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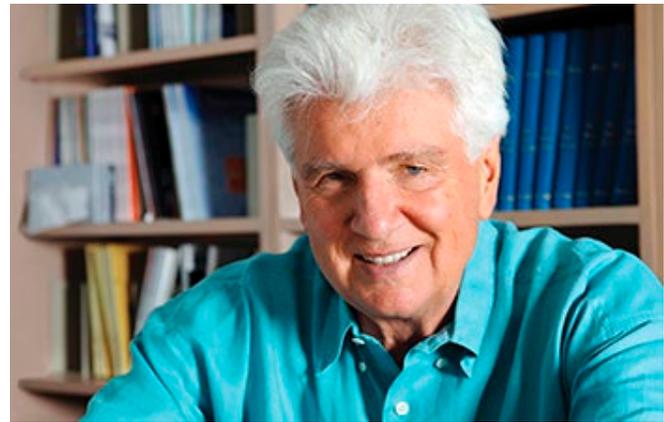
IN MEMORIAM

Günter Blobel: Pioneer of molecular cell biology (1936–2018)

Blobel Laboratory Trainees

Günter Blobel was a scientific colossus who dedicated his career to understanding the mechanisms for protein sorting to membrane organelles. His monumental contributions established research paradigms for major arenas of molecular cell biology. For this work, he received many accolades, including the Nobel Prize in Medicine or Physiology in 1999. He was a scientist of extreme passion and a nurturing mentor for generations of researchers, imbuing them with his deep love of cell biology and galvanizing them to continue his scientific legacy. Günter passed away on February 18, 2018, at the age of 81.

Günter was a *Lebenskünstler*—a master of the art of living. He had a tremendous presence; when he walked into a room, he couldn't help but command attention. A towering man, with a flock of white hair and a jovial nature, he loved to tell stories: of experiences in the laboratory, of life in New York, and of his time growing up in Germany. Günter was born in 1936, as the son of a veterinarian. He considered his early childhood to be idyllic, raised with his seven siblings in a rural and remote part of Silesia, then in the eastern part of Germany. There, largely isolated from the horrific developments overtaking Europe, Günter recalled long afternoons in beautiful manor houses adorned with hunting trophies, winter days of sleighing, and summer days in horse-drawn carriages. However, this life was shattered during the final months of the Second World War, when he and his family were forced to flee the advancing Soviet army. Passing through Dresden at this chaotic time, Günter experienced two life-changing events: seeing for the first time the magnificent baroque splendor of this “great jewel of a city,” and days later, witnessing its leveling by Allied firebombing. The family picked up the threads of their former lives in Freiberg, a medieval town located to the west of Dresden. Again, Günter has fond recollections of his time there. He immersed himself in the town's rich cultural legacy, centered around baroque and classical music performances in its Gothic cathedral. Unfortunately, the new communist regime of East Germany was inhospitable to the Blobel family, who were seen as part of the bourgeois class and were therefore denied access to higher education. Forced to leave Freiberg by these circumstances, Günter moved to Tübingen in West Germany to study medicine. Though he completed an MD degree, he was not inspired to become a practicing physician. Instead, he decided to try a career in research science.



Günter Blobel.

Accordingly, Günter joined the laboratory of Van Potter at the University of Wisconsin to obtain a PhD. As a graduate student, he developed an interest in cell structure and function, and became engaged with a problem that subsequently became an obsession: how the cell sorts secretory proteins to the rough ER. To pursue these interests, he moved to the laboratory of George Palade at the Rockefeller University for postdoctoral studies. Günter's time with Palade was a critical and formative experience, as the Palade laboratory was an intellectual epicenter for study of the ER. Palade and coworkers had described the secretory pathway during the 1960s in an exquisite series of studies that combined cell fractionation, biochemistry, and electron microscopy. Indeed, this and related work on the functions of membrane organelles earned Palade, Christian De Duve, and Albert Claude the Nobel

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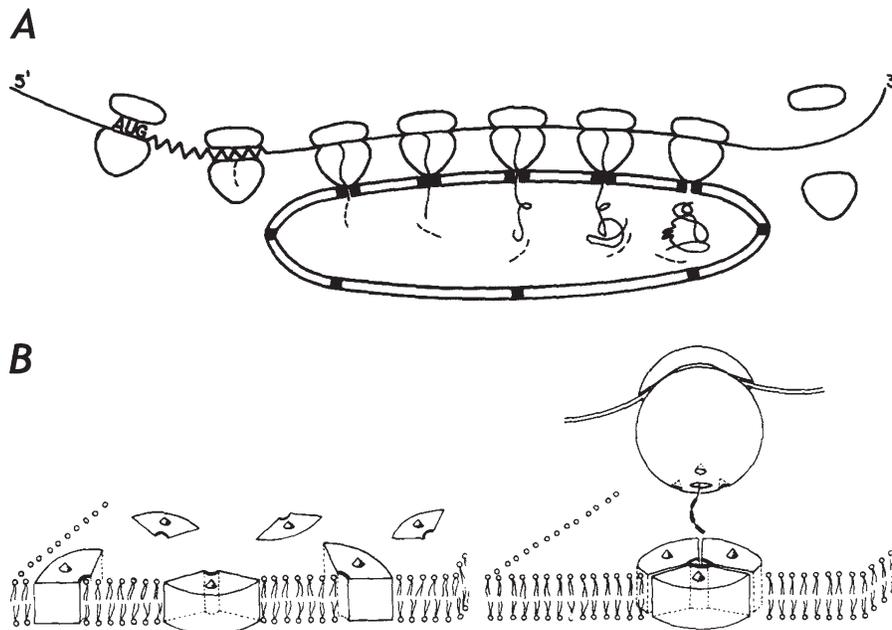


Figure 1. The original signal hypothesis (adapted from reference 2). (A) Illustration of the essential features of the signal hypothesis for the transfer of proteins across membranes. Signal codons after the initiation codon AUG are indicated by a zigzag region in the mRNA. The signal sequence region of the nascent chain is indicated by a dashed line. Endoproteolytic removal of the signal sequence is indicated by the presence of signal peptides (indicated by short dashed lines). (B) Model for the formation of a transient proteinaceous tunnel in the membrane through which the nascent chain is transferred.

Prize in Medicine or Physiology in 1974. Günter revered Palade, and used his mentor's work and conceptualization of cell function as a touchstone throughout his career.

As a newly promoted faculty member at the Rockefeller University, Günter set out to address the question of how mRNAs that encode secretory proteins are selected for synthesis on the rough ER. Despite a lack of experimental evidence, in 1971, Günter and David Sabatini proposed an amazingly prescient hypothesis (1). They suggested that ER targeting of secretory proteins is specified by the presence of a signal peptide on the N terminus of proteins destined to be translocated into the ER. This concept was subsequently termed the “signal hypothesis” (2). “At first it was just a wonderful idea,” Günter remembered; “It was quite a bold thing to say because nothing hinted at a signal sequence. But it was by far the best thing we could come up with.”

Inspired by his work with Palade, Günter knew that only a complete *in vitro* reconstitution of the ER translocation process with purified components—a Herculean challenge at the time—could validate this hypothesis. Encouraged by studies from several laboratories suggesting that the IgG light chain might be synthesized as a precursor polypeptide larger than the secreted form (1972–1975), he worked intensively to develop methods to biochemically dissect the process of ER secretory protein translocation. He provided the first strong experimental support for the signal hypothesis in two landmark papers in 1975 (2, 3). As with many of his most remarkable discoveries, these were published in the *Journal of Cell Biology*. By analyzing IgG light chain synthesis using *in vitro* assays with isolated ER membranes, he obtained striking evidence for the presence of an N-terminal signal sequence on the nascent chains that was involved in cotranslational translocation across the ER membrane, and that was cotranslationally removed (Fig. 1). He predicted that the signal sequence communicated with specific receptor proteins to mediate ER attachment of ribosomes and induce transient assembly of a proteinaceous channel to conduct the nascent polypeptide

across the ER membrane. He also proposed that the mechanisms and components involved in secretory protein translocation could be used for the insertion of membrane proteins into the ER. These ideas were controversial, particularly the notion that a proteinaceous channel was involved in signal sequence-mediated membrane translocation, and became the focus of heated scientific debates for the subsequent two decades. But, as was typical of Günter's outlook on science, he was unfazed: “I thought my ideas were reasonable. So why not propose them?”

Indeed, the controversies surrounding the signal hypothesis were laid to rest, one by one, by a series of elegant papers from the Blobel laboratory and from others over the next two decades. This work identified a cytosolic ribonucleoprotein particle (termed the SRP) that interacts with the signal sequence, a specific SRP receptor in the ER membrane involved in ribosome targeting, a signal peptidase that cotranslationally removes the signal peptide, and, finally, the proteinaceous channel (Sec61 complex) that mediates movement of the nascent chain across the membrane (4). The evolutionary significance of signal-mediated protein translocation across the ER was resoundingly underscored by the identification of conserved systems in diverse organisms including yeast and bacteria. The experimental approach used by Günter and associates for these studies, involving the use of an *in vitro* assay with a quantitative functional output for mechanistic analysis by fractionation and reconstitution, helped usher in the era of modern molecular cell biology.

With the first experimental evidence for the signal hypothesis in hand, Günter prophetically speculated that systems analogous to those for ER translocation also might be deployed for targeting proteins to other membrane organelles (2). These speculations were decisively borne out in subsequent studies on protein transport into the mitochondrion, the chloroplast, the peroxisome, and the nucleus (Fig. 2), providing the ultimate vindication for these bold conjectures. This was a major element of Günter's principle of “protein topogenesis”: that information for sorting

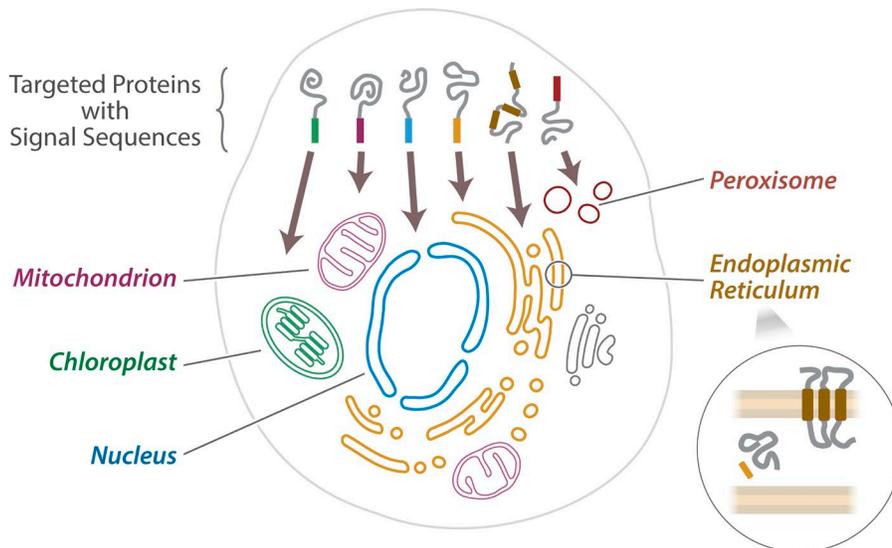


Figure 2. **The principles of protein targeting directed by signal sequences.** A schematic of a cell is shown, with different membrane-bound organelles illustrated and labeled. Newly synthesized proteins carry signal sequences, often but not always at one end of the protein, which can direct that protein to the correct organelle within the cell and allow them to cross the organellar membranes. The lower right inset depicts how additional classes of topogenic sequences (5) can specify the membrane integration of proteins (brown shading) instead of simple membrane translocation with signal sequence cleavage (orange shading).

of proteins to different membrane compartments, as well as for integration into membranes, is encoded in discrete classes of “topogenic” sequences recognized by various receptor systems, membrane-spanning protein conduits, and other effectors (5).

Propelled by his polymathic character, Günter brought his enthusiasm and brilliance to bear on several other areas, particularly nucleocytoplasmic transport and the nuclear lamina. In the mid-1970s, Günter and coworkers made the remarkable discovery that nuclear pore complexes (NPCs) remain as intact structures after detergent extraction of nuclear envelopes, attached to a proteinaceous “nuclear lamina” derived from the inner surface of the nuclear envelope (Fig. 3, A and B; 6, 7). These results suggested that NPCs could be isolated and biochemically analyzed, a goal that was realized many years later by trainees from his laboratory. Günter enthusiastically promoted the view that the nuclear lamina is a widespread nuclear structural component of fundamental importance, albeit not evident in most cells by electron microscopy. These results initiated a stream of biochemical and functional studies that gradually became an experimental

torrent involving numerous laboratories. This effort has firmly established the importance of the lamina in myriad nuclear functions in higher eukaryotes, including nuclear mechanics, signaling, and the dynamic 3D functional organization of chromatin. The relatively recent discoveries linking mutations in nuclear lamina proteins to at least 15 human genetic disorders speak to Günter’s visionary intuition on the importance of the nuclear lamina.

In the arena of nucleocytoplasmic transport. Günter’s group was a frontrunner in the identification of nuclear transport factor proteins and NPC components, his competitive fervor helping to propel the field forward at an astounding rate during the 1990s. More recently, in keeping with his desire to understand the mechanistic principles of cells and fascinated by the molecular architecture of these players, he retailored his laboratory to solve the atomic structures for some of the proteins comprising these assemblies (Fig. 3 C). He drew great joy from solving molecular structures, seeing in them an elegance akin to that found in great architectural masterpieces, and he was adept in

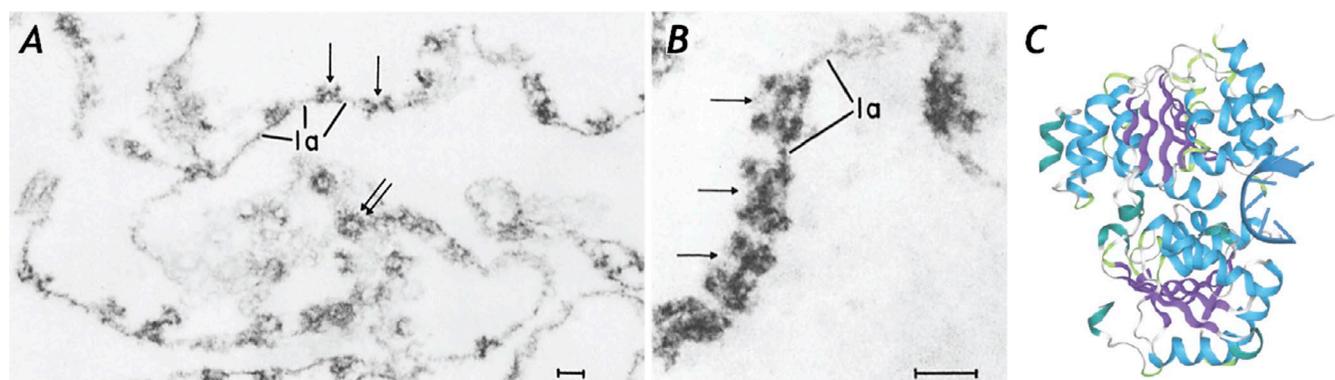


Figure 3. **The nuclear envelope, and associated structures. (A and B)** Transmission electron micrographs of a thin section through an NPC–lamina fraction isolated from rodent liver, showing NPCs in lateral (single arrow) and frontal (double arrow) views, and the associated nuclear lamina (la). Bars, 100 nm (adapted from reference 7). **(C)** An example of one of the most recent structures from the Blobel laboratory (PDB: 5SUP): the messenger ribonucleoprotein particles remodeling complex of Sub2 associated with an ATP analogue, RNA, and a C-terminal fragment of Yra1, required to process and package messenger ribonucleoprotein particles before export through the NPC.

