



Bruce Stillman, Ph.D.

**B**ASIC RESEARCH IS A CENTRAL, DEFINING activity of Cold Spring Harbor Laboratory. It is the wellspring of both new knowledge and technological advances that make new discoveries possible.

This discovery science is expensive, and getting more expensive every year. Inflation in the biomedical sector outpaces that of the broad US economy, mostly due to the wide use of advanced technologies that require expensive equipment or reagents. We also pay a premium for the highly trained personnel who are needed to offer ever increasingly high-tech methods to our faculty, postdoctoral fellows and students. At the same time, we continue to see an erosion of the total amount of support any highly meritorious scientist can obtain from federal sources such as the National Institutes of Health and the National Science Foundation. At the NIH and NSF, policies implemented in the last four years have deliberately limited the type and amount of funding that can be awarded to the nation's very best and most productive scientists in order

to “spread the grants as widely as possible,” a form of scientific socialism that does not bode well for the future of American science. At the same time, however, opportunities abound in many areas of science, including cancer, neuroscience, plant biology and quantitative biology—areas of focus at Cold Spring Harbor.

Fortunately, with very strong support from our Board of Trustees and supporters of CSHL, we have seen a dramatic increase in our endowment. But this precious resource should support the core of what makes CSHL one of the leading research institutions in the world of basic discovery science. Having recognized this, it has been increasingly obvious that there are many instances in which we can add value to our science and translate these discoveries they will make an impact in the clinic—and this is particularly the case for cancer.

With this background, we took the initiative in 2015 of entering into a strategic affiliation with Northwell Health, previously known as the North Shore-LIJ Health System. It's an alliance that I expect to be transformative. It will provide an unprecedented opportunity to add value to certain of our discoveries and multiply the impact of our research.



*It's an alliance that I expect to be transformative.  
It will provide an unprecedented opportunity to  
...multiply the impact of our research.*





*The urgency of speeding the translation of basic research into clinical advances is captured in this picture of pancreatic cancer patient Gail Poinelli conferring with CSHL's Dr. David Tuveson (right) and Northwell Health's Dr. Craig Devoe. The brave Ms. Poinelli, who lost her battle with the illness in 2016, is one of over 40,000 Americans whose lives are claimed by pancreatic cancer annually.*

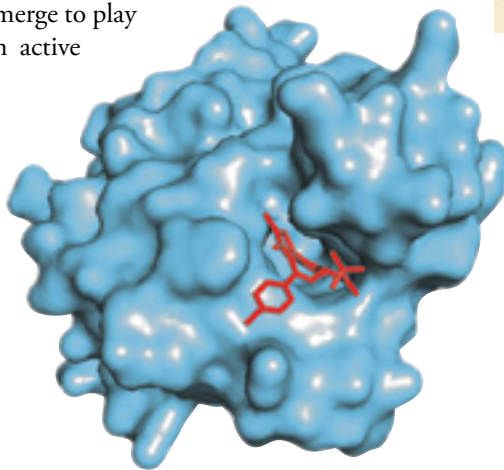
The sweet spot of the affiliation is translational cancer research, which includes the development of new cancer diagnostics and therapeutics and the training of a new generation of research-capable cancer clinicians. Northwell Health is one of the largest integrated health systems in the nation. Its recently expanded Cancer Institute, with over 200 academic oncologists and clinicians, is part of a system of care encompassing 21 hospitals and 400 outpatient physician practices throughout the New York metropolitan area. Serving more than 8 million people, Northwell treats some 19,000 new cancer cases annually. This makes it one of the most important sites of cancer treatment in the US.

As CEO Michael Dowling has noted, Northwell's oncologists will make CSHL's most promising pre-clinical research available to cancer patients in the form of

innovative trials. Patients will be receiving advanced treatments and diagnostics they would not otherwise be offered, and benefit from them years before they would be available to patients elsewhere. At the same time, Northwell's large patient intake provides our scientists with opportunities to perform cancer research using tumor samples from precisely defined subsets of patients. As we move further into the era of targeted therapy, assembling appropriate patient cohorts becomes ever more critical if we want to speed the time it takes to evaluate new treatments.

Clinician-scientists at Northwell have already begun teaming up with faculty at CSHL. Each team has a specific disease focus, or a focus within broad types of cancer such as particular subtypes of breast or prostate cancer. Under the leadership of Dr. David Tuveson, deputy

director of CSHL's NCI-designated Cancer Center and a talented clinician-scientist, Northwell-CSHL teams have begun to gather periodically at our Banbury Center to plan and assess their work. Our agreement additionally supports the education and training of Oncology Fellows. In this aspect of the alliance, the clinical training of oncologists in the Northwell Health system, in conjunction with the Hofstra University-Northwell Health School of Medicine, will include an elective period of laboratory research at CSHL. Via summer and full-year fellowships, a cadre of cancer doctors in training will emerge to play an active



role in translating the next wave of fundamental discoveries about cancer into new diagnostics and therapies.

It's reasonable to ask how the new alliance will change the way research is done at CSHL. I want to make clear that it in no way alters our core commitment to basic research, which is unshakable. This collaboration adds to our capabilities in a manner illustrated by two con-

*...our commitment to basic research is unshakable.*

trasting stories about basic research. One of these stories came to a happy conclusion early in 2015 when the Food and Drug Administration approved palbociclib (Ibrance) for the treatment of metastatic breast cancer. It's a first-in-class inhibitor of CDK4 and CDK6, enzymes called protein kinases that help regulate the cell division cycle. In 1991, David Beach, then a highly

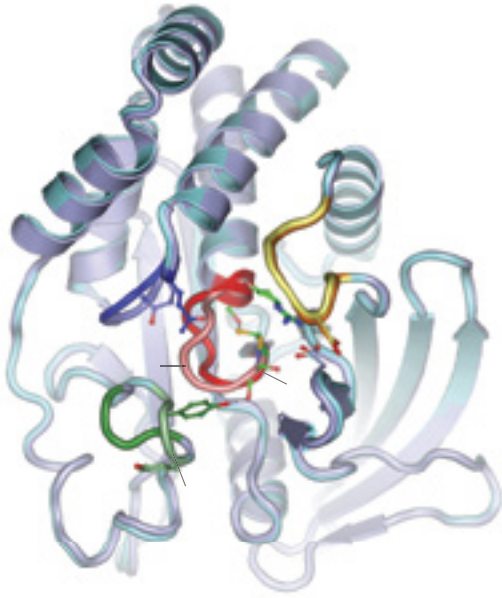


Christopher Vakoc (above) and colleagues in 2011 discovered a powerful drug target called *BRD4* for AML, an often deadly form of leukemia. Translational work has already led to clinical trials testing the effectiveness of a drug called JQ1 (red) that "hits" the target (left).

productive CSHL basic scientist who had already published many papers helping to identify the molecular players involved the control of cell division, reported the discovery of yet another factor, which he called Cyclin D. This discovery coincided with the same finding by former CSHL Trustee Charles Scherr of St. Jude's Children's Hospital, who went on to discover the protein kinase CDK4, which forms a complex with Cyclin D. It soon became clear from the work of Beach and Scherr that the Cyclin D-CDK4 complex is a critical node in the fundamental decision of whether a cell keeps dividing or rests from proliferation. Importantly, it became apparent that most cancer cells have mutations in this control pathway, thereby pushing tumor cells on the road to unchecked growth and aggressive cancer.

What is poignant about the 2015 approval of Ibrance is the fact that it came a quarter-century after the fundamental discoveries by Beach and Scherr. In the early 1990s, we simply did not know enough about cancer to convert their newly generated knowledge into an effective anticancer drug. Now we do. Another basic research discovery at CSHL, this one quite recent, makes the point about our progress vividly, suggesting why the time is ripe for a clinical alliance that enhances our ability to promptly take our basic insights into the clinic. In 2011, Christopher Vakoc, Scott Lowe and CSHL colleagues performed a screen using RNA interference

(RNAi) technology developed at CSHL by Gregory Hannon and his team. The 2011 discovery, which Vakoc has carried forward, revealed a drug target—a protein called BRD4—of unusual potential in the treatment



of aggressive forms of leukemia called acute myeloid leukemia (AML). Vakoc discovered that a drug—developed for another purpose by collaborating scientists at the Dana Farber Institute—hit the target, virtually eliminating AML in mouse models. These studies induced a number of pharmaceutical and biotech companies to initiate clinical trials that target AML, some of which are now in Phase II, with positive results already reported from Phase I studies. This is precisely the kind of rapid translation of an important basic scientific result that our new alliance with Northwell Health and its vast clinical system is designed to facilitate. It will enable us to pursue translational science with a vigor we otherwise could not while keeping our basic discovery engine primed.

Two of our faculty are now reaping the rewards of decades of meticulous basic research. Adrian Krainer's research on RNA splicing—which began in the 1990s and grows out of earlier Nobel Prize-winning work by Louise Chow and Richard Roberts at CSHL and by Sue Berget and Phillip Sharp at MIT—has made possible the development of a drug, now in Phase III trials, for the serious



*The power of basic research is exemplified in discoveries made by Nicholas Tonks and his team. Twenty-five years ago Tonks discovered an enzyme called PTP1B (illustration above) that is now the focus of several drug development efforts with potential applications in breast cancer, diabetes, obesity and Rett syndrome.*

children's disease, spinal muscular atrophy (SMA). Nicholas Tonks' fundamental discovery 25 years ago of the first of what proved a large family of enzymes called protein tyrosine phosphatases (PTPs) was the beginning



*Basic research in plant biology in several CSHL labs has led to discoveries that have the potential to significantly increase the yield of tomatoes, maize, and other food crops.*

of a scientific odyssey in which Nick has persisted in the face of doubters in the pharmaceutical industry. Tonks' team has recently demonstrated their ability to target PTP1B—with a drug Nick developed years ago—in cellular signaling pathways that play a key role in HER2-positive breast cancer. Phase 1 trials will begin at Northwell in the spring of 2016. Other PTP1B-targeting compounds in Tonks' lab are being evaluated by a major pharmaceutical firm for treatment of diabetes and obesity. It's another illustration of how basic science can pay off in ways that are not contemplated at the outset. We see similar promise in other fields: for instance, in Zachary Lippman's basic research on the process of branching morphogenesis in plants, which now points to a way of significantly increasing fruit yields; and in Steven Shea's fundamental research on social behavior in rodents, which has led to unexpected insights into Rett syndrome, an autism spectrum disorder.

Basic research has made all of these opportunities possible. To keep our discovery science robust, we were pleased this past year to have been asked to join the ranks of select institutions named as beneficiaries of the Scientific Philanthropy Alliance. The SPA serves as an impartial advisor to major philanthropists, promoting basic research as the driver of new ideas, of new economic wealth and for the education of a new generation of talented new scientists. Several benefactors of the Laboratory organized the SPA, which we thank for providing another line of support for the basic research that is the lifeblood of Cold Spring Harbor Laboratory.

A handwritten signature in black ink that reads "Bruce Stillman". The signature is written in a cursive, flowing style.

April 2016