
<tbody>
<tr>
<td>up to 7</td>
<td>1</td>
</tr>
<tr>
<td>7 to 14</td>
<td>2</td>
</tr>
<tr>
<td>15 or more</td>
<td>3</td>
</tr>
</tbody>
</table>

Grade calculated by adding the scores. Grade correlates with survival.

<table>
<thead>
<tr>
<th>Grade</th>
<th>Score</th>
<th>5-year survival (%)</th>
<th>7-year survival (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>3 to 5</td>
<td>95</td>
<td>90</td>
</tr>
<tr>
<td>2</td>
<td>6 or 7</td>
<td>75</td>
<td>65</td>
</tr>
<tr>
<td>3</td>
<td>8 or 9</td>
<td>50</td>
<td>45</td>
</tr>
</tbody>
</table>

Treatment

- Primary or Local-Regional Therapy
  - Breast Conservation Surgery
    (lumpectomy + axillary dissection + XRT)
- Systemic Therapy
  - Hormonal Therapy
    Tamoxifen (nonsteroidal anti-estrogen and estrogen-like activity)
    Megace (progestational agent)
    Halotestin (an androgen)
- Chemotherapy
  - Adriamycin (A), cyclophosphamide (C), methotrexate (M), 5-fluorouracil (F), vincristine/vinblastine, mitomycin, cisplatin
  - most common combination CMF, CAP, or AC

Stage in Breast Cancer is Based on Size and Degree of Spread.

<table>
<thead>
<tr>
<th>Stage</th>
<th>Definition</th>
<th>Survival (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>I</td>
<td>Tumor ≤ 2 cm, N0, M0</td>
<td>96 72</td>
</tr>
<tr>
<td>II</td>
<td>Tumor &gt;2≤5 cm, N1,M0</td>
<td>81 71</td>
</tr>
<tr>
<td></td>
<td>or &gt;5, N0,M0</td>
<td></td>
</tr>
<tr>
<td>III</td>
<td>Tumor any size,N2, M0</td>
<td>52 39</td>
</tr>
<tr>
<td>IV</td>
<td>Tumor any size, N0-3, M1</td>
<td>18 11</td>
</tr>
</tbody>
</table>

Lecture Notes page 6

J.L. Salisbury, Ph.D.
salisbury@mayo.edu
Appendix G

Current Topics in Tumor Biology

(TBiol 5151)
2:30 - 3:30 p.m. Wednesdays

Journal Club Topics and (Presenter) Academic Year 1997-1998


(Ms. Nohelia Canales)


(Ms. Gwen Lomberk)

(Ms. Colleen Schehl)

(Ms. Julie Johnson)

(Mr. Andy Danielsen)

(Mr. Kun Xu)

(Ms. Gwen Lomberk)

(Mr. Eric Calhoun)
(Mr. Jonathan Hoyne)

(Mr. Kurt Krummel)

3/25 p53-Dependent Apoptosis Modulates the Cytotoxicity of Anticancer Drugs.
Uncoupling of S Phase and Mitosis Induced by Anticancer Agents in Cells Lacking p21.
(Ms. April Blajeski)

(Dr. Jill Reiter)

4/15 Trans receptor inhibition of human glioblastoma cells by erbB family ectodomains. D.M.
The enhanced tumorigenic activity of a mutant epidermal growth factor receptor common in
human cancers is mediated by threshold levels of constitutive tyrosine phosphorylation and
(Mr. Jonathan Hoyne)

4/22 Matrix adhesion and Ras transformation both activate a phosphoinositide 3-OH kinase and
16:2783-2793.
(Mr. Kun Xu)

4/29 A serine/threonine kinase gene defective in Peutz-Jeghers syndrome. A. Hemminki et al.,
Cloning and characterization of a novel serine/threonine protein kinase expressed in early
(Ms. Susan Barrett)

5/6 Forced degradation of Fas inhibits apoptosis in adenovirus-infected cells. A.E. Tollefson
(Dr. Cecelia Boardman)

5/20 Serial Analysis of Gene Expression. V. Velculescu, L. Zhang, B. Vogelstein, and K.
Gene Expression Profiles in Normal and Cancer Cells. L. Zhang, W. Zhou, V.
Science 276:1268-1272.
(Mr. Eric Calhoun)
Faculty Research Interests

Advanced courses, tutorials, seminars and journal clubs provide the depth of knowledge you will require to become an expert in your chosen field of study. Because our student-to-faculty ratio is low, classes frequently use small-group, interactive tutorial settings.

Listed below are brief descriptions of each of the areas of specialization offered by Mayo Graduate School.

Biochemistry and Molecular Biology

This program has 23 primary appointees and 29 faculty (representing 16 different departments and divisions) with secondary appointments. Students can choose from a diverse array of research opportunities including the regulation of gene expression and cell growth, steroid hormone action, DNA replication, oncogenesis, human molecular genetics and molecular biophysics.

The faculty contact person is:

Frank M. Rusnak, Ph.D.
Department of Biochemistry and Molecular Biology
507-284-2289
rusnak@mayo.edu

Biomedical Imaging

This program's 25-member faculty and their laboratories offer training in the methods of biomedical imaging science and clinical imaging investigations. Particularly strong components include computer visualization, confocal microscopy, computed tomography (including dynamic), magnetic resonance imaging (including functional), ultrasound imaging and virtual reality.

The faculty contact person is:

Richard A. Robb, Ph.D.
Department of Physiology and Biophysics
507-284-2997
rar@mayo.edu

Immunology

A faculty of 17 investigators provides students with access to this rapidly growing field. Research opportunities include allergy, cellular immunology, immunogenetics, immunopharmacology, molecular biology, molecular immunology, immunochemistry, neuroimmunology, tumor immunology, autoimmunity, immunoparasitology and rheumatology.

The faculty contact person is:

Larry R. Pease, Ph.D.
Department of Immunology
507-284-9891
pease@mayo.edu
Molecular Neuroscience

This inter-departmental program has a 35-member faculty and 30 independent laboratories that focus on the neurobiological processes related to human disease. Strengths include molecular biology, membrane and channel biophysics, signal transduction, receptor pharmacology, neural networks, three-dimensional imaging and neural cell biology.

The faculty contact person is:

Anthony J. Windebank, M.D.
Molecular Neuroscience Program
507-284-8729
windebank.anthony@mayo.edu

Pharmacology

A faculty of 16 scientists and research laboratories associated with this program focus on the biochemical and physiological processes that underlie the action of drugs. Current areas of investigation include intracellular transduction mechanisms, pharmacogenetics, molecular neuropharmacology, cancer chemotherapy, receptor biology, cardiovascular pharmacology, muscle contraction, immunopharmacology, pharmacokinetics, mass spectrometry, drug metabolism and membrane function.

The faculty contact person is:

Cynthia T. McMurray, Ph.D.
Department of Pharmacology
507-284-2747
mcmurray@mayo.edu

Physiology and Biophysics

A 22-member faculty leads a variety of investigative projects in cellular and systemic physiology. The program emphasizes the study of ion channels, fast microscopic imaging of cells, smooth muscle physiology, gastrointestinal physiology, cardiovascular and pulmonary physiology, renal physiology and hypertension.

The faculty contact person is:

Richard A. Robb, Ph.D.
Department of Physiology and Biophysics
507-284-2997
rar@mayo.edu

The Biology of Breast Cancer and Tumor Biology

This integrated, multi-disciplinary program offers specialized training in the biology of cancer, especially women's cancers. The 32-member faculty have research and clinical appointments in a broad range of medical and research specialties. The program's research strengths include gene regulation, cell cycle control, oncogene and tumor suppressor action, tumor immunology, signal transduction, anti-tumor pharmacology, and the biology of breast, ovarian, uterine, lung, gastrointestinal and prostate cancers.

Supported in part by a grant from the USAMRMC Breast Cancer Research Program: DAMD 17-94-J-4116.

The faculty contact person is:

Jeffrey L. Salisbury, Ph.D.
Department of Biochemistry and Molecular Biology
507-284-4070
salisbury@mayo.edu

[Next...M.D.-Ph.D. Program]
UNIVERSITY OF MINNESOTA

Medical School
Assistant Professor in Cell Biology

The Department of Cell Biology and Neuroanatomy seeks applications for a tenure track assistant professor position. All areas of cell biology will be considered, but particular consideration will be given to research directed at signal transduction mechanisms. Candidates will be expected to develop a strong, independent research program that leads to external funding. Applicants must possess a Ph.D. or equivalent degree and have at least two years of postdoctoral experience. Candidates must be U.S. citizens or be able to secure permanent resident status and must provide verification of highest degree. Strong candidates will have significant publications in high quality peer-reviewed journals and will be engaged in research that complements the departmental strengths in cell biology, developmental biology and neuroscience. A demonstrated ability to interact and collaborate will be favored. The teaching assignment will be in appropriate departmental courses that are offered to professional students, undergraduates, or graduate students in several interdepartmental programs. Candidates should send curriculum vitae, statement of research interests, and arrange for three reference letters to be sent to: Hasina Hason, University of Minnesota, Department of Cell Biology and Neuroanatomy, 4-144 Jackson Hall, 321 Church St. S.E., Minneapolis MN 55455. The last date for receipt of applications is March 1, 1998.

The University of Minnesota is an equal opportunity educator and employer.

Cato Research Ltd. (CRL)

is a contract research organization with locations in Research Triangle Park, NC, Washington, DC, and Montreal, Canada. CRL engages in the planning and execution of regulatory strategy for pharmaceutical and biological product development. Our goal is to assist clients in developing and gaining regulatory approval for products in conventional and cutting-edge therapeutic areas and novel drug delivery technologies.

Our company has exciting non-laboratory opportunities at all corporate locations for self-starters with scientific backgrounds who want to learn and grow in a creative environment. These positions require excellent written and verbal communication skills, strong initiative and interpersonal abilities, and sound critical and analytical thinking. All positions require an M.D. or Ph.D. with expertise in toxicology, pharmacokinetics, infectious disease, immunology, oncology, pharmacology or other biological sciences.

Clinical Research Scientists

We are seeking individuals with clinical and scientific training to provide leadership in ensuring that client contracts are met. These opportunities require the ability to design, direct, coordinate and manage drug development and regulatory activities. The ideal candidates will have 5–8 years of experience in the pharmaceutical or biotech industry and knowledge of the entire drug development process.

Clinical Research Fellows

The Fellows receive training in the scientific, regulatory, medical and financial aspects of pharmaceutical drug development. The ideal candidates will possess a Ph.D. in Biological Sciences and the ability to manage multiple tasks.

If you are an excellent communicator and a team player with problem-solving skills, and wish to expand your career, you will find these opportunities challenging and professionally rewarding.

Cato Research Ltd. offers complete salary and benefits packages. For application to all CRL offices, please send your resume, cover letter and writing sample to: Job 400, Cato Research Ltd., 200 Westpark Corp. Ctr., 4364 S. Alston Ave., Durham, NC 27713-2280. No phone calls, please. EOE

Maihle, N.J.
Appendix I
Ph.D. Degree

Students in the Tumor Biology Track must complete the following, in addition to the 13 core credit requirement:

<table>
<thead>
<tr>
<th>Course Code</th>
<th>Course Title</th>
<th>Credits</th>
</tr>
</thead>
<tbody>
<tr>
<td>TBio 5000</td>
<td>Tumor Biology I: Introduction to Tumor Biology</td>
<td>3 cr.</td>
</tr>
<tr>
<td>TBio 5100</td>
<td>Research Seminars in Tumor Biology</td>
<td>1 cr.</td>
</tr>
<tr>
<td>TBio 5150</td>
<td>Current Topics in Tumor Biology</td>
<td>1 cr.</td>
</tr>
<tr>
<td>TBio 5300</td>
<td>The Business of Science and the Science of Business</td>
<td>1 cr.</td>
</tr>
<tr>
<td>TBio 5858</td>
<td>Laboratory Rotations in Tumor Biology (2 cr./rotation - 3 rotations req.)</td>
<td>6 cr.</td>
</tr>
<tr>
<td>TBio 8000</td>
<td>Tumor Biology II: Origins of Human Cancer</td>
<td>2 cr.</td>
</tr>
<tr>
<td>TBio 8005</td>
<td>Tumor Biology III: Growth Factors, Oncogenes, and Tumor Suppressors</td>
<td>3 cr.</td>
</tr>
<tr>
<td>TBio 8200</td>
<td>Cell Biology of Cancer</td>
<td>3 cr.</td>
</tr>
</tbody>
</table>

In addition, all students in the Tumor Biology Program shall register for the Animal Care and Use Training Sessions and the American Association of Cancer Research (AACR) course in Histopathology of Cancer. Additional advanced elective courses in any area may be taken to fulfill the overall degree requirements of 42 credits.

The Tumor Biology program is supported in part by a predoctoral training award from the USAMRDC entitled “Biology of Breast Cancer.”
TUMOR BIOLOGY

TBio 5000f. TUMOR BIOLOGY I: INTRODUCTION TO TUMOR BIOLOGY. (3 cr) Maible, Tindall
Material to be covered includes fundamental concepts and methods in
tumor biology, as well as normal tissue histology and tumor pathobiology.

TBio 5100f, w, s, su. RESEARCH SEMINARS IN TUMOR BIOLOGY. (1 cr/yr)
Maible, Salisbury
Informal presentation of intramural research findings from the laboratories
involved in relevant research investigations. Discussions will be based on
chalk-talk format with the open research notebook. In addition, speakers
from outside the institution will present throughout the year. All Tumor
Biology trainees will be expected to present their research plans/findings
in this forum annually and will be encouraged to actively participate in
this highly multidisciplinary exchange of ideas and information.

TBio 5150f, w, s. CURRENT TOPICS IN TUMOR BIOLOGY. (1 cr) Salisbury,
Maible
This journal club will discuss current primary literature covering all
aspects of tumor biology with an emphasis on women's cancers. The
journal club will meet once per week and be conducted under the open
discussion format with directed student and faculty presentations. During
the fall quarter, journal articles of fundamental and historic interest in the
area of tumor biology will be read and discussed. Topics to be covered
include: cell cycle, oncogenes, tumor suppressors, growth factors, signal
transduction, metastasis, DNA tumor viruses, retroviruses.

TBio 5200f. PRINCIPLES OF PanCREATIC CANCER. (1 cr) Urtrutta
Anatomy, fine structure, and embryology of the pancreas. Basic cell
biology and regulation of pancreatic gene expression. Cellular and animal
models for the study of normal and neoplastic pancreatic cell
differentiation. Epidemiology, etiology, diagnosis and management of
pancreatic cancer.

TBio 5250w. GENE THERAPY AND CANCER. (1 cr; odd yrs) Federspiel,
Salisbury
Current papers in the area of gene therapy and cancer will be reviewed
and discussed in the journal club format. Students in the Tumor Biology
program will participate in all sessions and will present a paper during the
quarters that they are enrolled in this journal club.

TBio 5300su. THE BUSINESS OF SCIENCE AND THE SCIENCE OF BUSINESS.
(1 cr; offered even years) Bennet, Maible
This course reviews concepts fundamental to the commercial potential of
biotechnology. Topics include current patent issues in biotechnology,
regulatory issues in biotechnology and research funding mechanisms, as
well as the grant review process.
TBio 5858f.w.s.su. LABORATORY ROTATIONS IN TUMOR BIOLOGY (2 cr) Staff
Tutorial course involving general techniques, instrumental analysis, and special procedures undertaken in the laboratory of choice. In addition, the student will assimilate the general research area of the laboratory through readings, lab meetings, and discussion. Students and faculty shall use these rotations to determine the degree of general mutual interest in research topics for potential thesis projects.

TBio 8000w. TUMOR BIOLOGY II: ORIGINS OF HUMAN CANCER. (3 cr; prereq TBio 5000) Maihle, Tindall
Topics to be covered include: basic tumor biology, oncogenes, tumor viruses, anti- oncogenes (tumor suppressors), tumor immunity, cancer chemotherapy, and biological response modifiers. Also listed under Molecular Biology 8250.

TBio 8005s. TUMOR BIOLOGY III: GROWTH FACTORS, ONCOGENES, AND TUMOR SUPPRESSORS. (3 cr; prereq TBio 5000, TBio 8000) Maihle, Tindall
This course will focus on the mechanisms by which growth factors and oncogenes influence cell growth and division. Topics include: transmembrane signal transduction; cell cycle and regulation of cell division; ontogeny of oncogenes; mechanisms of oncogene activation; the insulin receptor family; PDGF/sis and PDGF receptor; EGF receptor/c-erb B 1 and 2 (neu); introduction to hematopoietic growth factors/receptors; receptors which lack intrinsic kinase activity, ras family of oncogenes; introduction to nuclear signal transduction; chromosome/DNA-binding proteins; development and differentiation; wound-healing and angiogenesis; carcinogenesis in humans; and anti-oncogenes. Also listed under Molecular Biology 8370.

TBio 8200w. CELL BIOLOGY OF CANCER. (2 cr; offered even years; prereq TBio 5000) Salisbury, Gendler, Lingle
This course will cover normal histology and the histopathology of neoplasia and will consist of one lecture and one laboratory session each week. Normal development and microscopic anatomy of the four basic tissue types will be covered, followed by a detailed examination of integument, hemopoietic system, male and female reproductive tracks, respiratory system, and GI tract. Specific primary and metastatic tumors of each system will also be covered. The laboratory session will involve study of microscopic slide preparations and problem set discussion sessions.

TBio 8305s. BIOLOGY OF BREAST CANCER. (1 cr; offered odd years) Maihle, Salisbury
This course will cover the cell and developmental biology of the breast and the histopathobiology of breast tumors. Experimental models for breast cancer, growth factors, oncogenes, and tumor suppressers in breast cancer will be covered. Clinical topics including radiation and chemotherapy, surgical treatments, diagnosis, and experimental therapies in breast cancer will also be presented.
TBio  8400  MASTER'S PROJECT IN TUMOR BIOLOGY (3 cr) Staff  
Readings and/or research in Tumor Biology culminating in the submission of the Master's Project. Topics will be chosen by the student in consultation with the adviser and the student's advisory committee.

Research

TBio  8840f,w,s,su. RESEARCH IN TUMOR BIOLOGY.  Staff  
Graduate thesis research for Master's students under supervision of staff.

TBio  8900f,w,s,su. RESEARCH IN TUMOR BIOLOGY.  Staff  
Graduate thesis research under supervision of staff.
## Biology of Breast Cancer Program Extramural Training Support

<table>
<thead>
<tr>
<th>Grant #</th>
<th>P.I.</th>
<th>Source</th>
<th>Term</th>
<th>Total Direct</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>T32 CA 75926</td>
<td>Salisbury, J.L.</td>
<td>NCI</td>
<td>7/1/98-6/30/03</td>
<td>$805,414</td>
<td>Biology of Cancer: A Predoctoral Training Program</td>
</tr>
<tr>
<td>NRSA</td>
<td>Baines, J.E.</td>
<td>NIH</td>
<td>9/15/97-5/31/02</td>
<td></td>
<td>Novel Immunotherapeutic Approaches to Cervical Cancer</td>
</tr>
<tr>
<td>NRSA</td>
<td>Canales, N.D.</td>
<td>NIH</td>
<td>5/1/97-5/1/02</td>
<td></td>
<td>Research Training in Breast and Ovarian Cancer</td>
</tr>
</tbody>
</table>
Biology of Breast Cancer:

A Predoctoral Training Program

"...for, if we would serve science, we must extend her limits, not only as far as our own knowledge is concerned, but in the estimation of others."

Rudolf Virchow, 1859
Biology of Breast Cancer:

A multidisciplinary predoctoral training program in the biology of breast cancer.

Mayo Clinic Foundation
Rochester, Minnesota 55905

Program Director:

Jeffrey L. Salisbury, Ph.D.
Guggenheim 1442 C
Phone: (507) 284-3326
FAX: (507) 284-1767
Email: salisbury@mayo.edu

Co-Director:

Nita J. Maihle, Ph.D.

Training Program Education Committee:

Jeffrey L. Salisbury, Ph.D., Tumor Biology
Mark J. Federspiel, Ph.D., Molecular Medicine
Diane F. Jelinek, Ph.D., Immunology
Donald J. Tindall, Ph.D., Urology Research
Nita J. Maihle, Ph.D., Tumor Biology
Sandra J. Gendler, Ph.D., Biochemistry and Molecular Biology
Jonathan E. Baines, M.D./Ph.D. Trainee

Tumor Biology Library - Guggenheim 1497
# Table of Contents

<table>
<thead>
<tr>
<th>Section</th>
<th>Page</th>
</tr>
</thead>
<tbody>
<tr>
<td>Goals for the Biology of Breast Cancer Program</td>
<td>1</td>
</tr>
<tr>
<td>Overview</td>
<td>1</td>
</tr>
<tr>
<td><strong>Program Structure</strong></td>
<td></td>
</tr>
<tr>
<td>Administrative Structure</td>
<td>2</td>
</tr>
<tr>
<td>Qualifications of the Program Faculty</td>
<td>2</td>
</tr>
<tr>
<td><strong>Program Curriculum</strong></td>
<td></td>
</tr>
<tr>
<td>Introduction</td>
<td>3</td>
</tr>
<tr>
<td>Summary of the Curriculum</td>
<td>3</td>
</tr>
<tr>
<td>Graduate School Core Offerings</td>
<td>3</td>
</tr>
<tr>
<td>Biology of Breast Cancer Course Offerings</td>
<td>3</td>
</tr>
<tr>
<td>Recommended Electives</td>
<td>3</td>
</tr>
<tr>
<td>Biology of Breast Cancer Track and Core Curriculum Schedule</td>
<td>4</td>
</tr>
<tr>
<td><strong>Program Implementation</strong></td>
<td></td>
</tr>
<tr>
<td>Innovative Strategies for Teaching</td>
<td>5</td>
</tr>
<tr>
<td>Course Structure</td>
<td>5</td>
</tr>
<tr>
<td>Balance of Historic and Current Scientific Perspective</td>
<td>6</td>
</tr>
<tr>
<td>Commitment, Accountability and Responsibility</td>
<td>6</td>
</tr>
<tr>
<td>Continuing Education</td>
<td>6</td>
</tr>
<tr>
<td>Integration of the Clinical Perspective</td>
<td>6-7</td>
</tr>
<tr>
<td>Additional Academic Activities</td>
<td>7-8</td>
</tr>
<tr>
<td><strong>Research Training</strong></td>
<td></td>
</tr>
<tr>
<td>Selection of Thesis Laboratory, Mentor, and Thesis Committee Members</td>
<td>8</td>
</tr>
<tr>
<td>Thesis Research</td>
<td>8</td>
</tr>
<tr>
<td>Ph.D. Thesis</td>
<td>8</td>
</tr>
<tr>
<td><strong>Student Recruitment, Progress and Track Evaluation</strong></td>
<td></td>
</tr>
<tr>
<td>Trainee Candidates</td>
<td>9</td>
</tr>
<tr>
<td>Trainee Evaluation</td>
<td>9</td>
</tr>
<tr>
<td>Curriculum and Program Evaluation</td>
<td>9</td>
</tr>
<tr>
<td><strong>Syllabus Outlines for Selected Tumor Biology Track Courses</strong></td>
<td>11-19</td>
</tr>
<tr>
<td>Training Support</td>
<td>20</td>
</tr>
<tr>
<td>Program Predoctoral Trainees</td>
<td>20</td>
</tr>
<tr>
<td>Student Highlights</td>
<td>21</td>
</tr>
<tr>
<td>Faculty Listing</td>
<td>22-24</td>
</tr>
<tr>
<td>Mayo Research Facilities</td>
<td>25</td>
</tr>
<tr>
<td>Mayo Core Facilities</td>
<td>25-27</td>
</tr>
</tbody>
</table>
Biology of Breast Cancer:

A Program for Graduate Training in Tumor Biology

The goals of the Biology of Breast Cancer and Tumor Biology Training Program are four-fold:

- First, to provide trainees with a solid and uniquely multidisciplinary knowledge base in the biology of cancer using breast cancer as the paradigm.

- Second, to guide the development of each individual trainee so that they achieve their fullest academic and research potential.

- Third, to aid trainees in the establishment of their professional network of peers and colleagues in the field of breast cancer research.

- Fourth, to stimulate new working alliances between students, fellows, and staff participating in breast cancer research, education, and clinical endeavors at the Mayo Clinic and within the Mayo Cancer Center.

Overview

The “Biology of Breast Cancer” is a multidisciplinary predoctoral training program in the biology of cancer, with a specific emphasis on breast cancer. The focus of the program is to provide an educational environment that stimulates excellence in scientific thought and training while simultaneously providing exposure to all of the major fields of study relevant to tumor biology. While a defining feature of this program is its research focus and integral link with clinical aspects of breast cancer, a general foundation in tumor biology is both important and essential in achieving this goal. The curriculum for this program is outlined in the course syllabus material, provided below, and the thesis research topics of the students matriculating in the program. This information clearly details and substantiates the major breast cancer research focus of this new training program. Research and training are broadly focused on gene regulation, cell cycle control, cancer genetics, oncogene and tumor suppressor action, tumor immunology, signal transduction, antitumor pharmacology, with a particular emphasis on breast cancer, but also includes investigators with research programs in ovarian, uterine, lung, G.I., brain, and prostate cancers. Students participate in laboratory-based research, as well as in a formal tumor biology curriculum that integrates current concepts in cell growth control with the natural history of human tumors.

The Biology of Breast Cancer Training Program has been supported by extramural training grants since its inception. Currently, these training grants include an award from the US Army Medical Research and Materiel Command in the “Biology of Breast Cancer” (DAMD17-94-J-4116), and a T32 Training Grant from the National Cancer Institute in “Tumor Biology” (CA75926). In addition, the program operates under the generous support of the Mayo Foundation through the Mayo Graduate School and the Mayo Cancer Center. Individual trainees also have been successful in competition for individual research awards.
Program Structure

Administrative Structure

The training program is administered through the Mayo Graduate School, and is closely allied with the Mayo Cancer Center. Day-to-day program administration operates largely through the Director (Dr. Salisbury) and Co-director (Dr. Maihle) of the Biology of Breast Cancer and Tumor Biology Training grants. Long range planning and administration operates through the Tumor Biology Education Committee (Drs. Salisbury, Maihle, Federspiel, Jelinek, and Tindall, and a trainee, Mr. J. Baines). The Education Committee meets each academic quarter to discuss student recruitment, student progress, and coordination within the Tumor Training Program curriculum. In addition, the directors of the three cancer-related pre- and postdoctoral training grants (Drs. Salisbury, Maihle, Getz, and David) and the Director of the Mayo Cancer Center (Dr. Prendergast) interact to coordinate ongoing programs and activities related to cancer research and education at Mayo in general. The Biology of Breast Cancer Training faculty also meet quarterly, in addition to frequent interactions through participation in program courses, journal clubs, research workshops, and a biweekly social hour called the “Tumor Biology Tea.”

Qualifications of the Program Faculty

The training faculty consists of approximately 50 full and associate members drawn from each of the basic science departments, as well as clinical faculty who participate in scholastic activities of the program but who do not have active research laboratories. The level of individual faculty participation varies each year for specific courses, topics and journal clubs. Nonetheless, a growing and enthusiastic cadre of participating faculty has emerged. In future years the program may elect to restructure its faculty based on degree of faculty participation, given the mounting enthusiasm for this program, as well as recent and ongoing recruitment of new staff in the area of cancer biology. At this time, however, faculty privileges will remain as they are in order to promote both the inclusively and multidisciplinary nature of this new training program. A listing of the participating faculty is given on pages 23-25. While most individual faculty are associated with traditional discipline-based basic science and clinical departments (such as Oncology, Molecular Biology, Experimental Pathology, Pharmacology, etc.) the administrative structure of the Program is that of an interdisciplinary programmatic unit. This programmatic structure reflects the interdisciplinary nature of the major research and academic efforts of the associated faculty.

- Full members of the training faculty have an established track record of accomplishment in biomedical research as demonstrated by significant publications of high scientific merit, excellence, and innovation. Overall, the faculty have consistent records of extramural funding in support of their individual research programs.
- The collective interests of this training faculty are quite broad, but all show direct breast cancer relevance. These interests include: cell signaling, cancer genetics, gene regulation, tumor immunology, oncogene and tumor suppressor action, cell cycle regulation, tumor virology, gene therapy, hormonal regulation, and molecular cytology. Most training faculty are also members of the NCI-designated Mayo Cancer Center.
- Faculty drawn from both clinical and basic science departments contribute to the Training Program through participation in a variety of relevant educational activities and as clinical instructors. Faculty from outside the program may serve as advisory members of qualifying examination and thesis committees, however, they may not serve as research mentors for students in the Biology of Breast Cancer Training Program.
Program Curriculum

Introduction

The program curriculum and thesis research is a predoctoral training program leading to the Ph.D. degree in Biomedical Sciences. Each year, 3 to 5 students are accepted into the program for an appointment term of 4 to 5 years. The program strives to maintain a steady state level of 15 to 20 students. Trainee stipends initially are supported through institutional funds. Students qualify for support from extramural training grants (third through fifth year), following successful completion of the qualifying examination. The training program curriculum includes didactic course work, journal clubs, research seminars and workshops, and tutorial and special clinical activities. Students who matriculate into the program must meet the general course requirements of the Mayo Graduate School in which a minimum of 15 credits are required from the Graduate School Core Curriculum. Students must also complete 20 credits from the didactic Tumor Biology Program curriculum and the remainder of their credit requirements can be completed through elective courses offered by the Tumor Biology Program, the Mayo Graduate School or by special topics courses given at other institutions and sanctioned by the Mayo Graduate School. For students in the Biology of Breast Cancer Program these electives must include a course entitled “Biology of Breast Cancer.” A student's program of core courses is individually developed by the Education Committee in consultation with the student and his/her advisor.

Summary of the Curriculum:

Graduate School Core Offerings (15 Credits, minimum)

- Genome Biology (3)
- Immunology (3)
- Principles of Cell and Tissue Design (3)
- Biochemistry (3)
- Genetics (1)
- Pharmacology (2)
- Developmental Biology & Statistics (1)
- Biology of Disease (1)
- Ethics (1)

Required Biology of Breast Cancer Course Offerings (20 Credits)

- Tumor Biology I: Introduction to Tissue and Tumor Biology (3)
- Tumor Biology II: Origins of Human Cancer (3)
- Tumor Biology III: Growth Factors, Oncogenes, and Tumor Suppressors (3)
- Biology of Breast Cancer (1)
- Current Topics in Tumor Biology: Journal Club (1) x 3
- Research Seminars in Tumor Biology and Tumor Biology Interest Group (1)
- AACR Course in Histopathology of Cancer, Keystone CO. (1)
- Laboratory Rotations in Tumor Biology (3 required rotations, 2 credits each = 6 total)
- Research in Tumor Biology (Thesis Research (0))

Recommended Electives

- Quantitative Biology I-III, Neuroscience (1), Integrated Physiology (5)
- Tumor Immunology (1), Business of Science and the Science of Business (1)
- Cytogenetics (2)
<table>
<thead>
<tr>
<th>Year I or II</th>
<th>M</th>
<th>T</th>
<th>W</th>
<th>Th</th>
<th>F</th>
</tr>
</thead>
<tbody>
<tr>
<td>Fall Quarter</td>
<td>Biochemistry</td>
<td>Immunology</td>
<td>Biochemistry</td>
<td>Immunology</td>
<td>Biochemistry</td>
</tr>
<tr>
<td>9:00-10:00</td>
<td>Genome Biology</td>
<td>Genome Biology</td>
<td>Genome Biology</td>
<td>Genome Biology</td>
<td>Genome Biology</td>
</tr>
<tr>
<td>11:00-12:30</td>
<td>Tumor Biology I</td>
<td>Tumor Biology Journal Club</td>
<td>Tumor Biology I</td>
<td>Tumor Biology I</td>
<td>Tumor Biology I</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Year I or II</th>
<th>M</th>
<th>T</th>
<th>W</th>
<th>Th</th>
<th>F</th>
</tr>
</thead>
<tbody>
<tr>
<td>Winter Quarter</td>
<td>Cell Biology</td>
<td>Genetics</td>
<td>Cell Biology</td>
<td>Cell Biology</td>
<td>Cell Biology</td>
</tr>
<tr>
<td>9:00-10:00</td>
<td>Cell Biology</td>
<td>Genetics</td>
<td>Cell Biology</td>
<td>Cell Biology</td>
<td>Cell Biology</td>
</tr>
<tr>
<td>11:00-12:30</td>
<td>Tumor Biology II</td>
<td>Tumor Biology Journal Club</td>
<td>Tumor Biology II</td>
<td>Tumor Biology II</td>
<td>Tumor Biology II</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Year I or II</th>
<th>M</th>
<th>T</th>
<th>W</th>
<th>Th</th>
<th>F</th>
</tr>
</thead>
<tbody>
<tr>
<td>Spring Quarter</td>
<td>Pharmacology</td>
<td>Pharmacology</td>
<td>Pathobiology</td>
<td>Pathobiology</td>
<td>Pathobiology</td>
</tr>
<tr>
<td>9:00-10:00</td>
<td>Pharmacology</td>
<td>Pharmacology</td>
<td>Pathobiology</td>
<td>Pathobiology</td>
<td>Pathobiology</td>
</tr>
<tr>
<td>10:00-11:00</td>
<td>Development</td>
<td>BioStatistics</td>
<td>Development</td>
<td>BioStatistics</td>
<td>Development</td>
</tr>
<tr>
<td>11:00-12:30</td>
<td>Tumor Biology III</td>
<td>Tumor Biology Journal Club</td>
<td>Tumor Biology III</td>
<td>Tumor Biology III</td>
<td>Tumor Biology III</td>
</tr>
</tbody>
</table>

Additional advanced elective courses may be chosen in any area from the Mayo Graduate School Bulletin to fulfill the overall degree requirement of 35 credits. In addition, all students in the training program are required to take formal classes in Radiation Safety, Animal Care and Use, and also participate in an NIH Grant Writing Workshop. These required courses are not administrated by the Mayo Graduate School, and, therefore, are not offered for Graduate School credit.
Program Implementation

Innovative Strategies for Teaching

Tumor Biology track courses establish a solid foundation in the biology of cancer. In practice, the biology of breast cancer is emphasized as an illustrative paradigm throughout the program curriculum. Issues of fundamental importance to understanding breast cancer are also relevant to other cancers, and the converse is also true. Likewise, there are many instances where the illustrating example of a relevant point may involve biological systems as diverse as yeast and frogs (not breast cancer research, but directly relevant to improving our understanding of breast cancer). Thereby, the tumor biology curriculum is grounded in basic cancer biology with the aim of leading students to a thorough understanding of all aspects at the forefront of breast cancer research. In order to accomplish this, our curriculum has been strategically developed to provide a strong foundation in the study of cancer using breast cancer as the model wherever possible. For example, to illustrate the significant emphasis on and integration of breast cancer in the overall curriculum, topics in breast cancer are featured in three 3 credit hour courses in the curriculum (see below), as well as in approximately one out of every three journal club presentations. Moreover, all trainees are required to register for a didactic course in “The Biology of Breast Cancer” (T BIO 8305, page 134 in the Mayo Graduate School Bulletin). The three major courses were organized and are taught using breast cancer as the principle paradigm for instruction and content: “Introduction to Tissue and Tumor Biology” (Tumor Biology I, T BIO 5000), “Origins of Human Cancer” (Tumor Biology II, T BIO 8000), and “Growth Factors, Oncogenes and Tumor Suppressors” (Tumor Biology III, T BIO 8005).

Tumor Biology Track courses share several distinguishing features and innovative strategies for teaching which enhance learning and student/faculty participation.

- **Course Structure**: Tumor Biology I, II, and III are given as an integrated series during the first three consecutive quarters following matriculation. Individually, these courses meet three times per week (1 1/2 hour per session) with an overview and historical review of a selected topic(s) for the current week presented in a didactic lecture format during the first class session. This is followed by a student presentation of a current or historically relevant research paper(s) in the area of the week’s topic during the second class session using the journal club format. Finally, a round table small group problem set discussion format is used to focus on questions and problems relevant to the week’s topic during the third class session. The research paper presentation and problem set discussions are carefully organized in order to thoroughly integrate research topics relevant to the theme of the week. The effectiveness of these active learning/teaching strategies is apparent to all participants in these courses. The active involvement of students in the learning and discovery process, in information processing, and in the application of information to problems requires that students are accountable for learned information on an immediate and ongoing basis. Problem centered learning also puts learning into context and facilitates learning transfer. These sessions allow students to organize and categorize information into meaningful units and to ‘discover’ novel relationships and extract and assimilate important points in an interactive and participatory manner.
- **Balance of Historic and Current Scientific Perspectives**: Given the rapid pace of progress in the biological sciences and the exponential rate of growth of relevant literature, the general philosophy that is promoted within the program is to *teach less better*. The objective here is to lay a strong foundation in cancer biology with the clear understanding that what is particularly relevant and important today, may not be so tomorrow. Therefore, emphasis is placed on developing effective learning skills, and the application of these skills to both historic paradigms, as well as critical review and evaluation of issues at the forefront of modern cancer biology.

- **Commitment, Accountability, and Responsibility**: Integral to the Tumor Biology Program teaching philosophy is *Peer Performance Assessment* and the *Team Learning Model*. These strategies create a climate in which all students are encouraged to grow. This results in a classroom environment where students from diverse backgrounds (including clinical fellows) feel welcome to fully participate in discussions and problem solving. In this way, desired student performances are tied directly to the efforts of the students themselves, to the involvement of students in the teaching-learning process, to the opportunities to make choices, and to the degree to which they interact with their peers and instructors. Emphasis is placed on organization and presentation skills, accountability tracking, and peer assessment and feedback. Our experience with this learning model is that trainees rapidly gain a level of professional expectation of their peers (and themselves) that both promotes and enhances the general level of academic and scholastic effort among trainees.

- **Continuing Education**: Senior students who have completed the formal didactic course requirements, and postdoctoral fellows who are supported by other cancer-related training grants actively participate in the journal club sessions that are an integral component of the Tumor Biology curriculum. In this way, senior trainees contribute to the critical mass of the class and also enhance the sophistication and the multidisciplinary nature of the discussions. Through this mechanism, more advanced trainees revisit current topics in cancer biology throughout their advanced training years. In this manner, advanced trainees have the opportunity to reinforce key concepts, to help new trainees better understand these concepts, and also to remain current with the rapid pace of development in the dynamic field of cancer biology. Likewise, Program faculty (and their laboratory personnel) actively participate in these regular weekly journal club sessions.

- **Integration of the Clinical Perspective**: The training program is designed to give the trainee a broad and well-rounded understanding of cancer from the basic science,
population science, and clinical perspectives. Integration of the clinical activities of the Mayo Cancer Center into the training program is achieved in several ways:

◊ First, clinical fellows and residents from a broad spectrum of cancer relevant programs (e.g., oncology, hem/onc, orthopedic research, gynecologic oncology, pediatric oncology, etc.) formally enroll and participate in training program courses and journal clubs. Active participation by clinical residents and fellows adds considerably to the multidisciplinary perspective of the student body, and to stimulating class discussions.

◊ Second, clinical staff give didactic lectures in areas of their specialty in Biology of Breast Cancer Training Program courses. For example in Tumor Biology II and III, individual lectures are given by clinical staff practicing in Surgery, Medical Oncology, Radiation Oncology, and Surgical Pathology.

◊ Third, program trainees are required to attend Mayo Cancer Center Research Workshops, Mayo Cancer Center Grand Rounds, and appropriate departmental, Research Society, Oncology Society, and Hematology Society lectures, and receive course credit for participating in these activities (TBio 5101: Research Seminars in Tumor Biology). Students are made aware of relevant speakers through Email, direct mailings, and weekly announcements during the weekly journal club.

◊ The program curriculum is integrated with clinical practice wherever possible through special course related activities. For example, small group tours of the Surgical Pathology Suite (in TBio I) allow students to observe gross dissection of surgical specimens (including breast tumors), rapid freezing and cryomicrotomy, microscopic examination (via video monitors) and diagnosis by staff pathologists, and reporting to operating room surgeons. Additionally, Tumor Biology Program trainees are required to attend clinical rounds with a Mayo staff member (any division or department). Direct exposure to clinical activities such as these are useful for students to understand, based on first-hand observation, the intensity, dedication, and skill involved in the clinical care and treatment of cancer patients. These activities also promote the involvement of our clinical faculty in Biology of Breast Cancer Training Program activities.

◊ Finally, Tumor Biology trainees may have one or more clinical staff advisors participate as members of their Thesis Advisory Committee. Sometimes this involvement is fairly technical, e.g., participation by a Mayo Cancer Center biostatistician in study design and analysis. In other instances, however, clinical advisors may be directly involved in helping the trainee define a clinically relevant question, and/or assist them with tumor specimen acquisition and/or data analysis.

Additional Academic Activities

Seminars by Students, Faculty and Invited Speakers: Extensive institutional resources support seminars by nationally and internationally recognized scientists and clinicians on the Mayo Rochester campus. Approximately 350-400 speakers come to the Rochester campus each year. Trainees are, therefore, exposed to diverse biomedical research opportunities, and institutionally and departmentally-based research seminars throughout the year. In December of 1997, students of the Tumor Biology Program hosted Dr. Judah Folkman (Harvard University) who presented the Annual Findling Lecture of the Mayo Graduate School. The 1998-1999 academic year will feature a series of visiting speakers who will focus on the broad topic of Epigenetics and Cancer. During each of their research years, Tumor Biology Trainees also present research seminars and research posters in multidisciplinary research workshops and retreats (e.g., the Mayo Graduate School Annual Research Symposium, the Joint Mayo Cancer Center/Laboratory Medicine Retreat). More recently, we have formalized the Tumor Biology Interest Group (TBIG) for Mayo Graduate
School course credit. This monthly research workshop provides the opportunity for all Tumor Biology trainees (pre- and postdoctoral) to regularly present their research plans, proposals, and results in a constructively critical internal forum.

**Attendance at National Research Meetings:** All students are supported to attend at least one national scientific meeting each year even if they are not presenting an abstract. If they are presenting their work, attendance at additional meetings is encouraged and supported by the research mentor's laboratory. Mentors take an active role in introducing students to the professional culture and ‘networking’ critical to success in any biomedical research career through this mechanism. In recent years, Biology of Breast Cancer Training Program trainees have attended and presented at the following national meetings: AACR, ASCB, FASEB, Annual Oncogene Meetings, Annual Human Cancer Meeting, Cold Spring Harbor Cancer Genetics Meetings, Salk Tyrosine Phosphorylation Meetings, and various Gordon and Keystone Conferences.

**Research Training**

**Selection of Thesis Laboratory, Mentor, and Thesis Committee Members:** Trainees typically matriculate in June through August and are required to complete three laboratory-based (minimum 8 weeks each) rotations during their first year. Any laboratory-based Mayo Graduate School faculty member may serve as a mentor for these research rotations. Selection of the thesis mentor follows completion of successful laboratory-based rotations by mutual consent of the student and mentor with the sanction of the Training Program Education Committee and Mayo Graduate School Education Committee. The qualifying examination consists of a written thesis proposal, an oral presentation (TBIG forum), and its defense before a thesis committee consisting of a minimum of 4 faculty (including the thesis advisor), and when appropriate, an extramural committee member from outside the institution. Typically clinical or extramural committee members' research specialties are related to the general area of the student’s thesis topic. Following successful completion of the qualifying examination, research progress is assessed through regular Thesis Advisory Committee meetings (minimum of one Thesis Advisory Committee meeting per year). While Thesis Advisory Committee members are available for advice, technical assistance, and consultation throughout the year, these meetings provide a formal opportunity for input by the Thesis Advisory Committee on progress and experimental aspects of the thesis project. The chair of the Thesis Advisory Committee formally reports the outcome of each committee meeting in writing to the Training Program Education Committee and to the Mayo Graduate School.

**Thesis Research:** The Biology of Breast Cancer Training Program places strong emphasis on thesis research. All laboratory-based faculty have demonstrated records of research training at both the predoctoral and postdoctoral levels. The specific details of an individual student’s research training plan are developed following the selection of a thesis advisor and in consultation with the Thesis Advisory Committee. The thesis research project must be hypothesis driven and experimental in nature and must, in addition, have a direct relevance to the biology of cancer.

**Ph.D. Thesis:** The thesis is the most important document that the Ph.D. candidate prepares during the course of graduate study, and is a record of the scientific accomplishments that justify the awarding of the Ph.D. degree. The thesis is archival. Consequently, the Mayo Graduate School has developed standards for its format and style, and our trainees adhere to these guidelines. The thesis examination consists of a formal thesis research seminar open to all members of the Mayo community followed by a meeting with the Thesis Examining Committee during which the scientific merit and accomplishments of the candidate are evaluated. Successful completion of a research thesis typically also results in two or more research manuscripts submitted for publication in peer-reviewed journals of high scientific standards.
Student Recruitment, Progress and Track Evaluation

Trainee Candidates: Students recruited into the Biology of Breast Cancer Training Program are selected on the basis of outstanding academic credentials, a stated desire to study and conduct research in the area of breast cancer biology, and an assessment of individual research potential by the training faculty. Many of the applicants to the Mayo Graduate School have had research experience within the Mayo system through summer undergraduate research internships. Typically, candidates for admission to Mayo’s graduate programs apply directly to the Graduate School where their academic credentials, letters of recommendation, and personal statements are placed on record. Applicants are selected for on-site interview by the Mayo Graduate School Admissions Committee. The interview process involves faculty and student assessment of each applicant’s research and academic interests. The Biology of Breast Cancer Training Program also has placed special emphasis on recruitment and training of under-represented minorities. Currently the predoctoral class consists of a total of 16 students, 3 of whom are minorities (19%, including one Native American, and two Hispanic students).

Trainee Evaluation

Trainee evaluation takes place at several levels and is assessed by comparison of established and objective data relating recruitment credentials, program completion, academic performance, placement, and ultimately career achievement.

- Academic performance of trainees, including coursework evaluation and consideration of reports from the trainee’s Qualifying Exam and Thesis Advisory Committees.
- Successful completion of the degree program.
- Success in gaining competitive pre- and postdoctoral fellowships and/or extramural funding.
- Ultimately, the appointment of these trainees to independent research positions with evidence of ongoing research activities relevant to cancer biology.

Curriculum and Program Evaluation

Curriculum and Program evaluation includes the areas listed below, as well as additional areas defined by the External Review Committee:

- Ability to recruit and retain outstanding Ph.D. candidates.
- Course content and appropriateness to the biology of breast cancer.
- Thoroughness of didactic and formal training in the biology of cancer.
- Effectiveness of teaching and examining methods and procedures.
- Vitality and effectiveness of student/faculty interactions in the academic components of the program.
- Evidence of faculty mentorship and establishment of intramural and extramural professional networks.
- Scope and role of individual faculty participation in the Biology of Breast Cancer Training Program.
- Integration of the clinical perspective and understanding of physician and patient concerns in the diagnosis and treatment of cancer.
- Overall effectiveness of the Program Director, Co-Director, Education Committee, and of the Graduate School in administrating the Biology of Breast Cancer Training Program.
In addition to these standards to be used for self-evaluation (through surveys of both trainees and faculty), the Biology of Breast Cancer Training Program has successfully completed two rigorous external reviews, each of which has resulted in an extramural training award (i.e., the US Army Medical Research and Materiel Command award for predoctoral training in the “Biology of Breast Cancer” (DAMD17-94-J-4116), and a T32 Training Grant from the National Cancer Institute for predoctoral training in “Tumor Biology” (CA75926)). One aspect of these predoctoral awards is a requirement for formalization of an External Advisory Committee and periodic external review by this Committee.

**Syllabus Outlines for Selected Tumor Biology Track Courses**

Mayo Tumor Biology track courses cover a series of topics of historic relevance and primary importance to cancer biology. In addition, each year course organization and content evolve according to current trends and in order to incorporate breaking forefront issues in this dynamic field. Cancer Course syllabus outlines for 1998, indicating topic, format, and faculty are listed on the following pages.
<table>
<thead>
<tr>
<th>Date</th>
<th>Topic</th>
<th>Instructor</th>
</tr>
</thead>
<tbody>
<tr>
<td>September 29</td>
<td>Principles of Cell and Tissue Design</td>
<td>Salisbury</td>
</tr>
<tr>
<td>September 30</td>
<td>TBJC: Fundamentals of the Cell Cycle</td>
<td></td>
</tr>
<tr>
<td>October 1</td>
<td>Laboratory - Light and Electron Microscopy</td>
<td></td>
</tr>
<tr>
<td>October 6</td>
<td>Stem Cells, Differentiation, and Cancer</td>
<td>Maible</td>
</tr>
<tr>
<td>October 7</td>
<td>TBJC: Genomic Instability/Aneuploidy</td>
<td></td>
</tr>
<tr>
<td>October 8</td>
<td>Guest Lecture - Epigenetics and Genetics</td>
<td></td>
</tr>
<tr>
<td>October 13</td>
<td>Properties of Transformed Cells <em>in vitro</em></td>
<td>Maible</td>
</tr>
<tr>
<td>October 14</td>
<td>TBJC: Temin - Hayflick</td>
<td></td>
</tr>
<tr>
<td>October 15</td>
<td>Senescence and Immortalization</td>
<td></td>
</tr>
<tr>
<td>October 20</td>
<td>Properties of Transformed Cells <em>in vivo</em></td>
<td>Maible</td>
</tr>
<tr>
<td>October 21</td>
<td>TBJC: Transgenics and Knockouts</td>
<td></td>
</tr>
<tr>
<td>October 22</td>
<td>Xenografts</td>
<td></td>
</tr>
<tr>
<td>October 27</td>
<td>Tissue Biology - Epithelia</td>
<td>Salisbury</td>
</tr>
<tr>
<td>October 28</td>
<td>TBJC: Cell Polarity</td>
<td></td>
</tr>
<tr>
<td>October 29</td>
<td>Laboratory - Epithelia</td>
<td></td>
</tr>
<tr>
<td></td>
<td>[TERM PAPER OUTLINE DUE by 4:00 p.m.]</td>
<td></td>
</tr>
<tr>
<td>November 3</td>
<td>Tissue Biology - Connective Tissue</td>
<td>Salisbury</td>
</tr>
<tr>
<td>November 4</td>
<td>TBJC: Invasion and Metastasis</td>
<td></td>
</tr>
<tr>
<td>November 5</td>
<td>Laboratory - Connective Tissue</td>
<td></td>
</tr>
<tr>
<td>November 10</td>
<td>Endothelial Cells, Vascular Tissue and Lymphatics</td>
<td></td>
</tr>
<tr>
<td>November 11</td>
<td>TBJC: Angiogenesis</td>
<td>Salisbury</td>
</tr>
<tr>
<td>November 12</td>
<td>Laboratory - Vascular Tissue</td>
<td></td>
</tr>
<tr>
<td>November 17</td>
<td>Pathobiology of Cancer</td>
<td>Salisbury</td>
</tr>
<tr>
<td>November 18</td>
<td>TBJC: Epithelial / Mesenchymal Interactions</td>
<td></td>
</tr>
<tr>
<td>November 19</td>
<td>Surgical Pathology Tours</td>
<td></td>
</tr>
<tr>
<td>November 24</td>
<td>INDEPENDENT STUDY</td>
<td></td>
</tr>
<tr>
<td>November 25</td>
<td>INDEPENDENT STUDY</td>
<td></td>
</tr>
<tr>
<td>November 26</td>
<td>Thanksgiving</td>
<td></td>
</tr>
<tr>
<td>December 1</td>
<td>Normal Breast / Breast Cancer</td>
<td>Salisbury</td>
</tr>
<tr>
<td></td>
<td>[TERM PAPER DUE by 4:00 p.m.]</td>
<td></td>
</tr>
<tr>
<td>December 2</td>
<td>TBJC: Breast Tumor Staging and Grade</td>
<td></td>
</tr>
<tr>
<td>December 3</td>
<td>Laboratory - Breast Pathology</td>
<td></td>
</tr>
<tr>
<td>December 8</td>
<td>Normal Intestine / Colon Cancer</td>
<td>TBA</td>
</tr>
<tr>
<td>December 9</td>
<td>TBJC: HNPCC - MIN Mice</td>
<td></td>
</tr>
<tr>
<td>December 10</td>
<td>Laboratory - GI Tumors</td>
<td></td>
</tr>
<tr>
<td>December 17</td>
<td>[TERM PAPER EVALUATIONS DUE by 4:00 p.m.]</td>
<td></td>
</tr>
</tbody>
</table>
Tumor Biology II:
Origins of Human Cancer (TBio 8000)
2:30 - 4:00 p.m. Tue. Thur. Winter Quarter 1998
[50% participation/50% term paper]

January 7  Origins of Human Cancer: An Overview  Maihle
January 8  Tumor Biology Journal Club:
January 9  Problem Set (Maihle)

January 14 Origins of Human Cancer: Etiology and Genetics  Smith
January 15 Tumor Biology Journal Club:
January 16 Problem Set

January 21 Origins of Human Cancer: Progression and Metastasis  Gendler
January 22 Tumor Biology and Journal Club: Angiogenesis
January 23 Problem Set (Gendler and Maihle)

January 28 Origins of Human Cancer: Epidemiology and Prevention  Yang
January 29 Tumor Biology Journal Club: Intestinal Polyposis and COX-2
January 30 Tumor Immunology: An Overview  Mitchell

February 4 Problem Set (Epidemiology and Prevention)
February 5 Tumor Biology Journal Club (Tumor Immunology)  Jelinek
February 6 Problem Set (Tumor Immunology)

February 11 Paraneoplastic Syndromes (in Breast and Ovarian Cancer)  Lennon
February 12 Tumor Biology Journal Club: Paraneoplastic Autoimmunity
February 13 Problem Set

February 25 Introduction to Clinical Research  O'Fallon
February 26 Tumor Biology Journal Club: Phase I Trial of Dolastatin-10
February 27 Problem Set

March 4 Introduction to Chemotherapy  Ames
March 5 Tumor Biology Journal Club: Inhibitors of Farnesyl Transferase
March 6 Problem Set

March 11 Tumor Imaging: An Overview  Robb
March 12 Experimental Tumor Imaging  Ehman
March 13 Problem Set (Maihle)

March 18 Introduction to Surgical Oncology  Nelson
March 19 Tumor Biology Journal Club: Surgical Procedures in Colon Cancer

March 25 Introduction to Radiation Therapy  Bonner
March 26 Breast Cancer Patient Vignettes
March 27 Experimental Gene Therapy  Raffel
## Tumor Biology III

**Growth Factors, Oncogenes, and Tumor Suppressors (TBIO 8005)**  
2:30 - 4:00 p.m. Tue. Thur. Winter Quarter 1998  
[50% participation/50% term paper]

<table>
<thead>
<tr>
<th>Date</th>
<th>Topic</th>
<th>Instructor</th>
</tr>
</thead>
<tbody>
<tr>
<td>April 7</td>
<td>Cell Cycle and Cell Growth Control</td>
<td>Salisbury</td>
</tr>
<tr>
<td>April 8</td>
<td>Tumor Biology Journal Club</td>
<td></td>
</tr>
<tr>
<td>April 9</td>
<td>Regulation of Immediate Early Gene Expression</td>
<td>Getz</td>
</tr>
<tr>
<td></td>
<td>(AACR Meeting 3/28 - 4/1)</td>
<td></td>
</tr>
<tr>
<td>April 14</td>
<td>Growth Factors/GF Receptors</td>
<td>Maihle</td>
</tr>
<tr>
<td>April 15</td>
<td>Tumor Biology Journal Club</td>
<td></td>
</tr>
<tr>
<td>April 16</td>
<td>Student Discussion Problem Set</td>
<td></td>
</tr>
<tr>
<td>April 21</td>
<td>Intracellular Mediators: Kinases and Phosphatases</td>
<td>Maihle</td>
</tr>
<tr>
<td>April 22</td>
<td>Tumor Biology Journal Club</td>
<td></td>
</tr>
<tr>
<td>April 23</td>
<td>Student Discussion Problem Set</td>
<td></td>
</tr>
<tr>
<td>April 28</td>
<td>Intracellular Mediators: G Proteins</td>
<td>Karnitz</td>
</tr>
<tr>
<td>April 29</td>
<td>Tumor Biology Journal Club</td>
<td></td>
</tr>
<tr>
<td>April 30</td>
<td>Student Discussion Problem Set</td>
<td></td>
</tr>
<tr>
<td>May 5</td>
<td>Oncogenes and Viral Oncogenes</td>
<td>Maihle</td>
</tr>
<tr>
<td>May 6</td>
<td>Tumor Biology Journal Club</td>
<td></td>
</tr>
<tr>
<td>May 8</td>
<td>Student Discussion Problem Set</td>
<td></td>
</tr>
<tr>
<td>May 12-14</td>
<td>INDEPENDENT STUDY</td>
<td></td>
</tr>
<tr>
<td>May 19</td>
<td>Introduction to Tumor Suppressors</td>
<td>James</td>
</tr>
<tr>
<td>May 20</td>
<td>Tumor Biology Journal Club</td>
<td></td>
</tr>
<tr>
<td>May 21</td>
<td>Student Discussion Problem Set</td>
<td></td>
</tr>
<tr>
<td>May 26</td>
<td>Cancer Genetics</td>
<td>Jenkins/Lloyd</td>
</tr>
<tr>
<td>May 27</td>
<td>Tumor Biology Journal Club</td>
<td></td>
</tr>
<tr>
<td>May 28</td>
<td>Student Discussion Problem Set</td>
<td></td>
</tr>
<tr>
<td>June 2</td>
<td>P53 - Guardian of the Genome</td>
<td>James</td>
</tr>
<tr>
<td>June 3</td>
<td>Tumor Biology Journal Club</td>
<td></td>
</tr>
<tr>
<td>June 4</td>
<td>Student Discussion Problem Set</td>
<td></td>
</tr>
<tr>
<td>June 9</td>
<td>Retinoblastoma and Rb</td>
<td>Smith</td>
</tr>
<tr>
<td>June 10</td>
<td>Tumor Biology Journal Club</td>
<td></td>
</tr>
<tr>
<td>June 11</td>
<td>Student Discussion Problem Set</td>
<td></td>
</tr>
<tr>
<td>June 16</td>
<td>To Die Or Not To Die - Apoptosis</td>
<td>TBA</td>
</tr>
<tr>
<td>June 17</td>
<td>Tumor Biology Journal Club</td>
<td></td>
</tr>
<tr>
<td>June 18</td>
<td>Student Discussion Problem Set</td>
<td></td>
</tr>
</tbody>
</table>
**Biology of Breast Cancer**

(TBiol 5200)
Guggenheim 1093
1:30-2:30 p.m. Fridays

(50% participation/50% final exam)

- This course is aimed at integrating basic concepts in developmental, cellular and molecular biology of the breast together with current information on the etiology, diagnosis and treatment of breast cancer.
- The faculty include members from diverse basic science and medical disciplines including cell and molecular biology, pathology, oncology and surgery.

<table>
<thead>
<tr>
<th>Date</th>
<th>Topic</th>
<th>Instructor</th>
</tr>
</thead>
<tbody>
<tr>
<td>April 11</td>
<td>Breast Cancer: The Magnitude of the Problem</td>
<td>Inge</td>
</tr>
<tr>
<td>April 18</td>
<td>Development, Anatomy and Histology and Cell Biology of the Breast</td>
<td>Salisbury</td>
</tr>
<tr>
<td>April 25</td>
<td>Histopathology of the Breast</td>
<td>Wold</td>
</tr>
<tr>
<td>May 2</td>
<td>Experimental Models of Breast Cancer</td>
<td>Gendler</td>
</tr>
<tr>
<td>May 9</td>
<td>Oncogenes, Growth Factors, and Breast Cancer</td>
<td>Leof</td>
</tr>
<tr>
<td>May 16</td>
<td>Tumor Suppressors and Breast Cancer</td>
<td>Mr. Ritland</td>
</tr>
<tr>
<td>May 23</td>
<td>Radiation Therapy for Breast Cancer</td>
<td>Peterson</td>
</tr>
<tr>
<td>May 30</td>
<td>Surgical Treatment of Breast Cancer</td>
<td>Donohue</td>
</tr>
<tr>
<td>June 6</td>
<td>Breast Cancer Diagnosis and Imaging</td>
<td>Johnson</td>
</tr>
<tr>
<td>June 13</td>
<td>Systemic Therapy for Breast Cancer</td>
<td>Inge</td>
</tr>
<tr>
<td>June 20</td>
<td>Experimental Therapies for Breast Cancer</td>
<td>Maille</td>
</tr>
<tr>
<td>June 27</td>
<td>Final Examinations Due (by 5:30 p.m.)</td>
<td></td>
</tr>
</tbody>
</table>
Business of Science, Science of Business

TBio 5300
K. E. Bennet, M.B.A. and N. J. Maihle, Ph.D.

Summer Quarter (even years)

1) Introduction (August 2) Orientation and Objectives [KEB/NJM]
2) Administrative Structures in Support of Research (August 5) [KEB]
   Not-for-Profit & For Profit
3) Overview of Research Accounting (August 7) [KEB]
   Research Budgets
   Direct versus Indirect costs
4) Sources of Financial Support for Research (August 9) [NJM]
   Intramural & Extramural Support
   Federal & Private
5) Sources of Financial Support for Research (August 12) [KEB]
   Corporate
   Strategic Alliance, Joint Development
   Licensing/Venture Capital
6) Introduction to Intellectual Property (August 14) [KEB]
   Definition of Intellectual Property
   Protection of Intellectual Property
   Patents & Trade Secrets
   Ownership of Intellectual Property
7) Introduction of Cases (August 16) [KEB]
8) Commercialization of Research Discoveries (August 21) [KEB]
   Licensing
   Market Value of Invention
9) Independent Study on Cases (August 19)
10) Laws and Policies Governing Conduct of Research (August 23)
    Institutional [NJM]
    State, Federal, and International [KEB]
11) Case Presentations "Levamisole" (August 26) [KEB/NJM]
12) Case Presentations - "University of Florida" (August 28) [KEB/NJM]
13) Course Wrap-Up (August 30) [KEB/NJM]
Current Topics in Tumor Biology

(TBiol 5151)
2:30 - 3:30 p.m. Wednesdays

Journal Club Topics and (Presenter) Academic Year 1997-1998


(Ms. Nohelia Canales)


(Mr. Jonathan Hoyne)


(Mr. Kurt Krummel)


(Ms. April Blajeski)


(Dr. Jill Reiter)


(Mr. Jonathan Hoyne)


(Mr. Kun Xu)


(Ms. Susan Barrett)


(Dr. Cecelia Boardman)


(Mr. Eric Calhoun)

(Ms. Colleen Schehl)


(Mr. Andrew Danielson)


(Ms. Gwen Lomberk)

12/12 Special Seminar: The Findling Lecture. "Tumor Angiogenesis"
(Dr. Judah Folkman)
Biology of Breast Cancer Program Extramural Training Support

<table>
<thead>
<tr>
<th>Grant #</th>
<th>P.I.</th>
<th>Source</th>
<th>Term</th>
<th>Total Direct</th>
</tr>
</thead>
<tbody>
<tr>
<td>T32 CA 75926</td>
<td>Salisbury, J.L.</td>
<td>NCI</td>
<td>7/1/98-6/30/03</td>
<td>$805,414</td>
</tr>
<tr>
<td>NRSA</td>
<td>Baines, J.E.</td>
<td>NIH</td>
<td>9/15/97-5/31/02</td>
<td></td>
</tr>
<tr>
<td>NRSA</td>
<td>Canales, N.D.</td>
<td>NIH</td>
<td>5/1/97-5/1/02</td>
<td></td>
</tr>
</tbody>
</table>

Biology of Cancer: A Predoctoral Training Program

Novel Immunotherapeutic Approaches to Cervical Cancer

Research Training in Breast and Ovarian Cancer

Program Trainees

<table>
<thead>
<tr>
<th>Name</th>
<th>Date Entered</th>
<th>Research Mentor</th>
<th>Qualifying Examination</th>
<th>Thesis Defense</th>
</tr>
</thead>
<tbody>
<tr>
<td>Baines, Jonathan</td>
<td>6/3/96</td>
<td>Dr. Persing</td>
<td>passed</td>
<td></td>
</tr>
<tr>
<td>Calhoun, Eric</td>
<td>7/7/97</td>
<td>(rotations)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Canales, Nohelia</td>
<td>1/6/97</td>
<td>Dr. Gendler</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Eley, Gregory</td>
<td>7/8/96</td>
<td>Dr. James</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Holmen, Sheri</td>
<td>9/5/95</td>
<td>Dr. Federspiel</td>
<td>passed</td>
<td></td>
</tr>
<tr>
<td>Johnson, Julie</td>
<td>7/8/96</td>
<td>Dr. Maihle</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Lomberk, Gwen</td>
<td>8/6/97</td>
<td>Dr. Smith</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ritland, Steve</td>
<td>7/1/96</td>
<td>Dr. Gendler</td>
<td>passed</td>
<td>7/98</td>
</tr>
<tr>
<td>Rogers, Michael</td>
<td>8/16/93</td>
<td>Dr. Strehler</td>
<td>passed</td>
<td></td>
</tr>
<tr>
<td>Schehl, Colleen</td>
<td>8/4/97</td>
<td>Dr. Couch</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Xu, Kun</td>
<td>6/18/97</td>
<td>Dr. Prendergast</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Melissa Adriance</td>
<td>8/1/98</td>
<td>(pending entry)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Jessica Faupel</td>
<td>7/1/98</td>
<td>(pending entry)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Shannon Myers</td>
<td>6/1/98</td>
<td>(rotations)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Denise Walters</td>
<td>6/1/98</td>
<td>(rotations)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

"Strain dependent modulation of mammary tumor latency and phenotype in C-erbB2 transgenic mice."
Student Highlights:

Steve Ritland (Dr. Gendler’s laboratory) has completed all track requirements. He is nearing completion of his thesis work, and he will to defend his thesis July 10 of 1998. Steve’s thesis project includes a study entitled: “Strain dependent modulation of mammary tumor latency and phenotype in C-erbB2 transgenic mice.” He is presently exploring postdoctoral positions in several high caliber cancer research laboratories.

Sheri Holmen (Dr. Federspiel’s laboratory) has completed all track course requirements and her qualifying examination. Her research on retroviral vectors and soluble receptors is progressing well. Sheri is also an active member of the Mayo Graduate School Education Committee; during January she conducted an extensive survey and analysis of graduate student opportunities for extramural funding for the Mayo Graduate School. Sheri completed the AACR workshop on “Histopathology of Neoplasia” during the summer of 1997.

Mike Rogers (Dr. Strehler’s laboratory) has completed his qualifying examination and course requirements, and has recently identified CLP-binding proteins using the yeast two hybrid screen. Completion of this work will complement his earlier studies on CLP expression in human breast tumors, and these studies will serve as a basis for his thesis. Mike completed the AACR workshop on “Histopathology of Neoplasia” during the summer of 1995.

Jonathan Baines (Dr. Persing’s laboratory) has completed his course requirements and qualifying examination and has recently begun his thesis studies on the utility of HPV antigens as potential cervical cancer vaccines. Jon was awarded an NIH NRSA Fellowship based on his research proposal on “Novel Immunotherapeutic Approaches to Cervical Cancer”, and he presented an overview of this project in the Tumor Biology Interest Group (TBIG) in March. Jon is an active member of the Training Program Education Committee

Nohelia Canales (Dr. Gendler’s laboratory) is completing her final coursework and is just initiating her thesis research on Muc-1 expression in normal tissue and mammary tumors using the transgenic and knockout mouse models. Nohelia was awarded an NIH NRSA fellowship based on training in the Tumor Biology Program, and recently received an AACR Travel Award for her participation in this year’s AACR meeting.

Greg Eley (Dr. James’ laboratory) is completing his coursework and will take his qualifying examination during the Summer Quarter. He has recently submitted a first author manuscript for publication describing a novel transcript that is co-amplified with the ErbB1 gene. Greg is enrolled in the Genome Analysis course to be held at Cold Spring Harbor Laboratory and the Medical and Experimental Genetics course to be held at the Jackson Laboratory this summer.

Julie Johnson (Dr. Maihle’s laboratory) has completed her coursework and will take the qualifying examination during the Summer Quarter. Her thesis project is on track and she has completed her first thesis advisory committee meeting. Julie’s thesis project is a study of alternate erbB signalling pathways. Julie completed the AACR workshop on “Histopathology of Neoplasia” during the summer of 1997. Julie also has served as the student representative to the Graduate School’s Admissions committee this past year.

THE FACULTY AND THEIR RESEARCH
Tumor Biology Program

Robert T. Abraham, Associate Professor; Ph.D., Pittsburgh, 1981. Signal transduction; cell-cycle regulation; leukemogenesis.


Margot P. Cleary, Visiting Scientist; Ph.D., Columbia, 1976. Breast cancer; obesity; nutrition.

Fergus J. Couch, Assistant Professor; Ph.D., University College Cork, Ireland, 1992. Identification and characterization of genes involved in familial and sporadic breast and ovarian cancer development. Functional analysis of the BRCA2 breast and ovarian cancer predisposition gene.

Chella S. David, Professor; Ph.D., Iowa State, 1966. Immunogenetic aspects of immune response, with emphasis on the major histocompatibility complex class II la genes and T-cell receptor gene.

Gordon W. Dewald, Professor; Ph.D., North Dakota, 1972. Cytogenetics and molecular cytogenetics of congenital disorders and hematologic malignancies.

Richard L. Ehman, Professor; M.D., Saskatchewan, 1979. Magnetic resonance imaging.

Charles Erlichman, Professor; M.D., Toronto (Canada), 1974. Pharmacology of drugs used in cancer therapy.

Mark J. Federspiel, Assistant Professor; Ph.D., Michigan State, 1987. Retroviral vectors; antiviral strategies; molecular medicine.

Lorraine A. Fitzpatrick, Professor; M.D., Chicago, 1980. Prostate cancer metastatic to bone; skeletal calcification; steroid regulation of metastatic disease.

*Sandra J. Gendler, Associate Professor; Ph.D., USC, 1984. Tumor cell biology; mucins in cancer and cystic fibrosis.

Michael J. Getz, Professor; Ph.D., Texas at Houston, 1972. Molecular biology of peptide growth factors; biology of tissue factor in tumorigenesis.


C. David James, Associate Professor; Ph.D., Wright State, 1986. Cancer genetics; cell cycle regulation.

Diane F. Jelinek, Assistant Professor; Ph.D., Texas Southwestern Medical Center, 1985. Cytokine-mediated signaling and gene expression in normal and malignant human B lymphocytes.


Larry M. Karnitz, Assistant Professor; Ph.D., Iowa, 1989. Signaling mechanisms of oncogenes and hemopoietic growth factors; molecular radiobiology.

Scott H. Kaufmann, Associate Professor; M.D./Ph.D., Johns Hopkins, 1981. Pharmacology of topoisomerase-directed antineoplastic agents; apoptosis; resistance to anticancer drugs.

Paul J. Leibson, Associate Professor; Ph.D., 1981, M.D., 1979, Chicago. Tumor immunology; lymphocyte activation; antiviral immunity.

Vanda A. Lennon, Professor; M.B.B.S., Sydney (Australia), 1966; Ph.D., Melbourne (Australia), 1973. Immunobiology of autoimmunity and cancer; ionic channel protein antigens in human neoplasms of lung, ovary, and breast (carcinomas), and thymic epithelium (thymoma).

Edward B. Leof, Associate Professor; Ph.D., North Carolina, 1982. Regulation of cellular proliferation; genetics of pneumocystis carinii.

Ricardo V. Lloyd, Professor; M.D./Ph.D., Wisconsin, Madison, 1975. Endocrine tumor biology, especially pituitary and thyroid.
John A. Lust, Assistant Professor; M.D./Ph.D., Boston University, 1983. Role of IL-6 and IL-6R in pathogenesis of multiple myeloma; detection of minimal residual disease in myeloma transplant patients by PCR.

L. James Maher, Associate Professor; Ph.D., Wisconsin, 1988. Nucleic acid biochemistry; triple helix DNA.

Nita J. Mahrle, Associate Professor; Ph.D., Yeshiva (Einstein), 1983. Molecular basis of cancer; human breast, ovarian, and prostate carcinomas; gliomas.

David J. McKeon, Professor; Ph.D., Johns Hopkins, 1972. Signaling and gene transcription events in T helper lymphocytes; MHC class II protein transport.

Michael J. McManus, Assistant Professor; M.D., Georgetown, 1983. Molecular pediatric oncology; growth factor receptors; tyrosine kinase signal transduction pathways.

Mark A. McNiven, Associate Professor; Ph.D., Maryland, 1987. Cytoskeletal dynamics in mammalian cells; molecular basis of cellular migration during metastasis; vesicle-based transport in epithelial cells.

L. Joseph Melton, III, Professor; M.D., LSU, 1969. Chronic disease epidemiology.

Heidi Nelson, Associate Professor; M.D., Washington (Seattle), 1981. Colorectal cancer; immunotherapy.


Dennis J. O’Kane, Assistant Professor; Ph.D., SUNY at Stony Brook, 1979. Telomerase activity as a diagnostic marker for cancer; translational research on new tumor markers.

David H. Persing, Associate Professor; M.D./Ph.D., California, San Francisco, 1988. Precise promoter mutations in hepatic tumors; immunogenetic determinants of chronic papillomavirus infections and cervical cancer; association of chronic infections with lymphoproliferation.

Mark R. Pittelkow, Professor; M.D., Mayo, 1979. EGF-related growth factor/receptor function: epidermal keratinocyte and melanocyte regulation of growth and differentiation.

Karl C. Podratz, Professor; M.D./Ph.D., St. Louis, 1974. Molecular prognostic determinants in gynecologic malignancies.

Gregory A. Poland, Professor; M.D., Southern Illinois. Expertise in vaccine development and evaluation, adjuvant development and evaluation; vaccine antigen processing and HLA presentation; and vaccine immunogenetics.

Franklyn G. Prendergast, Professor; M.B.B.S., West Indies, 1968; Ph.D., Minnesota, 1977. Fluorescence spectroscopy; protein structure and dynamics; biochemistry and bioluminescence.

Corey Raffel, Associate Professor; M.D./Ph.D., California, San Diego, 1980. Pediatric neuro-oncology; gene therapy and cancer.

Jeffrey L. Salisbury, Professor; Ph.D., Ohio State, 1978. Cell cycle control; centrosomes; mitotic spindle poles; breast cancer.

David J. Smith, Professor; Ph.D., Wisconsin, 1978. Chromosomal fragile sites; molecular genetics of cancer development.

Thomas C. Spelsberg, Professor; Ph.D., West Virginia, 1967. Steroid action on early (c-myc) gene transcription, steroids and TGF-β action on bone cell functions, and early gene expression.

Emanuel E. Strehler, Associate Professor; Ph.D., ETH Zurich (Switzerland), 1981. Intracellular Ca²⁺ homeostasis and signaling; molecular mechanisms of disease.

Stephen N. Thibodeau, Professor; Ph.D., Washington (Seattle), 1979. Cancer genetics; colon and prostate cancer.


Raul Urrutia, Assistant Professor; M.D., Cordoba (Argentina), 1987. Cell differentiation.

Peter J. Wettstein, Professor; Ph.D., North Carolina at Chapel Hill, 1977. Role of minor histocompatibility antigens in allograft rejection.


Lester E. Wold, Professor; M.D., Chicago, 1977. Immunocytochemistry; bone tumors and tumor-like conditions; breast diseases.

Charles Y-F. Young, Assistant Professor; Ph.D., Brigham Young, 1984. Calpain inhibitor-induced apoptosis in human prostate adenocarcinoma cells.

*Scottsdale campus.
Research and Resource Facility Information Sheet

Mayo Research Facilities

The research training environment and research facilities at Mayo are exceptional. There are approximately 150 career scientists and clinicians engaged in research at Mayo, along with their associated research personnel. The annual Mayo research budget is approximately $140 million, which is derived in roughly equal proportions from extramural and Mayo Foundation funds. In 1997, approximately two-thirds of Mayo's extramural support was provided through grants and contracts awarded through the National Cancer Institute. Thus, research at Mayo is both well supported by the institution and recognized for its excellence through a competitive extramural peer-review process. Internal research resources are administrated by the Mayo Foundation Research Committee (Rochester) and the Mayo Cancer Center Advisory Committee.

Mayo Core Facilities

The Mayo Foundation supports most major research instrumentation needs, and, in addition, provides resources required for shared equipment core facilities. These include 10 'core' research laboratories that serve as major scientific and educational resources for students and their respective laboratories. Each Shared Mayo Research Resource is overseen by a Mayo staff investigator with expertise in the appropriate area in addition to a staff of professional research assistants who are expert in the use of instrumentation specific for the particular facility. Faculty and students can access these resources at many levels. An 'occasional' use or analysis can be performed by the facility staff. Students whose research projects require more extensive use of a facility can be trained by the technical staff in a given facility and, thereby, develop technical expertise themselves. Thus, students can truly follow their research question wherever it may take them, even if it goes beyond the technologies available in their mentor's laboratory.

Mayo Clinic Shared Research Resource Facilities.

<table>
<thead>
<tr>
<th>Analytical NMR Resource</th>
<th>Biomedical Imaging Resource</th>
</tr>
</thead>
<tbody>
<tr>
<td>Electron Microscopy Resource</td>
<td>Flow Cytometry/Optical Morphometry</td>
</tr>
<tr>
<td>Mass Spectroscopy Resource</td>
<td>Mathematical Methods Resource</td>
</tr>
<tr>
<td>Molecular Biology Resource</td>
<td>Protein Sequencing/Peptide Synthesis</td>
</tr>
<tr>
<td>Radioimmunoassay Resource</td>
<td>Research Computing Facility</td>
</tr>
<tr>
<td>Pharmacology Core</td>
<td>Cytogenetics Core</td>
</tr>
<tr>
<td>Cancer Biostatistics Core</td>
<td>Pathology Core</td>
</tr>
</tbody>
</table>
Analytical nuclear Magnetic Resonance Facility (NMR) - (625-341-9800)
Dr. S. Macura, Director (4-6937). Analytical Nuclear Magnetic Resonance Facility supports applications of high-field nuclear magnetic resonance (NMR) spectroscopy to biochemical and biological systems, in particular, structure-function studies of biomacromolecules. The facility contains high-field state-of-the-art NMR spectrometers and computers for processing and analyzing NMR data. The facility is staffed by experts in various fields of NMR experimentation who carry out core research and development and spend part of their time consulting with and training users. Facility staff may collaborate with users who have projects of mutual interest.

Biomedical Mass Spectrometry Facility (BMSF) - (625-025-9800)
Dr. S. Naylor, Director (4-5220; pager 4-6091) Additional Contact Person - Dr. A. J. Tomlinson (4-1040; pager 4-7506). The Biomedical Mass Spectrometry Facility is a resource for Mayo investigators to isolate and structurally characterize a wide range of biological relevant compounds. The activities within the BMSF are focused on four specific services; 1) Consultation and education involving the isolation and characterization of biopolymers and metabolites; 2) Routine molecular weight service of molecules including biopolymers, metabolites, and lipids; 3) CE and HPLC separation of complex mixtures; and 4) long-term collaborative interactions with Mayo investigators involving the isolation and structural characterization of biopolymers and metabolites from complex biological matrices.

Electron Microscopy Core Facility (EMCF) - (625-347-9800)
Dr. J. L. Salisbury, Director (4-3326) Additional Contact person - Jon Charlesworth (4-3148 or 4-1616). The Electron Microscopy Core Facility provides specimen preparation, microscopy, and photography services to investigators from both clinical and basic science laboratories. In addition to standard transmission and scanning electron microscopy, the facility performs x-ray probe microanalysis and immuno-gold labeling procedures.

Flow Cytometry/Optical Morphology Core Facility - (625-350-9800)
Dr. P. J. Leibson, Director (4-4866) Additional Contact Person - Jim Tarara (127-0183 or 4-4241). Instrumentation of flow cytometry/cell sorting is provided for high speed cell analysis and cellular fractionation. Samples are prepared in the investigator’s laboratory and can be run by core facility personnel. Complete image analysis software is available. Other instruments and services provided include confocal microscopy and computer controlled ratio imaging.

Immunohistochemical Laboratory Core Facility (ICl) - (560-091-9800)
Dr. G. G. Klee, Director (4-8213; pager 4-7406) C. M. Schimek, Supervisor (5-4772; pager 127-4046) Technical Coordinator - Don Heser (5-7855). The Immunohistochemical Core Laboratory (ICl) is a Mayo Research Committe core facility for providing laboratory testing at minimal cost to Mayo researchers. When capacity is available, testing is also provided for investigators outside Mayo on a collaborative basis and for MML clinical trials clients. The ICL staff is actively involved in new assay development and improvement of current assay methodology. The laboratory is located at 5-223 Joseph, Saint Mary’s Hospital, next to the GCRG. Requests for ICL service are prioritized based on volume, number of protocols requiring the test(s), availability of an alternative source for testing and development effort required. First priority for laboratory testing is given to GCRG protocols and other NIH funded protocols.

Material and Structural Testing Resource (MSTR) - (409-103-9900)
Dr. K-N An, Director (4-2589) Additional Contact Person - Tricia Neale (4-1460). The MSTR is a facility designed to thoroughly assess the mechanical characteristics of a multitude of engineering and biologic materials. The facility is capable of developing custom tailored solutions to analyze a wide array of problems utilizing both experimental and theoretical approaches.
Biomathematics Resource - (640-151-9800)
Drs. Z. Bajzer (4-8584) and A. Manduca (4-8163), Directors. The Biomathematics Resource was established to provide advanced mathematics and computing expertise to research programs and to the Mayo community as a whole. The facility is staffed by experts in mathematical modeling, image processing, and machine learning algorithms who carry out research and development, collaborate, and provide consulting. Services provided include: 1) collaboration with Mayo investigators and/or clinicians on specific projects which require advanced mathematical and/or computational methods, 2) development of new methodologies, 3) consultation and education in mathematical and computational methods, and 4) custom development of software for specific needs.

Mayo Biomedical Imaging Resource (BIR) - (640-193-9800)
Dr. R. A. Robb, Director (4-4937; pager 4-7744) Additional contact persons: Mahlon Stacy (4-6174), Dennis Hanson (4-6103) Jon Camp (4-3870). The Mayo Biomedical Imaging Resource (BIR) is dedicated to the advancement of research in the biomedical imaging and visualization sciences. The BIR provides expertise, advanced technology and comprehensive software related to biomedical imaging and scientific visualization, including image display, processing and analysis; image databases; virtual reality; computer graphics; video animation; computer workstations; computer networks; and computer programming. The BIR also provides specialized services in computer maintenance and back-up, custom software development and imaging system design.

Mayo Protein Core Facility - (625-288-9800)
Dr. D. J. McCormick, Director (4-4992) Additional Contact Person: Benjamin Madden (4-2457). The Mayo Protein Core Facility is a resource which provides services and methods related to the isolation, characterization, and analysis of proteins. Current services provided by the facility include: 1) N-terminal amino acid sequencing; 2) Carboxyl (C) terminal sequencing; 3) amino acid analysis; 4) protein isolation by conventional liquid chromatography (FPLC); 5) solid phase peptide synthesis at both small and large scales; 6) production of peptide immunogens by conjugation to protein carriers and MAP resins; 7) peptide nucleic acid synthesis (PNA); 8) proteolysis and peptide mapping. 9) peptide purification by reverse phase HPLC; 10) analysis of protein sequences from data bases for structural homology, synthetic peptide design, and molecular modeling For a complete description of the facility and its services use the local Web page at the site address: http://www-rcf.mayo.edu/protein/

Molecular Biology Core - (625-205-9800)
Dr. B. C. Kline, Director (4-7489); Additional Contact persons: oligonucleotides - Maryjane Doerge (4-8186), DNA sequencing - Bruce Eckloff (4-3797), training - Ross Aleff (4-3794). The Molecular Biology Core synthesizes oligonucleotides, performs semi-automated, fluorescence-based DNA sequencing, and provides laboratory training in molecular biological techniques. For more information, use the local Web page http://www-rcf.mayo.edu/molecular/

Research Computing Facility (RCF) - (653-000-9800)
Dr. R. A. Ghanbari, Director (4-1817) Additional contact person: Robert Bleimeyer (127-4656). The goal of the Research Computing Facility is to guide, coordinate, and enable the effective use of computing and information management technologies by the Mayo investigator. Resources available consist of a staff of computing professionals well versed in addressing the needs and interests of research using hardware and software tailored to the needs of the investigator. Five integrated services are available which include: 1) collaborative consulting; 2) education on the application of computing technology; 3) management and coordination of computing resources; 4) services for desktop computers in cooperation with other Mayo service groups; and 5) shared hardware, software, and reference data resources.
June 26, 1998

Nita J. Maihle, Ph.D.
Department of Biochemistry and Molecular Biology
Mayo Foundation
200 First Street SW
Rochester, Minnesota 55905

Dear Nita:

This is a response to your request for an evaluation of your predoctoral training program in breast cancer research supported by a USAMRDC training grant. I do have significant familiarity with predoctoral training programs. I have been chair of a basic science department and thus responsible for a predoctoral training program for the past 19 years and director of a National Cancer Institute training grant supporting predoctoral trainees for almost the same length of time.

From my review of the submitted materials, it is clear that you have successfully used this award to initiate a multidisciplinary training program with a particular focus in breast cancer. The administrative structure, governance and didactic course work is appropriate for training scientists to pursue breast cancer research. You have a very competitive, interactive faculty representing sufficient breadth and depth in areas relevant to breast cancer research. The best measure of a training program is the long term success of its trainees. This is difficult to evaluate in such a brief time after its initiation. However, it appears that you have been successful in recruiting excellent students, most of whom are progressing well in the program. I have had the opportunity to meet one of your trainees, Steve Ritland, who is nearing completion of his predoctoral training. He is truly impressive and shows promise for developing into an outstanding independent investigator. If you are able to produce a few more as impressive as Steve Ritland, your program will be an unqualified success.

Overall, you program has been successful in accomplishing the goals of the original US Army Breast Cancer Integration Panel of which I was a member. That goal was to increase the number of high quality scientists working on the breast cancer problem. Another very important measure of the success of your program is the fact that you have been able to continue the program by obtaining a training grant from the National Cancer Institute. I congratulate you and your colleagues on your accomplishments.

Sincerely,

[Signature]

Harold L. Moses, M.D.
B.F. Byrd, Jr. Professor of Oncology
Director, Vanderbilt Cancer Center
Professor and Chair, Department of Cell Biology
Dear Nita:

I just discovered that I hadn't sent my very short write-up of your program to you. I think in the confusion of a trip (I do a lot of work in the planes!), I must have left it behind. Here is even a shorter version of what I wrote:

The program in Biology of Breast Cancer is timely and competitive. The integration of basic and clinical research is ultimately what is needed in all these programs. The breadth of the background of the faculty is refreshing and would, if used properly, provide an appropriate base of knowledge for the trainees. The resources would be the envy of other national programs!

The curriculum, as stated, is excellent. The inclusion of developmental biology, principles of cell and tissue design and ethics as well as the more standard courses should be commended. However, it would have been helpful to have included the CV of the faculty and a history of their training students and fellows.

Sincerely,

Mina

---

Appendix M

Printed for maihle.nita@mayo.edu (Nita Maihle Ph.D.)

The Biology of Breast Cancer Training Program of the Mayo Clinic is a predoctoral training program in tumor biology. The focus of the program is to provide an educational environment that stimulates excellence in scientific thought and training while simultaneously providing exposure to all of the major fields of study relevant to tumor biology with a particular emphasis on breast and other women’s cancers. The goals of the program are four-fold: First, to provide trainees with a solid and uniquely multidisciplinary knowledge base in the biology of cancer. Second, to guide the development of each individual trainee so that they achieve their fullest academic and research potential. Third, to aid trainees in establishment of their professional network of peers and colleagues in the field of cancer research. And, fourth, to stimulate new working alliances between students, fellows, and staff participating in cancer related research, education, and clinical endeavors at the Mayo Clinic and within the Mayo Cancer Center. Students participate in laboratory-based research, as well as in a formal tumor biology curriculum that integrates current concepts in cell growth control with the natural history of human tumors. The Biology of Breast Cancer Training Program courses establish a solid foundation in the biology of breast cancer and share several distinguishing features and innovative strategies for teaching which enhance learning and student/faculty participation. In particular, active involvement of students in the learning and discovery process, in information processing, and in the application of information to problems requires that students be accountable for learned information on an immediate and ongoing basis. Problem-centered learning also puts learning into context and facilitates learning transfer. These sessions allow students to organize and categorize information into meaningful units and to “discover” novel relationships and extract and assimilate important points in an interactive and participatory manner. Supported by the US Army Medical Research and Materiel Command (DAMD17-94-J-4116) and the Mayo Clinic Foundation.

Type font preferred: New Times Roman 12 point • Justification on right and left is preferred

Jeffrey L. Salisbury, Ph.D.  
(Signature of PRESENTER)

Gugg-14, Mayo Clinic, Rochester, MN 55905  
(907) 284-3326

As the 1st AUTHOR of this abstract, on behalf of all the authors, I hereby transfer its copyright to the American Association for Cancer Education

Signature of 1st Author  
Jeffrey (S Salisbury  
E-mail address Salisbury@mayo.edu

Gugg-14, Mayo Clinic, Rochester, MN 55905  
(507) 284-3326

(address)  
(city and state, ZIP code, country)  
(area code, telephone number)
List of Personnel Receiving Pay From this Effort
(1994-1998)

Margaret A. Adelsman
Jonathan E. Baines
Nohelia D. Canales
Gregory D. Eley
Sheri L. Holmen
Julie L. Johnson
Sharon L. Jones
Gwen Lomberk
Steve R. Ritland
Michael S. Rogers
Colleen M. Schehl
Kun Xu