GENES AND PROTEINS THAT CONTROL THE SECRETORY PATHWAY

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Introduction

George Palade shared the 1974 Nobel Prize with Albert Claude and Christian de Duve for their pioneering work in the characterization of organelles interrelated by the process of secretion in mammalian cells and tissues. These three scholars established the modern field of cell biology and the tools of cell fractionation and thin section transmission electron microscopy. It was Palade’s genius in particular that revealed the organization of the secretory pathway. He discovered the ribosome and showed that it was poised on the surface of the endoplasmic reticulum (ER) where it engaged in the vectorial translocation of newly synthesized secretory polypeptides (1). And in a most elegant and technically challenging investigation, his group employed radioactive amino acids in a pulse-chase regimen to show by autoradiographic exposure of thin sections on a photographic emulsion that secretory proteins progress in sequence from the ER through the Golgi apparatus into secretory granules, which then discharge their cargo by membrane fusion at the cell surface (1). He documented the role of vesicles as carriers of cargo between compartments and he formulated the hypothesis that membranes template their own production rather than form by a process of de novo biogenesis (1).

As a university student I was ignorant of the important developments in cell biology; however, I learned of Palade’s work during my first year of graduate school in the Stanford biochemistry department. Palade was a close friend of my graduate advisor Arthur Kornberg, who won the Nobel Prize in 1959 for his discovery of DNA polymerase, the first enzyme found to take its instructions from a DNA template (2). At first glance Kornberg and Palade had little in common. Palade was a classical anatomist and physiologist who used the electron microscope as his primary tool of analysis. Kornberg was a classical biochemist who cared deeply about the chemistry of life, which he probed exclusively through the study of pure enzymes. However, in the late 1960s as the study of DNA synthesis began to focus on the possible role of a membrane surface in organizing the segregation of replicating chromosomes, Kornberg took a keen interest in membrane biochemistry and in 1969, the year before I started graduate school, Kornberg traveled to several laboratories of membrane biologists including Palade’s, who was then at The Rockefeller University. On return to Stanford, Kornberg turned his attention to membrane enzymes in the hope that a membrane surface may provide a crucial link to the problem of DNA replication. Just then, in the summer of 1969, the field of DNA replication was shaken with the discovery by John Cairns, then Director of the Cold Spring Harbor laboratory, that Kornberg’s DNA polymerase was not required for chromosome replication. I visited Cold Spring Harbor that summer and was swept up in the excitement of the Cairns isolation of an E. coli pol1 mutant, lacking polymerase activity, but which grew
normally and yet was sensitive to UV irradiation, a clear sign that the classic polymerase could not be the enzyme responsible for replication but instead played a role in DNA repair (3).

The power of genetics and biochemistry combined

Kornberg was a dominant figure with a powerful personality and intellect. His focus on enzyme chemistry shaped a generation of students of DNA enzymology, including several former postdoctoral fellows and associates who joined him to form the core of what was to become the preeminent biochemistry department in the country at Stanford Medical School, where he moved from Washington University, St. Louis in 1959, the year in which he was awarded his Nobel Prize. With the pure DNA polymerase, Kornberg proved that it took its instructions from a template strand and copied DNA in an antiparallel direction, as predicted from the Watson-Crick model of the DNA duplex (4). The most persuasive evidence that it could be the replication enzyme came in 1967 with the demonstration that polymerase alone copied the circular single strand template of the bacteriophage \( \phi \)X174 to make a complementary strand, which then also served as a template to make infectious viral strand DNA (5,6). Thus the enzyme could faithfully take instructions from a template of around 5500 nucleotides and form, essentially error-free, a complement to reproduce the viral infectious cycle in a living cell.

However, several features of the polymerase left some investigators skeptical that it was the authentic replication enzyme. DNA chain elongation by the polymerase was quite slow in comparison to the progression of a chromosome replication fork. The enzyme had properties that suggested an ability to repair DNA damage, for example in the excision of thymine dimers on DNA isolated from cells exposed to UV light (7). Another puzzling feature was the requirement for a complementary oligonucleotide that forms a short duplex, which serves to launch the polymerase from a 3’OH provided by the primer (5). Nonetheless, an enzyme much like the \( E. coli \) polymerase is encoded by the T4 bacteriophage and in that case phage mutations in the polymerase gene show that it is clearly required for viral chromosome replication (8).

Quite independently, bacterial geneticists found genes essential for chromosome replication by the isolation and characterization of temperature sensitive (ts) mutations that arrest DNA synthesis in cells warmed at 42C (9,10). Cells carrying the \( dna \) mutations can grow at 30 C but cease growing at 42 C. The “\( dna \)” genes thus represented candidates for the authentic replication machinery quite distinct from the \( pol1 \) gene identified as non-essential in the Cairns mutant. A grand union of the genetics and biochemistry first developed through a twist of fate with the discovery by Tom Kornberg, Arthur’s middle son, then a graduate student in the laboratory of Malcolm Gefter at Columbia University, of another replication activity detected in lysates of the Cairns mutant (11). Gefter and Kornberg went on to discover that the authentic polymerase is encoded by the \( dnaE \) gene, one of the approximately half-dozen genes then known to be required for chromosome replication (12).

In 1970 I joined Arthur’s lab powerfully influenced by the two strands of investigation, enzymology as practiced by the Kornberg school, and molecular biology and genetics, as best described in James Watson’s textbook \textit{Molecular Biology of the Gene} (13). I had read and reveled in the details in the first
edition of this book when I was a freshman at UCLA, and although I was drawn to the Kornberg approach for graduate training, I was mindful that genetics and cellular physiology must inform the biochemistry.

A stunning precedent for the value of a combined genetic and biochemical approach came from the pioneering work of Robert Edgar, a bacterial geneticist who dissected the process of T4 phage assembly with the isolation of mutations in the genes that encode subunits of the phage coat (14), and William Wood, a new faculty colleague of Edgar’s at Cal Tech. Wood had trained with Paul Berg, a former post-doctoral fellow of Kornberg’s and then a colleague in the new Biochemistry Department at Stanford. At Cal Tech in the fall of 1965, Wood and Edgar joined forces to perform one of the classic experiments in molecular biology. Edgar had found that some of the viral coat mutants accumulated incomplete viral heads and tails within infected cells. Edgar used the standard cis-trans genetic complementation test, first developed by Seymour Benzer for the characterization of phage rII genes (15), to characterize the genes involved in T4 phage morphogenesis. Wood imagined that biochemical complementation might be achieved by mixing extracts of different phage assembly mutant-infected cells. Indeed, starting with separate extracts that had essentially no detectable infectious virions as assayed by the phage plaque test, Edgar and Wood found that mixing lysates of genetically complementing mutants (i.e. biochemical complementation) produced a thousand-fold increase in infectious particles (16). The team went on to identify functional assembly intermediates and to map the pathway of virus assembly. Clearly, this approach had the potential to dissect complex pathways and to reveal molecular details that might not otherwise be elucidated by a strictly genetic or biochemical analysis.

In 1971, Doug Brutlag, a talented graduate student in Arthur’s lab, discovered that the conversion of the M13 phage single stand circle to the double strand replicative form was blocked in infected cells by an inhibitor of the transcription enzyme RNA polymerase, this in spite of the fact that no viral or host gene expression is required at the first stage of chromosome replication. Brutlag and Kornberg suggested that RNA polymerase might provide the missing primer to initiate the growth of a DNA chain (17). Brutlag then established a replication reaction in a concentrated lysate of uninfected E. coli cells and found that this faithfully reproduced the requirement for RNA polymerase in the conversion of M13 single strand template to the duplex replicative form (18). A similar concentrated extract of E. coli had been developed in the laboratory of Friedrich Bonhoeffer in Tübingen Germany, and found by one of Bonhoeffer’s postdoctoral fellows, Baldomero Olivera, to be capable of replicating φX174 single strand circular template (20). Both concentrated lysates contained membranes and cytosolic proteins and it seemed possible that the reaction would require a membrane contribution. However, at the same time, Bruce Alberts, then at Princeton University, found that soluble cytosolic lysates of T4 phage infected cells replicated T4 DNA and applying the logic of Wood and Edgar, Jack Barry and Alberts showed biochemical complementation of soluble protein fractions obtained from different T4 replication mutant cells (21). At Stanford, a new postdoctoral fellow in the Kornberg group, William (Bill) Wickner, found that the lysate capable of replicating M13 DNA could be centrifuged to produce a soluble fraction with no loss of replication activity (18). All interest in membranes and DNA replication seemed to evaporate with that result.
I joined the effort initiated by Brutlag and Kornberg, first on the replication of M13 DNA and then using the cell-free reaction Doug Brutlag had developed, I found that ϕX174 double strand formation was insensitive to the drug that blocks the standard RNA polymerase, suggesting perhaps an alternative RNA polymerase for primer synthesis (18, 19). David Denhardt at Harvard University, with whom I had worked for a summer, had reported that ϕX174 double strand formation was dependent on the *E. coli* dnaB gene; thus it seemed possible that the cell-free reaction might provide a functional assay for the purification of the dnaB protein and for the remaining dna proteins. Indeed, it did, and this reaction permitted the detection and fractionation of the full set of *E. coli* chromosome replication proteins (22). One of the Dna proteins, DnaG, was found to catalyze a novel RNA synthesis reaction that provides the primer for ϕX174 as well as for *E. coli* chromosome fork replication. Going forward I was confident that the combined genetic and biochemical approach could prove crucial in the elucidation of other complex cellular processes.

The cell division cycle represented one such complex pathway that was just beginning to be probed by molecular genetic approaches. I was particularly taken by the efforts of Leland Hartwell who had exploited the classical genetic tools available for baker’s yeast, *Saccharomyces cerevisiae*, to probe the essential series of events that lead to yeast cell division. The key was a set of genes, identified by the isolation of *ts* lethal mutations that focused attention on crucial control elements in the progression of the cell cycle (23). Subsequent molecular genetic discoveries by Paul Nurse, Tim Hunt and others illuminated the molecular basis of cell cycle control. Here again, the molecular insights that started with a classical genetic approach proved crucial to the discovery of a protein kinase that controls the decision to initiate the cell division cycle and then acts repeatedly in transitions throughout the division cycle. A billion years of evolution conserved a similar pathway in mammals. From this it seemed most likely that studies on yeast could pave the way for a mechanistic understanding of many, if not all, other essential eukaryotic intracellular processes.

**Investigating biological membranes as a macromolecular assembly**

Although the replication reactions he investigated were not directly connected to membranes, Kornberg remained interested in the problem of how to purify membrane enzymes and thus he was eager to welcome an experienced membrane enzymologist, Bill Wickner, who joined the lab as a postdoctoral fellow in 1971. Bill had trained as a medical student with Eugene Kennedy at Harvard Medical School where he lost interest in clinical medicine but gained an abiding passion for biological membranes. He and I shared endless hours in conversation about our work but importantly, I learned a great deal from him about what was or was not known about how membranes are put together. In this context, I read the work of Palade and his associates David Sabatini and Phillip Siekevitz who were then exploring the mechanism of vectorial membrane translocation of secretory proteins as they are made on ribosomes associated with the ER (24, 25). The Stanford biochemistry department was a focal point for visits by all the leading figures in modern biology. I met Hartwell and two other memorable men who represented different approaches to the study of membrane function: Efraim Racker who shared Arthur’s passion for enzymes in his dissection of the mechanism of mitochondrial oxidative phosphorylation and Daniel Koshland, who had exploited a genetic approach pioneered by Julius Adler to probe the mechanism of
bacterial chemotaxis, a process intimately linked to the detection of chemical gradients at the bacterial cell surface.

As I considered my future research career directions, I was motivated by a desire to break away from the field of DNA replication but to appropriate the tools and logic that had propelled the Kornberg group to a successful resolution and reconstitution of the enzymes of the replication process. As I concluded graduate work in 1974, the beginnings of a revolution in genetic engineering and recombinant DNA were just emerging, largely from the work of Stanford biochemists Dale Kaiser and Paul Berg, Stanford microbiologist Stanley Cohen and the UCSF biochemist Herbert Boyer. The tools of molecular cloning were in prospect, thus it was appealing to consider how they may be applied to uncover essential genes in any number of cellular processes.

And yet I was uncomfortable with the frenzy of activity that focused on all things DNA. I did not enjoy the pressure of competing with other laboratories doing the same experiments. I nervously unwrapped each new issue of the Proceedings of the National Academy of Sciences (PNAS) to see if our competitors had beaten us to key discoveries. Basking in the glow of Kornberg’s influence had its advantages, but in facing my own independent career I resolved to strike off in a new direction where I might have the chance to establish my own identity and not be dependent on or overshadowed by Kornberg’s reputation.

In making a choice for future research, I was impressed with the work of S. Jonathan Singer at UC San Diego, particularly with his greatly influential paper on the Fluid Mosaic Model of Membrane Structure (26). Here was a grand synthesis that provided a conceptual framework to think about how a membrane might be constructed. Singer’s lab had assembled tools to explore the topology of membrane proteins using electron microscopy. His associate Kiyoteru Tokuyasu had developed an impressive cryoelectron microscopic approach to the detection of antigens on membranes (27). They had demonstrated that glycans on glycoproteins and glycolipids are asymmetrically displayed on the extracellular surface of red cells and on the luminal surface of the ER membrane, thus fulfilling the prediction that transbilayer movement of hydrophilic proteins and glycans was thermodynamically unlikely, but at the same time explaining how the asymmetry of the plasma membrane may be achieved at the outset of the secretory pathway (28). Singer enjoyed the warm support of my mentors at Stanford so I set off with my bride Nancy, whom I had met through my friendship with Bill Wickner, to join Singer’s lab as a postdoctoral fellow in the fall of 1974.

Singer was so different from Kornberg that I experienced a bit of culture shock while trying to identify a research project of mutual interest. Although he made his career as a physical chemist, he had evolved into a cell biologist focusing on questions of cellular organization. I was keen to use the reconstitution approach of Kornberg to probe some aspect of membrane assembly or endocytosis but Singer pressed me to pursue a morphological study using electron microscopy. Of course it was important to learn a new discipline as well as a different approach so I took up a project to investigate the unusual behavior of neonatal human erythrocytes, which unlike mature red cells are able to internalize antibody or lectin molecules clustered on the cell surface (29). I found the work frustrating, and in spite of Tokuyasu’s patience, my technical skills in thin section electron microscopy left much to be desired. It took two years
to obtain one precious Rh control sample of newborn cord blood. The prospect of a satisfying molecular understanding of this process seemed remote and my dependence on cord blood from the local labor and delivery ward slowed progress to a snail's pace. I was spoiled by my previous experience with microorganisms and the slow and cumbersome approaches then available for work with human samples or even with cultured mammalian cells simply could not compare. So I had time to read and think, which was perhaps the greatest benefit of my postdoctoral years.

Shortly after I started in Singer's lab, the annual meeting of the American Society for Cell Biology (ASCB) convened in San Diego. At the time the ASCB was a small and quite personal organization, much more of a cottage industry than was the larger and more influential American Society for Biological Chemistry (ASBC), subsequently renamed the ASHMB. Palade had just returned from Stockholm to deliver a special lecture to an adoring crowd who rose to a standing ovation at the end of his presentation.

Although I knew then, and learned even more so later, how brilliant and broad Palade was in his scholarship, I came away from the meeting feeling that cell biology had yet to enter the molecular world of biochemical mechanism. Here was an enormously complex pathway of membrane transformation in the secretory pathway and yet not a single protein had been ascribed a specific role in this essential process.

The first crucial breakthrough that delivered Palade's pathway into the molecular era came with the report in 1975 by the Palade protegé Günter Blobel, of a cell-free system that reproduced the initiation and translocation of a secretory precursor protein into the interior of isolated ER membranes. Two papers in the Journal of Cell Biology by Blobel and Dobberstein paved the way to a mechanistic understanding of the link between protein synthesis and the vectorial discharge of secretory proteins through what must surely be a hydrophilic channel protein in the ER (30, 31). Although earlier work by Sabatini and Palade had demonstrated the completion of the translocation event in vitro using rough microsomes isolated from pancreatic tissue, Blobel's breakthrough allowed the entire process to be replicated with the discovery of an essential role for the N-terminal signal peptide in guiding the nascent chain to a special site on the ER membrane. The signal hypothesis and the beautiful work that followed garnered a Nobel Prize for Blobel in 1999.

Singer was quite excited by the Blobel discovery because it supported his view that the establishment of protein asymmetry in the membrane must depend on a special channel in the ER that would convey hydrophilic protein sequences through the hydrophobic bilayer. And yet, Singer remained skeptical that a biochemical reconstitution approach would yield an essential understanding of the process. But to me, this was precisely the way forward, though my own efforts in that direction would await an opportunity to take the initiative.

Increasingly, I believed a unique opportunity lay in the evaluation of plasma membrane assembly in S. cerevisiae and my reading of the literature focused on what was known before 1975, which outside of the work of Gottfried Schatz and Walter Neupert on mitochondrial biogenesis was essentially nil. I read about the organization of the yeast cell surface, particularly at the nascent division site, which had an intriguing intermediate filament ring abutting the cytoplasmic surface of the bud neck membrane and a
unique deposition of chitin in a ring embedded within the cell wall polysaccharide (32,33). Vesicles implicated in secretion were seen by thin section electron microscopy to localize to the cytoplasm of an early cell bud and then to appear near the cytokinesis furrow later in the cell cycle (Fig.1) (34,35). It seemed reasonable to suppose that these vesicles were responsible for secretion and localized plasma membrane assembly. These ideas excited me a great deal more than the tedious work I was doing on human neonatal erythrocytes.

![Figure 1. Thin section of wild-type yeast cell showing endoplasmic reticulum (er), vacuole (va) and secretory vesicles (ves).](image)

**Free speech and free inquiry at UC Berkeley**

Just as I left Stanford on my way to San Diego, I became aware of an Assistant Professor opening in the Biochemistry Department at University of California, Berkeley. Although I had no postdoctoral training, I decided to apply for this position just in case the Berkeley faculty would see my graduate record as an indication of my interests and abilities. Fortunately for me, the first person to whom they offered the job, turned it down, and in a call I will never forget, Michael Chamberlin, a Paul Berg-trained Stanford Biochemistry graduate and then Chair of the Berkeley search committee, conveyed the good news that I had the job. I was so excited that I foolishly accepted over the phone with no further negotiation! And so within the first few months of my postdoctoral training, I had the luxury of planning my future career without the responsibilities of the job.
The Berkeley Biochemistry Department was a perfect place for my interests. Daniel Koshland served as Chair and the faculty included a distinguished group of classical biochemists such as Esmond Snell, Jesse Rabinowitz, Clinton Ballou, Jack Kirsch and Howard Schachman, as well as a group with broader interests in genetics and molecular biology such as Allan Wilson, Stuart Linn, Ed Penhoet, Chamberlin, and Bruce and Giovanna Ames. Jeremy Thorner, a close friend from my Stanford years, had taken up a study of yeast pheromone biology as a beginning faculty member in the bacteriology and immunology department at Berkeley. Ballou was an expert in carbohydrate chemistry with a particular interest in the yeast cell wall. Koshland, whom I had met at Stanford, and Ames were most appealing because they blended genetics and biochemistry in a way that I found compatible with my temperament. I believed that my future colleagues would allow me the freedom to explore a new direction quite different from my graduate or postdoctoral work.

In the remaining time of my postdoctoral work, I completed a project and published a paper but all my thoughts were directed to my future at Berkeley. Of course, I had no experience with yeast and knew essentially no genetics so I planned to spend three weeks at the yeast genetics course offered at Cold Spring Harbor and taught by Fred Sherman and Gerald Fink. Sherman and Fink were master geneticists and were able to draw on all the major figures in the yeast community who dropped by to teach and remain for a day or two. It was a thrill to meet Lee Hartwell and to share my thoughts about how yeast cells may grow by vesicle traffic. On the other hand, like with thin section electron microscopy, my skills in yeast tetrad dissection were inadequate. I believe I held the record for fewest tetrads dissected until several years later when James Rothman took the course.

How to study secretion in yeast

As the time approached for the move to Berkeley, I worked feverishly to craft an NIH grant proposal that included a range of ideas on how to study secretion and membrane growth in yeast. Published evidence suggested that secretion was localized to the bud portion of the dividing cell but there were no tools available to study the localization of a newly synthesized plasma membrane protein. My ideas were fanciful but in the cold light of day, the NIH reviewing panel found my experience inadequate (I had no preliminary data) and my ideas unproven. The rejection was crushing and my colleagues must have wondered if their gamble on me was about to crash. Adding insult to injury, I was denied a Basil O’Connor starter grant from the March of Dimes where the interviewer found me intelligent but regretted that I had not proposed to work on cell division in Lesch Nyhan syndrome! Fortunately, the NSF, and friendly reviews from Lee Hartwell and Susan Henry, a young yeast geneticist who studied phospholipid regulation, rescued me with a grant in the princely amount of $35,000 for two years. With this and a small internal University grant, a modest effort took shape.

What to do first? In the fall of 1976, two graduate students joined my lab: Janet Scott and Chris Greer. Janet had transferred from another lab so she had to find something that would work quickly. I felt that in order to study the yeast plasma membrane it would be necessary to have a clean way to remove the cell wall avoiding the use of crude snail gut enzymes, Glusulase, that were used to convert cells to spheroplasts. Another lytic enzyme secreted by a soil bacterium, *Oerskavia xanthineolytica*, seemed a
good source to begin a purification effort. Janet perfected the conditions of induction and purification of an enzyme we called lyticase (36). Subsequently the bacterial gene was cloned and lyticase is still used as a recombinant enzyme for experiments that require undamaged membranes. Chris also wanted to pursue a biochemical project, so I set him off on an effort to purify yeast actin, which at the time seemed a logical choice for a protein that may be involved in vesicle traffic. Chris completed the project but it was not until years later that Peter Novick, then a postdoctoral fellow in David Botstein’s lab, showed that an actin ts mutation delayed and mislocalized secretion at a restrictive growth temperature (37).

With a small lab, a little money and time free from other responsibilities, I started a couple of my own projects to look at the localization of secretion with a focus on chitin, a polysaccharide in the division septum, and invertase an enzyme secreted into the cell wall. My first undergraduate research student, Vicki Brawley (now Chandler), helped me to study an unexpected surge in chitin synthesis that accompanied the arrest of the yeast cell division cycle in response to the mating pheromone α-factor. That work resulted in my first independent publication, a PNAS paper that was critically edited and communicated by my colleague Clint Ballou (38). The notion of localized deposition and activation of the plasma membrane enzyme chitin synthase seemed tractable but the subject excited little interest outside of a small and contentious community of yeast investigators. Fortunately, a breakthrough in the study of invertase secretion reinforced in my mind the importance of investigating a topic of general interest.

Within a few months, Peter Novick joined the group for his thesis work. Peter was quiet, focused and technically superior. His background was impressive, having trained as an undergraduate at MIT and during summers as a research student in the lab of Arthur Karlin at Columbia University where Peter’s father was a Professor of Physics. Peter focused his studies on invertase, an enzyme that hydrolyzes sucrose to glucose and fructose and which yeast cells use to mobilize hexose for uptake by active transport at the cell surface. Invertase synthesis is repressed in cells growing on a medium containing high (2%) glucose and is derepressed when cells are shifted to low glucose (0.1%). Peter found that secretion of invertase is rapid: The pool of intracellular intermediates in the secretion of invertase is depleted within five minutes after the addition of cycloheximide to block new protein synthesis. He then looked at chemical agents that were reported to block secretion in animal cells to see if they could be used in yeast. My first thought was to find a way to block the fusion of secretory vesicles at the cell surface to see if both secretion and plasma membrane growth were arrested. Those experiments failed and we were faced with a question of how to find secretion mutants.

During that first year, I followed up on an intriguing observation made by Susan Henry, then at Albert Einstein Medical School, who showed that starvation of a yeast inositol auxotroph led to cell death and a rapid arrest in cell growth. She demonstrated that starved cells increase in buoyant density, suggesting an imbalance in macromolecule biosynthesis and net cell surface growth (39). I tested the possibility that inositol may be required for secretion and cell surface growth by assaying invertase activity in intact cells, a measure of enzyme in the cell wall (yeast cells are impermeable to and can not transport sucrose), and in detergent lysed spheroplasts from which the cell wall material had been removed, a measure of intracellular intermediates in secretion. Another dead end; I found that inositol starvation did not block secretion.
Secretion mutants

During my postdoctoral years, I kept a box of cards with ideas about what to pursue in my lab at Berkeley. One of many ideas was a search for secretion mutants. In retrospect, we could have initiated that search right away, but I was not a geneticist and just did not think that way. And when Novick’s work inevitably turned to that approach, we assumed that a block to secretion would be lethal and that one would require a selection procedure to find what might be a rare ts lethal mutation. But what advantage could a dying secretion defective cell have over a viable one? One thought was to select against cells that could take up a toxic substance through a newly synthesized cell surface permease, one whose export would be blocked in a secretion mutant such that the mutant cell would survive exposure to the toxin. We settled on the yeast sulfate permease, which fails to discriminate sulfate and chromate. Under the right conditions, chromate kills cells that express the sulfate permease. Indeed in a screen of mutants that survived exposure to chromate at 37°C, a standard non-permissive temperature for yeast, Peter found a ts lethal mutation that also blocks invertase secretion. However, on reconstructing the conditions of the selection, he found that this mutant died at 37°C even more rapidly than the wild type strain in the presence or absence of chromate. So this was no selection at all! From this we concluded that the mutations may not be so rare after all and that a Hartwell style search among a set of random ts lethal mutations might turn up more secretion specific lesions.

The mutant, sec1, that came from the aborted attempt at a selection, turned out to conform to all the predictions we had made. At a permissive temperature, 24°C, mutant cells behaved like wild type cells in growth and rapid secretion of invertase and another conveniently assayed secreted enzyme, acid phosphatase. The induction and appearance of sulfate permease was also normal. However, on shift to 37°C, sec1 mutant cells arrested secretion of invertase and acid phosphatase (Fig. 2), which accumulated to a high level within dying cells, and the sulfate permease failed to appear in intact cells. These blocks were reversible and on return to 25°C, the accumulated invertase and acid phosphatase were secreted even in the absence of new protein synthesis. Thus, we concluded that the mutant Sec1 protein must be thermally, but reversibly, unstable.
In May of 1978, George Palade visited Berkeley for two lectures in a series sponsored by the pharmaceutical company Smith, Kline and French. This was the first opportunity I had had to meet Palade personally and it was a thrill to be able to share with him what we were doing to study secretion in yeast. He was not aware that yeast cells secrete glycoproteins. The graduate students hosted Palade for dinner and in the course of the conversation, Peter Novick spoke of his new results on the sec1 mutant. Palade encouraged Peter to examine the mutant by thin section microscopy. Shortly thereafter, Peter called my office from the EM lab in the basement of the Biochemistry building, urging that I come inspect the images of sec1 mutant cells. The picture was stunning; cells chock full of vesicles filling the entire cytoplasmic compartment (Fig. 3). An enzyme-specific cytochemical stain for acid phosphatase showed all the vesicles carried this enzyme and likely other proteins secreted by yeast cells. Mutant cells grown at 24°C behaved just like wild type cells did, with a small cluster of vesicles in the bud portion of the cell. Short of the moments I witnessed the birth of my children, nothing in my life compares to the excitement of that image in the EM room in the summer of 1978.
Peter and I assembled a paper for publication in the PNAS, which was communicated by Dan Koshland (40), and we continued a quest for more mutants of this sort because surely with a procedure that was unfavorable for the selection of sec1, many more genes might be found with no selection whatsoever. Peter collected 100 random ts mutant colonies by a standard mutagenesis protocol and found one more mutant, sec2, which phenotypically resembled sec1 in accumulating a uniform population of vesicles. Thus at least two proteins were implicated in some step in the delivery of vesicles to a target membrane, possibly the plasma membrane. But surely there must be more such genes and the prospect of generating thousands of ts colonies in a time before the robotic approaches we now enjoy, was a bit daunting.

For the next of what would be a brilliant string of observations, Peter noticed that sec1 mutant cells fail to enlarge, fail to divide and become phase refractile during an hour or more of incubation at 37C. This
contrasts with the behavior of Hartwell’s cell cycle ts mutants that arrest with a unique cell morphology characteristic of the cell cycle stage that is blocked, but that continue to enlarge into misshapen structures. Peter reasoned that secretion defective cells may continue to produce macromolecules but by failing to enlarge their buoyant density may increase, just as I had expected of the inositol auxotroph of Susan Henry. Peter then performed a beautiful experiment to test his theory. Mutant cells were constructed with a constitutively expressed form of acid phosphatase and an aliquot was incubated at 37C. A corresponding wild type cell sample with a normally repressed phosphatase gene was mixed in a ratio of 100:1 with the mutant cells and the mixed cells were centrifuged on a self-forming gradient of Ludox, a colloidal silica suspension that was then marketed as a commercial floor polish. Susan Henry had exploited the same preparation of Ludox to separate inositol-starved and normal cells. Fractions of the gradient plated on rich medium formed colonies that were then stained with a phosphatase-specific histochemical reagent to reveal the distribution of phosphatase-constitutive and -repressed colonies. The result was an absolute separation of sec1 mutant cells at the bottom of the gradient and wild type cells at the top (Fig. 4). This density gradient then provided the opportunity Peter needed to enrich and screen many more ts colonies for additional sec mutants.

![Figure 4. Ludox gradient of wild-type (light stipples) and sec1 mutant cells (heavy stipples) separated from bottom (high density) to top (low density) of centrifuge tube. Reproduced from ref. #41.](image)

Over the next 18 months, Peter with Charles Field, a technician who was an expert in yeast genetics, repeated the mutagenesis on a large scale with different mutagens and assembled a large collection of density enriched ts colonies, 220 of which proved to be defective in secretion. Genetic complementation tests uncovered 23 genes among these mutants and the distribution of alleles suggested yet more genes were likely to be discovered. Electron microscopic inspection revealed three different phenotypic categories of organelle disruption: Mutations in 10 genes, like sec1 and 2, accumulated secretory vesicles, mutations in another 9 genes caused accumulation and distortion of the ER membrane, and another two caused a toroid-shaped organelle, which Novick called the “Berkeley body”, to proliferate. One concern we had was that the sec mutations might not represent components of the secretory machinery, but merely defective biosynthetic cargo proteins that interfere with secretion. However, the simple complementation tests used to establish the genes showed all the alleles to be genetically recessive, and thus unlikely to
represent dominant inhibitors of the process. Novick and Field completed a morphological and physiological characterization of selected alleles of each of the 23 genes, and we put together a comprehensive paper for the relatively new journal, Cell, which through the force of the personality of the Editor, Benjamin Lewin, was changing the way life science research was evaluated and promoted (41).

In the following year, Novick and Susan Ferro, who later became Susan Ferro-Novick (the first of many marriages within my laboratory), teamed up to apply a classic genetic epistasis test to establish the order in which the SEC genes exert their function. In the course of this work Peter found that one of the mutants sec7 that accumulates the odd “Berkeley bodies” appeared to define a stage equivalent to that of the Golgi apparatus in mammalian cells. Quite by chance he found that this structure irreversibly blocked secretion unless cells were incubated in medium containing low glucose in which case mutant cells accumulate a classic, multi-cisternae Golgi structure (42). Some years later, Chris Kaiser, a talented postdoctoral fellow with considerable experience in yeast genetics, revisited the SEC genes that govern traffic early in the pathway and uncovered a distinct smaller vesicle species that mediates traffic between the ER and the Golgi complex (43). He classified a set of SEC genes that governs vesicle formation and another set required for vesicle consumption, presumably by a process of membrane fusion at the Golgi complex. Importantly, he showed that the two sets of genes show extensive genetic interactions, with mutations in each group exacerbating the mutant phenotype of other members of that group but not between the two groups. This behavior, referred to as synthetic lethal interaction, suggested that the members of each group function together, possibly by physical interaction with one another. These results led to a picture of the secretory pathway in yeast that was essentially the same as Palade had shown for mammalian cells, but with the crucial bonus that each step in the elaborate chain of events was now defined by genes and thus proteins that would surely illuminate the molecular mechanisms of this pathway (Fig. 5, 6).

![Figure 5. Yeast secretory pathway circa 1981. Reproduced from ref. #42.](image)
Two other studies added molecular detail to the emerging view of the secretory pathway in yeast. Brent Esmon, a graduate student in the lab, applied a histochemical stain for invertase on cell lysate samples that were electrophoretically resolved on a native polyacrylamide gel. He discovered that the mutants defective in protein transport from the ER accumulate discrete forms of glycosylated invertase, distinct from invertase that progressed to the Golgi compartment and into secretory vesicles. Using antibodies that diagnose the “outer chain” carbohydrate epitopes of yeast glycoproteins, Brent learned that the division of labor between the ER and the Golgi complex in yeast with respect to N-glycan maturation is much the same as in mammalian cells (44). Tom Stevens, a postdoctoral fellow, studied the traffic of a protein to the yeast vacuole and found that it is diverted from the Golgi complex, similar to the traffic of lysosomal proteins in mammalian cells (45). Stevens and another postdoctoral fellow, Scott Emr, took this part of the pathway to their own labs at the University of Oregon and Cal Tech, respectively, to develop powerful genetic selections to uncover the genes that govern this sorting limb of the secretory pathway. The \(VPS\) genes continue to illuminate the process of sorting from the Golgi complex to the endosome and on to the vacuole or lysosome in all nucleated organisms.

Given our finding that the yeast and mammalian secretory pathways are fundamentally conserved, the biotech industry was quick to exploit the fermentation possibilities of yeast culture to engineer the expression of commercial quantities of important human secreted proteins. Chiron, near Berkeley in the San Francisco Bay Area was the first to succeed. Recombinant expression of the hepatitis B surface antigen in yeast resulted in the production of virus-like membrane particles that proved to be highly immunogenic and which were commercialized as a potent hepatitis vaccine, the sole source of that product in use today (46). As hepatitis B is the major cause of primary liver cancer, the successful introduction of this product of the yeast secretory pathway could, if fully implemented, dramatically reduce the incidence of liver cancer. Indeed this commercial product is considered the first anti-cancer vaccine. Chiron next engineered the expression and secretion of human insulin in yeast and that product,
now marketed by Novo Nordisk, accounts for one-third of the world supply of human recombinant insulin.

I never patented any of our discoveries or thought to do work directed to commercial application in my laboratory because I was completely absorbed by the pursuit of fundamental knowledge. Nonetheless, as a consultant to Chiron I did benefit financially and was enormously gratified to see our work applied to such important practical goals. My view is that the work of drug discovery and practical application is best left to the private sector and that University scientists should focus on basic discovery.

**Important genes uncovered by other means**

Although the initial set of SEC genes revealed the broad outline of the secretory pathway, it became clear that key elements in the process were not reflected in the Novicks’ mutants. We had hoped to find mutations that block the insertion of secretory polypeptides into the lumen of the ER and thus to define genes that constitute the translocation channel predicted by the classic work of Palade, Sabatini and Blobel. The key prediction was that mutations in a putative channel would accumulate unglycosylated secretory precursor polypeptides in the cytoplasm. No such defects were found in the initial set of sec mutations. Susan Ferro conducted a wider search for mutants using the density gradient technique and turned up two that accumulated unglycosylated forms of invertase (47). However, on closer inspection these mutations identified genes involved in the biosynthesis of glycans on secretory proteins rather than bona fide catalysts of translocation (48, 49). Clearly, a different, more directed approach was needed.

Studies in *E. coli* and in yeast showed that the N-terminal signal peptide is necessary and sufficient for the translocation of a secretory protein across the cytoplasmic membrane or ER membrane, respectively (50, 51). The recombinant expression of a chimeric protein constructed by the fusion of a signal peptide coding sequence and the *E. coli* β-galactosidase gene, encoding a soluble cytoplasmic enzyme, result in the membrane translocation of the hybrid protein. Beckwith and colleagues found the expression of such a hybrid protein in *E. coli* provided a selectable growth phenotype, which they used to isolate translocation defective sec mutations, defining the novel cytoplasmic proteins SecA and SecB (52). Using other genetic approaches, Silhavy, Ito and colleagues identified a gene encoding a membrane protein, PrlA/SecY, a candidate for the bacterial translocation channel (53, 54).

Ray Deshaies, an unusually creative and confident graduate student joined the lab in the mid 1980s and after an initial effort with the existing sec mutants, he decided to revisit the translocation problem. In three brilliant but entirely independent efforts, he succeeded in defining a number of genes required in the translocation process. Ray reasoned that if a signal peptide were appended to a cytoplasmic enzyme required for the production of an essential nutrient, the enzyme would be sequestered in the ER, removed from contact with its substrate. In this situation, cells would grow on the nutrient but not on its substrate unless a mutation was introduced that blocked the translocation of the hybrid protein into the ER. Of course, a mutation in an essential channel protein would likely kill the cell, so the quest was for mutations that crippled but did not destroy proteins required for the assembly process. Temperature-sensitive lethal mutations often exert a partial effect at a permissive temperature, thus the search was for mutations that
grow at 30°C on the substrate, in this case histidinol, the substrate of the enzyme histidinol dehydrogenase, the last step in the biosynthesis of histidine, but which fail to form colonies at 37°C on rich growth medium. Ray’s first mutant was called sec61 and further searches using the same selection identified five other genes that encode additional functions essential for translocation, including other subunits of the channel complex and a subunit of the signal recognition particle (SRP) (55, 56). Subsequent cloning of these genes revealed that SEC61 is homologous to the PrlA/SecY gene of E. coli (57). Comparable genes are found in mammals and biochemical analysis demonstrated that the Sec61 protein constitutes the core of the channel protein through which secretory and membrane proteins pass during assembly in the ER (58, 59).

Deshaies also tackled the question of how certain secretory proteins may be translocated post-translationally in yeast. In contrast to the classical rule of co-translational translocation discovered by Blobel, Peter Walter, a protégé of Blobel’s, discovered that at least one substrate, the precursor of the yeast mating pheromone α-factor, could pass across the ER membrane after the completion of translation (60). The assumption was that something extrinsic or intrinsic to α-factor precursor held it in a form that could readily unfold during the translocation event.

In reading an influential review article by Hugh Pelham on the possible role of the heat shock protein family hsp70 in dispersing protein aggregates (61), Ray imagined that hsp70 might also serve to retain partially unfolded forms of post-translational substrates such as α-factor precursor. Fortunately, we were in a position to test this in vivo because Margaret Werner-Washburn and Elizabeth Craig had just constructed a yeast strain missing three members of the major hsp70 class of proteins and with a ts mutation in the remaining fourth gene such that the quadruple mutant was ts lethal. Ray established in short order that this mutant accumulated untranslocated α-factor precursor and as a bonus, he found that the β subunit of the mitochondrial F1-ATPase, also post-translationally translocated into that organelle, accumulated in the cytoplasm. Ray and independently Chirico and Blobel showed that the requirement for Hsp70 could be reproduced in the cell-free reaction that reconstitutes the translocation of α-factor precursor into isolated yeast ER membranes (62, 63).

In a third example of Deshaies’ creative instinct, he solved a problem that had bedeviled a postdoc, Peter Bohni, who had struggled for two years to devise a selection for a mutation in the yeast signal peptidase, the enzyme that Blobel demonstrated cleaves the signal on a secretory polypeptide as it emerges on the luminal side of the ER. Neither the enzyme nor the gene for the peptidase had been obtained, thus it was of interest to test the function of the protein, which at that time remained a candidate for a subunit of the translocation channel. We knew that a mutation at the yeast invertase signal peptide cleavage site delayed the secretion of active enzyme, which accumulates in a precursor form in the ER (64). Attempts to devise a selection for mutations in the peptidase based on that secretion delay proved futile. Ray suggested that some uncleaved cargo proteins might be delayed more seriously than others and that a peptidase mutant could be in our original collection of sec mutants and would have the unusual characteristic of blocking only a subset of cargo proteins. In Peter Novick’s last effort as a graduate student, he had devised a cell surface chemical labeling procedure to assess the full range of major cargo proteins and how their cell surface appearance is affected in sec mutant cells incubated at 37°C (65). Curiously, one mutant in the
original collection, \textit{sec11}, showed an anomalous effect with certain cargo proteins blocked and others less so. With this insight, Bohni immediately investigated the \textit{sec11} mutant and found that it accumulated uncleaved invertase at a restrictive temperature (66). The \textit{SEC11} gene was cloned and found to be the prototype of all eukaryotic signal peptidases (67).

**Cloning genes as an adjunct to functional analysis of Sec proteins**

With the advent of cloning yeast genes by complementation, pioneered by Hinnen and Fink in 1978 (68), we had the immediate prospect of a molecular description of the \textit{SEC} genes and a possible alignment of these genes with comparable functions in simple metazoans and perhaps even mammals. I resisted the temptation to launch in this direction because it seemed unlikely that the \textit{SEC} genes would look like anything else then known. After all, DNA sequencing was still in its infancy and genome databases were nonexistent. Almost from the outset of our characterization of the \textit{sec} mutants, my focus was on attempting to develop a cell-free reaction that reproduced the function of Sec proteins. Most students and fellows who joined the lab resisted my entreaties or took up only half-hearted attempts. One initial effort in this direction yielded a feeble signal that seemed unlikely to prove useful (69). And yet, just miles away in his new lab at Stanford, Jim Rothman had succeeded in developing a reaction that appeared to measure a significant limb of the Golgi traffic pathway reconstituted in a lysate of mammalian cells (70). My own efforts remained on hold until I found a courageous student to take up the challenge.

\textit{SEC53} was the first \textit{SEC} gene cloned and identified with a biochemical function. Although \textit{sec53} was isolated and initially characterized as a mutant defective in translocation, the gene sequence predicted a soluble protein (71), which on closer inspection proved to be to the enzyme phosphomannomutase involved in the production of GDP-mannose, the precursor of \(N\)- and \(O\)-glycans in yeast (48). Other \textit{SEC} genes were cloned but other than predicting that \textit{SEC12} encoded an ER membrane protein and \textit{SEC18} encoded a soluble cytoplasmic protein, no functional biochemical role could be seen in the sequences (72, 73).

The first real breakthrough with respect to vesicular traffic came in 1987 when Novick, now in his own lab at Yale, cloned and sequenced \textit{SEC4}, which he showed encoded a small GTP-binding protein of the \textit{RAS} family (74). Novick’s focus on \textit{SEC4} was no accident. We had agreed that he could take charge of the group of \textit{sec} mutants that block late in the pathway and accumulate mature secretory vesicles. Salminen and Novick found that \textit{SEC4} overexpression suppressed the growth defect of several members of the group of late acting \textit{sec} mutants, and that double mutants constructed among the members of this class displayed a synthetic lethal form of genetic interaction. As the genomes of other organisms were sequenced, it became clear that \textit{SEC4} was a prototype of what are now called Rab proteins, each of which defines a unique destination for the fusion of vesicles to a target membrane. Continuing on the brilliant path he established right from the start of his graduate work, Novick has built a substantial body of highly original work that reveals detailed mechanisms associated with the production, migration and fusion of transport vesicles at the yeast cell surface. And given the fundamental conservation of the \textit{SEC} gene sequences, it is no surprise that Novick’s insights extend to all comparable vesicle targeting/fusion events in metazoans and mammals. Indeed, \textit{SEC1} was found to be related to the \textit{unc-18} gene isolated in the
original collection of uncoordinated mutants of *C. elegans* isolated by Sydney Brenner (75, 76). And the Sec1 protein is known to play a universal role in the control of SNARE protein action in vesicle fusion.

Jim Rothman’s pioneering initial effort to purify proteins required for vesicle fusion yielded the soluble ATPase, NSF (NEM-sensitive factor), which on cloning revealed a striking similarity to the yeast Sec18 protein, a gene that had been cloned by Scott Emr in his own lab at Cal Tech (73, 77). At around the same time, Chris Kaiser in my lab had detected a vesicle intermediate between the ER and Golgi, whose consumption by fusion required the genetically interacting genes SEC18, SEC17 and SEC22 (43). In a joint paper, our labs showed that SEC17 encodes the yeast equivalent of α-SNAP, a protein Rothman’s lab discovered as the factor required for NSF to bind a membrane site, later defined as the SNARE protein (78). Later work showed that SEC22 encodes one such yeast SNARE protein. These results made it clear that the two labs were working on fundamentally the same problem and forged a persuasive link between the mechanism of vesicle targeting/fusion in yeast and mammalian cells.

The mechanism of secretory vesicle budding was now accessible to molecular analysis. Palade had seen coated vesicles at the ER exit site in sections of pancreatic exocrine cells and the view was that the mechanism of budding would involve a coat similar to the classic clathrin coat first visualized as a coated pit engaged in yolk protein internalization in insect oocytes and characterized molecularly by Barbara Pearse with isolated bovine brain clathrin coated vesicles (79, 80). Rothman had evidence to suggest a role for clathrin in the transport of vesicular stomatitis virus G protein from the ER in cultured mammalian cells (81). Thus, clathrin or a similar coat protein was a candidate for one or more of the SEC genes required for traffic from the ER.

Greg Payne decided to assess the role of clathrin directly by cloning the gene for the heavy chain and characterizing the phenotype of a clathrin gene knockout in yeast. Given the expected role of clathrin in vesicular traffic, we assumed the gene would – like the SEC genes – be essential for cell viability. Yet, after disruption of the heavy chain gene in a diploid strain, Greg was shocked to see 2 disrupted spores in each tetrad growing after a several day lag phase. Clathrin deficient cells were sickly but continued to secrete even when the gene was knocked out in a number of different genetic backgrounds (82). Lemmon and Jones reported a strain in which the heavy chain gene was essential but it now seems likely this strain carried an additional mutation that exerted a synthetic lethal effect in the absence of clathrin (83). Further analysis showed that clathrin was required for the proper sorting-retention of a Golgi-localized dibasic peptidase essential for the proteolytic maturation of α-factor precursor (84). These results conformed nicely to the suggestion by Lelio Orci that clathrin coats mediate the retrieval of the proinsulin processing protease from condensing granules in pancreatic β cells (85). The search continued for a coat mechanism in the formation of secretory transport vesicles.

**A yeast cell-free vesicular transport reaction**

I knew that the full potential of the *sec* mutant collection awaited the development of a cell-free reaction to recapitulate at least a portion of the pathway *in vitro*. Finally, in 1985, I recruited a brilliant and creative graduate student, David Baker, who shared my vision and had the talent to make it happen. Up to
that point we had relied on the accumulation of precursor glycoproteins in sec mutant cells arrested at
37C to serve as substrates in in vitro reactions. Immature glycoproteins become modified by specific
outer chain glycan decorations en route through the Golgi complex when cells are returned to the
permissive temperature, and we assumed the same would be true in vitro. This assumption proved wrong.
The first hint of a problem came in the evaluation of sec53 mutant phosphomannomutase, which proved
to be inactive even in lysates of cells that were grown at a permissive temperature (48). But without such
a block to accumulate substrates in the ER, the assay for traffic would have to rely on a low level of
immature glycoproteins radiolabelled for a brief time during biosynthesis. Rothman had succeeded with
just such an approach (71), but the transit time of glycoproteins in yeast is much quicker than in
mammalian cells.

David had a fresh idea. Peter Walter’s lab (as well as the labs of David Meyer and Blobel) had
reconstituted the translocation of radiolabeled α-factor precursor into ER membranes prepared by
mechanical disruption of yeast spheroplasts (61). The product of this incubation was a core N-
glycosylated species that migrated at a discrete position on SDS-PAGE separation. David guessed that
membranes prepared by a more gentle lysis procedure, basically a quick freeze-thaw of yeast
spheroplasts, might preserve membrane organization well enough to permit vesicular traffic of the core
glycan modified synthetic α-factor precursor. Within a few weeks of starting, David observed the
production of a heterogeneous spread of low electrophoretic mobility forms of the radioactive precursor,
which importantly was precipitated by antibodies directed against mannose epitopes added to N-glycans
in the yeast Golgi complex. The reaction required cytosol, ATP and incubation at a physiological
temperature. The results were most promising and the assay was amenable to quantification and easy
repetition with many samples.

The crucial test of Baker’s reaction was to examine the effect of an ER-blocked sec mutant in the cell-free
reaction. Linda Hicke, an ambitious and technically gifted graduate student had cloned SEC23, one of the
four genes Kaiser found to interact in the formation of ER-derived transport vesicles. She collaborated
with Baker to reproduce the α-factor precursor transport reaction in separate incubations containing
membranes from wild type (wt) cells mixed with cytosol fractions from wild type (wt), mutant and
mutant cells complemented with the wt gene. The results were stunning, with a clear ts defect in transport
complemented by a wt copy of Sec23p supplied in the mutant cytosol fraction (86) (Fig. 7). Amazingly,
Susan Ferro-Novick and her graduate student Hannele Ruohola, developed virtually the same
methodology yielding similar results in their laboratory at Yale (87).

Baker and Hicke’s results were precisely what I had dreamed of and the experimental design was
modeled on my own graduate research, in which I used complementation of mutant lysates as an assay to
purify functional DNA replication enzymes (19). With her assay, Linda was able to purify overexpressed
recombinant Sec23p and then to show that it copurified with another protein that was not represented in
our original mutant collection but which proved to be encoded by another essential gene, which we then
called SEC24 (88).
By itself the purified heterodimer of functional Sec23/24p did not offer any clues to its role in ER vesicle budding. For this we required purified forms of the other cytosolic components necessary for budding of α-factor precursor. The next factor came by a circuitous route. Akihiko Nakano, the first of a series of outstanding postdoctoral fellows from Japan, had cloned the ER membrane protein Sec12 that we knew from Kaiser’s genetic work was intimately connected to the set of soluble Sec proteins involved in vesicle budding (43, 59). We had in our collection a set of genes cloned by overexpression suppression of the sec12 mutation. One clone suppressed sec12 ts growth even when its copy number was only two-fold of normal. Aki took this gene back to his lab in Japan and found that it encoded another small GTP binding protein, though of a class distinct from SEC4. This gene, which he called SAR1, also proved essential for secretion (89). Christophe D’Enfert, a postdoc from Paris, found that membranes isolated from a strain overexpressing SEC12 were defective in the transport reaction unless the cytosol contained overexpressed Sar1p (90). This became the assay to purify functional Sar1 that we found could also be isolated by recombinant expression in E coli (91).

As the proteins required for budding were being lined-up, it became clear that the requirements for the full transport reaction were quite complex and it seemed reasonable to devise a simpler assay to focus only on vesicle formation. Michael Rexach joined the lab as a graduate student and brought considerable skill and stubborn determination to this goal. Using a simple technique of differential centrifugation, Michael observed that ER membranes remained intact during the course of the cell-free incubation, as measured by the rapid sedimentation of ER marker proteins. In contrast, he found as the incubation
proceeded that a substantial fraction of the core glycosylated α-factor precursor, which was initially contained within large ER envelopes, was transferred into a slowly-sedimenting vesicle species, which lacked translocation activity and other marker proteins of the ER membrane and lumen. Importantly, the formation of this vesicle species was blocked in the mutants that Kaiser showed to be defective in the production of the vesicle intermediate in vivo (sec12 and sec23), but not in mutants blocked later (sec18) (92). Again, similar results were obtained in Ferro-Novick’s lab (93). Rexach’s work provided us with the essential tool we needed to complete the purification and functional analysis of the proteins required for vesicle budding from the ER.

Two other genes required for ER vesicle formation remained to be functionally identified: SEC13 and SEC16. In his own lab at MIT, Kaiser cloned and characterized SEC16 and learned that it encodes a 240kD peripheral membrane protein, not readily released into the cytosol (94). Nancy Pryer, a postdoc in our lab, cloned SEC13 and found that it encodes a small cytosolic protein that contains a series of WD-40 repeats, very similar to the G protein β subunit (95). Members of this family have a 7-member β propeller structure common to proteins that engage in reversible multi-subunit protein interactions. Nina Salama, an effervescent graduate student in the lab, used Rexach’s budding reaction to purify a functional form of Sec13p and found that it co-purifies with an additional subunit, which we cloned and characterized as a novel SEC gene, SEC31 (96). With this last piece of the puzzle, we found that the budding reaction was sustained with isolated membranes and pure, recombinant Sar1p, Sec23/24p and Sec13/31p, with Sec16p presumably being supplied by the membrane fraction. A complete functional analysis of the mechanism of vesicle budding was now at hand.

**COPII mediates vesicle budding from the ER**

We had few clues as to the mechanism of vesicle budding mediated by the pure Sec proteins in our collection. Rothman’s lab had identified and characterized a novel coat protein complex, coatomer, required for vesicle budding in transport within the Golgi complex (97). He suggested that this coat may also be required for vesicular traffic from the ER, but we found no evidence for subunits of the coatomer in our purified set of Sec proteins. In addition, we had cloned and characterized a different SEC gene, SEC21, that encodes a subunit of coatomer and although the sec21 mutant is blocked in traffic from the ER, it did not fit neatly into one of Kaiser’s mutant classes, and we attributed its effect on traffic from the ER to a backlog of cargo that accumulates when Golgi function is disrupted (98).

Several key insights developed in the 1990s that consolidated our efforts. Two wonderful new postdocs in the lab, Charles Barlowe and Tohru Yoshihisa, discovered a cycle of GTP hydrolysis and exchange on Sar1p. Tohru found that the Sec23 subunit is a GTP hydrolysis catalyst (GAP) specific for Sar1p and Charlie found that the cytoplasmic domain of Sec12p catalyzes nucleotide exchange on Sar1p (99, 100). Several years later Bruno Antonny, a tremendously skilled and perceptive biophysicist, discovered that the Sec31 subunit of the 13/31 heterotrimer complex accelerates the GAP activity of Sec23 10-fold (101). Clearly, a coordinated assembly event controlled by GTP binding and hydrolysis, served to frame the budding process. Given Rothman’s discovery of a role for GTP binding in the control of coatomer
assembly and vesicle budding on Golgi membranes, we were primed for the prospect of a novel coat complex (102).

Fate intervened again in the form of a phone call from the maestro of membrane morphology, Lelio Orci at the University of Geneva Medical School. Orci was instrumental in the effort to discover the morphologic stages in vesicle formation and fusion in the Golgi complex uncovered in the Rothman lab cell-free transport reaction. His skills were so extraordinary that I had attempted, unsuccessfully, to engage his interest when our work uncovered a role for clathrin in the retrieval of a Golgi enzyme similar to his discovery of the organization of clathrin and proinsulin processing in β-cells of the pancreas (85). His call in 1990 was prompted by our recent publication of Kaiser’s analysis of the vesicle species that mediates traffic from the ER. Lelio took pity on us for the primitive standards of our thin section EM analysis and graciously offered his help in a collaboration to examine the organization of the Sec proteins involved in ER vesicle formation. His first success was in using our antibody against the yeast Sec23p to localize the mammalian homolog precisely at the ER exit site in sections of pancreatic tissue (103). But the greatest excitement came when he discovered a novel coat that surrounded the vesicles formed in a reaction with yeast membranes and our purified Sec proteins. Barlowe isolated these vesicles and we saw a hint of a coat in thin sections prepared by my skilled EM technician, Susan Hamamoto, but the images Orci produced were simply breathtaking (Fig 8). We called this novel coat COPII and suggested that the Rothman/Orci coat be referred to as COPI (104). I count it as one of the great privileges of my career to have enjoyed over 20 years of continuous collaboration with Orci, a scholar and experimentalist of the highest distinction.
In a crucial initial collaboration, Orci and a new postdoc in the lab, Sebastian Bednarek, defined the ER as the morphological site of COPII budding. Sebastian purified yeast nuclei as a source of pure ER membrane and with Orci showed that COPII proteins, and curiously also COPI, form buds and incorporate cargo molecules from the outer nuclear membrane. Using a sequential binding assay, Sebastian demonstrated that the COPII assembles in pieces with Sar1p and Sec23/24p binding first and constituting an inner layer of the coat with Sec13/31p forming the outer layer of the coat (105).

With a purified ensemble of cytosolic proteins in hand, we turned our attention to the contribution of membrane proteins and lipids in an effort to define the minimum requirements for vesicle budding.

Sec16p represented the most obvious part of the machinery not accounted for in our reconstituted reaction. Two successive postdoctoral fellows, Joe Campbell and Frantisek Supek, found conditions in which a role for Sec16p in the budding reaction could be observed (106, 107). Eugene Futai succeeded in purifying recombinant Sec16p and found conditions in which it controlled the GTPase cycle mediated by the interaction of the full set of COPII proteins and Sar1 (108). Yet even now, it is not clear if Sec16 participates actively in the cycle of vesicle budding or rather plays a regulatory role in organizing COPII proteins at the ER exit site.

We considered the possibility that coat assembly may be regulated by the availability of membrane cargo proteins. Tom Yeung, a graduate student in the lab, found that membranes isolated from cells treated with
cycloheximide, and thus purged of newly-synthesized cargo, were perfectly active in budding COPII vesicles as assayed by the incorporation of a SNARE protein (109). Although biosynthetic cargo may not be essential for vesicle budding, we suggested that proteins cycling between the ER and Golgi might constitute an essential element of the membrane contribution to the formation of a COPII bud (110).

To test directly the role of membrane proteins and lipids in the budding event, Yeung initiated an effort to solubilize the membrane with detergents to see if membrane proteins and lipids could be reconstituted into liposomes capable of budding synthetic COPII vesicles. To our surprise, Yeung and a meticulous new postdoctoral fellow, Ken Matsuoka, systematically documented that synthetic COPII vesicles bud and could be isolated by density gradient sedimentation from reactions conducted with pure phospholipid liposomes of defined composition provided the reaction was conducted in the presence of a non-hydrolyzable analog of GTP (111). Bruno Antonny developed an elegant real time light scattering assay to monitor the stepwise assembly and disassembly of the coat in incubations containing GTP or a nonhydrolyzable analog (101). Eugene Futai then showed that GTP could replace a nonhydrolyzable analog to produce a stable COPII coated membrane provided the reaction was supplemented with the cytoplasmic domain of the Sar1p nucleotide exchange catalyst, Sec12p, presumably stabilizing the coat by repeated rounds of GTP nucleotide exchange. Curiously, these reactions arrested with buds on liposomes, but few if any completed COPII vesicles (112). More recently, Kirsten Bacia, another postdoctoral fellow, reconstituted the budding reaction on giant unilamellar vesicles where the process may be visualized in real time by light and fluorescence microscopy without potentially damaging manipulation, e.g. centrifugation. In these conditions, incubations containing the COPII proteins and nonhydrolyzable GTP produce long, multi-lobed, coated tubules with regular points of constriction but with little evidence of vesicle fission (113). The nature of the COPII fission reaction remains unresolved but appears to hinge on spatial regulation of GTP binding and hydrolysis at the vesicle bud neck.

The initial event that leads to a bud may begin when Sar1p acquires GTP through interaction with Sec12. Structural analysis showed that the soluble GDP-bound form of Sar1 shields an N-terminal amphipathic helix in a cleft of the folded protein (114). Nucleotide exchange displaces the N-terminus and renders activated Sar1p highly insoluble and prone to membrane insertion. Marcus Lee, an insightful postdoctoral fellow, reasoned that the embedment of the N-terminus in the bilayer may laterally displace phospholipids and create a local asymmetry in the surface area of the two leaflets, much as Sheetz and Singer had proposed decades earlier in the bilayer couple hypothesis (115). In a series of elegant experiments conducted in collaboration with Orci, Lee showed that Sar1p promotes the formation of membrane tubules from synthetic liposomes dependent on the insertion of the amphipathic N-terminal helix and that this insertion is required for COPII vesicle formation in vitro and protein transport in vivo (116).

Tremendous progress has been made on the structural analysis of the COPII coat, principally by the laboratories of Jonathan Goldberg and William Balch (117,118,119). We now have a detailed understanding of the mechanism of polymerization of the two layers of the coat and a key insight concerning the scaffold complex that forms the outer layer, a regular polyhedral lattice that Balch discovered in a self-assembly reaction with purified mammalian Sec13/31 heterotetramer. A former
postdoctoral co-worker, Giulia Zanetti, using cryoelectron microscopy has now visualized the lattice network of COPII formed on the surface of a synthetic liposome (120). Although little evidence suggests any significant structural or functional differences between the yeast and mammalian COPII proteins, mammals have the capacity to regulate the size of the coat to accommodate large or irregularly-shaped cargo complexes such as lipoproteins and pro-collagen. A posttranslational modification, ubiquitylation of Sec31, may serve to regulate some aspect of coat assembly to create a more flexible carrier (121).

An unexpected connection developed between the structure and function of the two layers of the COPII coat in the discovery of a mutation in the human Sec23A subunit. Simeon Boyadjiev and Waffa Eyaid, a Saudi colleague, examined a Bedouin family in which children have a rare craniofacial disorder. The recessive mutation maps to an invariant phenylalanine residue corresponding to a position on the structure of yeast Sec23p facing away from the surface predicted to abut the cytoplasmic face of the ER, a residue not at that time known to have any particular role in coat function or assembly. Orci examined primary skin fibroblasts from one of the afflicted children and observed a profound distortion of the ER and an accumulation of procollagen consistent with a severe defect in secretion (122). Fortunately for us, Jinoh Kim, a courageous postdoc in the lab, had systematically perfected a COPII vesicle budding reaction using membranes isolated from cultured mammalian cells (123). Chris Fromme, another ambitious and skilled postdoctoral fellow, took up the effort to recapitulate the defect seen in the human F382L mutant Sec23A. Chris found conditions that reproduce a budding defect and showed that the defect could be suppressed by increasing the level of recombinant human Sec13/31 in a budding reaction. Further he showed, with all pure mammalian COPII proteins, that the F382L mutant Sec23A has trouble making contact with the Sec13/31 complex as reflected in reduced stimulation of Sar1p GTP hydrolysis (124). At the same time, Goldberg’s lab had solved the structure of the yeast Sec23/24 heterodimer in complex with a fragment of yeast Sec31 that stimulates the GAP activity of Sec23p (125). The point of closest contact between Sec31 and Sec23 was located within angstroms of the position corresponding to the human F382 residue. Thus, the structure of the yeast protein and the functional deficit resulting from mutation in humans could be perfectly reconciled.

The COPII coat guides cargo selection in yeast and in mammalian cells and tissues

In the early 90s the prevailing view was that sorting of secretory and ER resident proteins occurs after cargo exits the ER, mediated by retrieval receptors that return escaped resident proteins back to the ER. Powerful support for this model came with the discovery and characterization of a retrieval signal and a receptor for soluble resident ER proteins such as the luminal hsp70 chaperone, BiP (126). Measurements of the rate of traffic of certain artificial proteins introduced into the secretory pathway argue against the need for active sorting of secretory proteins en route through the pathway (127). Furthermore, two major proteins secreted in the liver appear not to be concentrated in buds that form at the ER exit site, but instead later at the point of COPI-mediated resident protein retrieval (128). Although this issue continues to be the subject of considerable disagreement (129), the results of our vesicle budding reaction where resident proteins are largely excluded from COPII vesicles formed in vitro support an alternative view that active protein sorting accompanies the budding reaction and that resident protein retrieval mediated by sorting receptors in the Golgi membrane may represent a back up mechanism to reinforce the primary
event in the ER (92, 104). Substantial evidence developed over the past 15 years documents a role for ER-localized secretory cargo receptors and one particular subunit of the COPII coat, Sec24p, in the concentrative sorting of membrane and soluble luminal cargo proteins into COPII transport vesicles (130).

If secretory proteins are actively sorted into COPII vesicles, it should be possible to define a sorting signal by the isolation of point mutant forms that produce properly folded precursors that persist in the ER lumen. In practice this has proved difficult because of the uncertainty that a mutant protein may be subject to the quality control retention of misfolded proteins in the ER. Irene Schauer, one of the early graduate students in my lab, isolated just such a mutant of invertase that accumulates in the ER in what appears to be a perfectly active, properly assembled and fully soluble enzyme, but which is secreted from the ER 4–5 fold more slowly than normal (65).

With respect to membrane cargo proteins, early evidence supported a direct interaction with the inner subunits of the COPII coat, Sar1p and Sec23/24p, prior to the complete formation of the coated vesicle. Meta Kuehn, a postdoc in the lab, detected an interaction of plasma membrane permease and SNARE proteins but not luminal ER resident proteins with the inner COPII subunits dependent on incubation of membranes in the presence of a non-hydrolyzable analog of GTP (131). Bill Balch’s lab observed a similar interaction of mammalian Sec23/24 with a transit intermediate of the VSV G protein, and discovered the interaction depends on a C-terminal sorting sequence, ..DxE., in the G-protein (132). Sebastian Springer reinforced this idea with the observation of a stable and selective complex of Sar1p, Sec23/24p and pure recombinant forms of the cytosolic domain of two ER SNARE proteins, Bet1p and Bos1p (133).

For secretory proteins, the best evidence for selective sorting comes from the discovery of ER sorting receptors. David Ginsburg’s laboratory identified the genes involved in a rare, combined hemophilia in which two blood-clotting factors, V and VIII, are delayed in the ER. One gene encodes a lectin-binding membrane protein, ERGIC53 (or LMAN1) (134), that cycles between the ER and Golgi and which is actively packaged into COPII vesicles in a cell-free budding reaction prepared from permeabilized cultured mammalian cells (previous ref). The heavily glycosylated protein domains of factors V and VIII are suggested to interact with the luminal lectin-binding domain of ERGIC 53 to promote their exit from the ER (135). An appreciation of the exact role of ERGIC53 as a sorting receptor awaits the development of an approach to measure the incorporation of a blood-clotting factor into transport vesicles.

A breakthrough in yeast came with the discovery by Charles Barlowe, now in his own laboratory at Dartmouth, of the sorting receptor necessary for the transport of α-factor precursor. In a survey of membrane proteins in isolated COPII vesicles, Barlowe characterized Erv29p, a protein that had not turned up in any genetic screen (136). Deletion of ERV29 produced a viable strain with a pronounced defect in the secretion of α-factor, the mature species produced by proteolytic processing of the precursor in the trans Golgi (137). Unfortunately, Erv29 had evaded detection in classic selections for pheromone deficient yeast mutants because even the 30-fold delay in secretion of α-factor seen in the erv29 deletion strain is inadequate to reduce the steady state level of secreted pheromone below that necessary to
produce an infertile strain of yeast. Other work showed that ERV29 speeds the transport of a vacuolar protease from the ER and likely several other secreted proteins, though notably not invertase (138). Barlowe went on to demonstrate that Erv29p is required to package α-factor precursor into COPII vesicles in vitro and to map the residues responsible for Erv29p interaction with the precursor (137).

One could argue that ERGIC53 and Erv29p serve primarily as species-specific folding chaperones that accompany cargo molecules into the cis Golgi and then are recycled for reuse in the ER and that in their absence the cognate cargo molecules remain subtly unfolded and subject to quality control retention. Such appears to be the case for a large number of species-specific ER membrane chaperones, e.g. Shr3p required for the transport of amino acid permeases in yeast, which remain in the ER and do not accompany cargo into COPII vesicles (139). However, Per Malkus, a graduate student in my lab, showed that α-factor precursor is chemically concentrated 3-fold with respect to a soluble bulk flow marker, a glycotripeptide, within COPII vesicles produced in a budding reaction. This result favors a model of active sorting as opposed to bulk flow in the capture of cargo proteins into COPII vesicles (140).

A complementary line of evidence demonstrates that the COPII coat, specifically the Sec24p subunit, directs the selection of cargo molecules during the budding event. Yeast has three paralogs of Sec24, mammals have four, and genetic and biochemical evidence shows that several are responsible for the capture of particular subsets of cargo membrane proteins. Chris Kaiser’s lab at MIT was the first to recognize the important role of the SEC24 paralog he called LST1 in the transport of the major plasma membrane ATPase, Pma1p (141). Although deletion of LST1 is not a lethal event, cells are sickly and deficient in the surface presentation of the ATPase, which instead accumulates in the ER. Kaiser suggested that Lst1p might form an alternate complex with Sec23p to favor the packaging of Pma1p and that in its absence, the normal Sec24p may not properly sort Pma1p into COPII vesicles. Yuval Shimoni, a postdoc in my lab, proved that directly using a Pma1p budding reaction programmed with either Sec23/Sec24p or Sec23/Lst1p. Lst1p dramatically promoted the packaging of Pma1p into COPII vesicles in vitro (142).

Liz Miller, a wonderfully enthusiastic and talented postdoc, joined the lab to explore the details of cargo sorting mediated by COPII. Following Shimoni’s isolation of functional Sec23/Lst1p, Liz found a remarkably different spectrum of membrane proteins packaged into vesicles produced by the alternative heterodimer, including a defect in the incorporation of α-factor precursor, presumably because Lst1p is not required to recognize most membrane cargo or sorting receptor proteins, including Erv29p (143). Liz then undertook a detailed mutagenesis study designed to identify the residues of Sec24p devoted to the sorting of particular cargo proteins (144). She found mutant alleles that were synthetically lethal when one or both of the other SEC24 paralogs was deleted. One mutation mapped to a binding pocket Jonathan Goldberg had defined structurally on a lateral surface of Sec24p that interacts with the ..DXE.. sorting signal Bill Balch had discovered as important for the traffic of mammalian VSV G protein from the ER (132, 145). Using the budding reaction, Liz showed that Sec24p mutations in this binding site were fully capable of budding certain cargo but not those dependent on the ..DXE.. sorting motif. Per Malkus had identified a ..DXD.. motif in the C-terminal cytoplasmic domain of the yeast amino acid permeases which Liz showed was recognized perfectly adequately by a Sec24p mutant that failed to recognize the ..DXE..
motif (140). These and many results since then have built up a picture of a Sec24p coat subunit with multiple independent cargo binding sites which combined with the two Sec24p paralogs, helps to explain how the diverse repertoire of cargo molecules may be deciphered by a combinatorial code.

An even greater range of cargo proteins is encountered in the mammalian ER. Two striking examples of cargo specificity in sorting mediated by mammalian SEC24 paralogs have been reported. Chain terminating mutations in the mouse SEC24B gene cause an extreme form of neural tube closure defect referred to as craniorachischisis (146, 147). The same arrest is seen in deletions of the neural forms of such signaling receptors as Frizzled and Vangl, two neural epithelium surface proteins that are assembled on the distal and proximal plasma membranes of neural epithelial cells, respectively (148). Using permeabilized cultured mammalian cells, Devon Jensen, a graduate student in my lab collaborating with the laboratory of David Ginty, found that the Sec24B protein stimulated the packaging specifically of Vangl2 protein into COPII vesicles, again consistent with the sequence or structure-selective sorting of membrane proteins at the ER (146). Mutant alleles of human SEC24B may appear in children afflicted with a genetic form of spina bifida. Xiaowei Chen in David Ginsburg’s lab found a striking cargo preference mediated by another paralog, SEC24A. Deletion of SEC24A in the mouse leads to a striking decrease in cholesterol levels in the blood that Chen was able to attribute to a defect in ER transport and secretion of a soluble serum protein, PCSK9, which controls the itinerary of the LDL receptor (149). Lower levels of PCSK9 allow the LDL receptor to cycle efficiently and control cholesterol biosynthesis, thus explaining the low cholesterol in animals deficient in PCSK9 secretion. Chen’s results argue that the export of PCSK9 from the ER is mediated by a sorting receptor that is recognized and packaged into COPII vesicles by SEC24A. The nature of this receptor and its role in sorting of other cargo molecules remain to be discovered. It seems likely that many other such sorting receptors in the ER will be found, adding to the picture of an active process of cargo selection by the COPII coat, and by extension, by other coats involved in the intracellular traffic of membrane and soluble proteins.

**Lessons learned and credit given**

Summarizing almost 40 years of work is a daunting experience, but if I may, three key conclusions follow from the work I have described:

1. Secretion and plasma membrane assembly are physically and functionally linked through a series of obligate organelle intermediates.
2. The polypeptide translocation and vesicular traffic machinery has been conserved over a billion years of evolution.
3. The COPII coat sorts cargo molecules by the recognition of transport signals and physically deforms the ER membrane to create budded vesicles.

Limitations of space and time have made it impossible to acknowledge all the many contributions of the nearly 200 students, fellows and colleagues with whom it has been my privilege to collaborate over the years. Although I end this story here, the work continues in my lab in spite of the many distractions that the call from Stockholm has brought to my life. I am grateful to the present members of my lab for their
patience with me this year but even more importantly for the enthusiasm and dedication they bring to the work at hand. None of this would have been possible without the steadfast love and support of my family and friends, and the wise investment that the U.S. and California made in building educational and research opportunities second to none.

The subject of membrane assembly and vesicular traffic is rich with opportunity and remains an area with great potential for molecular and even atomic resolution in the years ahead. The connections between basic discovery and practical, medical application are certainly more tangible now than when I began my independent work in 1976. However, I trust the pursuit of basic discovery unconnected to any practical application will continue to motivate young scholars and that the agencies, government and private, that made discovery an adventure for me will continue to do so for as long as we thirst for knowledge of the natural world.
REFERENCES


