19. Trial for Setting-up of Biotechnology Laboratory in SUCs in the Philippines for Kuroshio Science Research Network

Satoshi Kubota, Sam Edward Manalili, Joshua Vacarizas, Teresa N. Avila, Katrina L. Canon and Plutomeo M. Nieves.

1. Collaborative Research under Kuroshio Science Network

After a Memorandum of Understanding (MOU) was signed between the University of the Philippines System, Bicol University in the Philippines, and Kochi University in Japanese FY2007 (November 24, 2005 and March 31, 2006, respectively), many joint research projects have been conducted, mainly in the natural and social sciences with a focus on fieldwork. Phylogenetic classification based on DNA sequence analysis is an important technique for providing basic information for such kind of research on biological resources; however, no researcher from the partner universities in the Philippines has published papers using this technique in "Kuroshio Science" journal and "Kuroshio Symposium." When Satoshi Kubota (first author) visited the Bicol University Tabaco Campus (BUTC) for the first time in March 2017, Plutomeo M. Nieves (last author), a professor from College of Fisheries, was planning to submit an application titled "Upgrading of Research Facilities/Laboratories of the Bicol University Tabaco Campus PCARRD Building" to Department of Science and Technology - Philippine Council for Agriculture, Aquatic, and Natural Resources Research and Development (DOST-PCAARRD), and we discussed about the equipment and reagents required for biotechnological experiments. In August 2017, the application was accepted and funded by the DOST-PCAARRD. To strengthen research collaborations in the Kuroshio Science Network, we set up a biotechnology laboratory and conducted seminars for young faculty and students; we published the first paper on outcomes from these activities in April 2021¹. In this chapter, we summarize the history and current status of biotechnological research, policy of conversion to Outcome-Based Education (OBE) for higher education in the Philippines, and current status of biotechnology-related seminars at the University of the Philippines Diliman. In addition, details of the setting up of the biotechnology laboratory and seminars for young faculty and students in BUTC are presented. Finally, we propose improvements for the stable implementation of biotechnological research in State Universities and Colleges (SUCs) in the Philippines.

2. History and Current Status of Biotechnological Research

Biotechnology, a life science research field, has a significant impact on the society because it is an important applied science. Even if we focus only on "Next-Generation Sequencing," which has made rapid progress in the 2000s, its market size was about 500 million USD in 2018 and is expected to reach 1 billion USD by 2024^2 .

Mendel's law of heredity (Mendel, 1866) was rediscovered in 1900 in independent papers by three botanists: de Vries, Hugo, von Tschermak, E., and Correns, C. The search for the identity of hereditary material began in the early 20th century. Since the beginning of the 20th century, it had been speculated that the genetic material could likely be proteins or nucleic acids. However, it was not until Avery *et al.* showed in 1941 that inoculation of the DNA from a highly virulent strain of *Streptococcus pneumoniae* into mice caused transformation, which strongly suggested that the DNA was the genetic material³. Many biochemists, organic chemists, and physical chemists have been trying to understand the mechanisms underlying DNA replication and conversion of genetic information into proteins during inheritance and reproduction, but the details remain unclear.

Watson and Crick integrated many related discoveries by other researchers and found that DNA strands have a double-helical structure and that the pairing of bases in these strands is pre-determined (complementarity). They also suggested that this structural feature allows DNA to make an exact copy of itself and pass the genetic instructions to the next generation. In addition, Crick advocated for the existence of "central dogma," a scheme for the flow of genetic information (DNA \rightarrow RNA \rightarrow Protein)⁵. In 1961, the first evidence of a triplet code was presented through experiments using T4 bacteriophages⁶. In the same year, Nirenberg and Matthaei showed that polyuridylic acid contains information for the synthesis of poly Lphenylalanine⁷, and subsequently deciphered the entire genetic code, thus winning the 1968 Nobel Prize in Physiology or Medicine along with Holley and Khorana. On the other hand, for proteins, which are functional macromolecules, such as enzymes as biocatalysts, the amino acid sequence of insulin was deciphered for the first time in the early 1950s⁸⁻¹¹. From the late 1960s to the 1970s, DNA ligase¹², which has DNA binding activity, and restriction enzymes 13,14, which cleave specific sequences of DNA strands, were discovered, and "recombinant DNA," a chimeric DNA constructed from DNAs obtained from different organisms, was successfully produced in vitro for the first time¹⁵. On the other hand, a specific strain of E. coli was found to take up DNA in the presence of CaCl₂¹⁶; using this phenomenon, a cyclic plasmid was successfully introduced into the bacterium and the drug resistance gene encoded in the plasmid was successfully expressed 17. By combining these techniques, in 1974, plasmid DNA artificially inserted with eukaryotic (Xenopus laevis) ribosomal DNA was successfully transformed into E. coli (the first report of recombinant DNA)¹⁸. Subsequently, recombinant protein synthesis of mouse dihydrofolate reductase in 1978 and human growth hormone in 1979 was reported in E. coli^{19,20}. Because this recombinant DNA technology could provide bacteria with abilities that they did not originally have, there were concerns about various dangers that could arise due to the advancement of biotechnology. In 1976, the National Institute of Health, USA, established guidelines that severely restricted development beyond what was necessary²¹. However, since 1978, the guidelines have been revised to be more appropriate from the viewpoint of scientific development²², and recombinant DNA experiments can now be performed in ordinary laboratories. In 1979, the postgraduate training course, "Molecular Cloning of Eukaryotic Genes," was held at the Cold Spring Harbor Laboratory, and the technology was widely introduced to young scientists who would lead life sciences in the future. The

manuals used in this training course were revised and published as "Molecular Cloning: A Laboratory Manual" in 1982²³. This series has become a best seller, selling more than 200,000 copies in all editions²⁴. The topics in this manual, which is intended for the beginners, include "isolation of single colonies of bacteria," "verification of strains through genetic markers," and "recovery of purified bacteriophages from a cesium chloride gradient." However, molecular cloning of eukaryotes, whose genome is hundreds of times larger than that of bacteria, is still not easy. The polymerase chain reaction (PCR) technique was developed to specifically amplify and analyze a portion of the DNA sequence of a large genome sequence²⁵, however, because the reaction employed Klenow fragment, enzymes had to be added after each cycle. The usage of Taq polymerase, a DNA polymerase produced by the thermophilic bacterium *Thermus aquaticus*, eliminated the need for the addition of enzymes during the reaction, thus opening the way for reaction simplification and automation²⁶. Immediately after the publication of the PCR method, the cycle sequencing method, which is still used today, was established by combining it with the enzymatic DNA sequencing method described by Sangar et al. in 1977²⁷. In addition, various application methods, such as the detection of mutations and length polymorphisms in DNA sequences²⁸ and quantification of mRNA²⁹, have also been published. A PubMed search using the two keywords "PCR" and "polymerase" yielded 264 hits in 1989, but the number of hits increased rapidly to over 7,200 in 1999. As a subsequent development of the technique, real-time PCR was reported in 1996³⁰ and digital PCR was reported in 1999³¹, and the number of hits exceeded 15,000 in 2012. In the human wholegenome sequencing data published in 2001³², the old Sanger method²⁷ was used. However, with the increasing demand for processing large amounts of DNA sequencing data, various parallel sequencing analysis methods with completely different principles have been developed^{33,34}, which has led to the development of "next-generation sequencing" as described above.

3. Introduction of Outcomes-Based Education (OBE) in Higher Education and Current Status of Biotechnology Education in the Philippines

In the Philippines, where traditional education (input-based and process-based education) has been the mainstream, "Policy-Standard to Enhance Quality Assurance (QA) in Philippine Higher Education thorough an Outcome based and Typology-based QA (Committee of Higher Education [CHED] Memorandum Order, No. 46, 2012)" was released in 2012 with the aim of improving the quality of higher education by shifting the educational system to "outcome-based education" The goal of this policy is to increase productivity through human resource development and enhance international competitiveness. Higher education institutions are required to provide educational services that meet the needs of academia and industry, as well as to develop high levels of academic skills, thinking, and behavior that meet international standards. In the field of biotechnology, it is essential to provide not only knowledge-based lecture-based education, but also practice-based experimental education. To achieve this, a systematic curriculum must be established, many instructors with sufficient abilities to conduct practical training must be trained, and the facilities and equipment in universities and colleges must be improved. In 2017, CHED presented the Policy, Standards, and Guidelines (PSG) in the Biology Department at the undergraduate level³⁶.

Based on CMO No. 46, series of 2012, the PSG specifies the "core competencies" expected of biology department graduates. Regardless of the type of university or college, it is necessary to develop a system that meets the minimum requirements set forth in this standard. Table 1 shows the components of the bachelor's

curriculum and number of credits. Of these, "Biology Tool Courses" and "Specialization Courses," which are related to biotechnology education, are excerpted (Tables 2 and 3).

In biology tool courses, the requirement is to offer half or more of the lecture credits or laboratory courses (Table 2). In specialization courses, more than 30 courses in fields such as cellular and molecular biology, microbiology, systematic biology, biotechnology, and genetics are listed. It is expected that each university will establish its own bachelor's degree.

The requirements for human resources, laboratories, and facilities are also presented in this PSG. As for human resources, unit heads are required to have a doctoral degree and department heads are required to have a master's degree. Each faculty member in charge of specialized subjects is also required to have a master's degree. In terms of laboratories and facilities, appropriate maintenance is required to achieve program outcomes; however, no specific conditions have been specified. In practice, it seems difficult for faculty members who have little experience in biotechnology experiments to provide adequate education in the Department of Biology, which consists only of master's degree graduates. In addition, many of the reagents and tools required for biotechnological experiments are expensive; even a simple series of experiments, such as DNA extraction without genetic recombination, fragment amplification by PCR, and species estimation by restriction fragment polymorphism (RFLP), requires a budget of several hundred thousand pesos, which cannot be maintained easily if sufficient budget is not secured.

Table 4 shows a list of biotechnology-related seminars conducted at the University of the Philippines Diliman in 2018-2019. Since the University of the Philippines is a leader in the academic field in the Philippines, there is a need to update the information pertaining to advanced technologies and their applications, which can be obtained from a variety of sources. Many of these are expected to be useful for the education of faculty members in charge of state universities. Table 5 shows a list of laboratories in the Philippines that require extensive equipment for biotechnological research. Although researchers and collaborators in these institutes can conduct biotechnological research, they are unlikely to have the capacity

Table 1 Components of the BS Bio Curriculum and Their Corresponding Units

COMPONENTS	UNITS
a. General Education Curriculum	36
b. Biology Tool Courses	18
c. Fundamental Courses	50
d. Specialization Courses	25
e. Free Electives	6
f. Undergraduate Thesis	6
g. Practicum or On-The-Job Training or equivalence or apprenticeship	3
h. Physical Education (PE)	8
i. National Service Training Program (NSTP)	6
Total	158

Table 2 Biotechnology Related Courses in Biology Tool Courses (Selected)

Area	Course	Lecture	Laboratory	Units
Chemical Biology	Chemical Biology I (Organic Molecules)	2	1	3
	Chemical Biology II (Analytical Methods for Biology)	2	1	3
	Chemical Biology III (Biomolecules)	3	2	5
BioPhysics	Biophysics	2	2	4

to collaborate with faculty members in biology departments across the country; therefore, it is essential to invest heavily in state universities as regional hubs or to establish biotechnological research institutes with facilities that can be shared.

Table 3 Biotechnology Related Suggested Courses in Specialization Courses (Selected)

Radiation Molecular Molecular Bioinform Genomics Microbiology Virology Microbial	nation DNA Techniques
Radiation Molecular Molecular Bioinform Genomics Wicrobiology Virology Microbial	
Molecular Molecular Bioinform Genomics Microbiology Virology Microbial	To 1 1
Molecular Bioinform Genomics Microbiology Virology Microbial	Biology
Bioinform Genomics Microbiology Virology Microbial	r Genetics
Microbiology Virology Microbial	r Systematics
Microbiology Virology Microbial	natics
Microbial	and Proteomics
	Taxonomy
1,110100101	Physiology
Microbial	Ecology
Microbial	Genetics
Industrial	Microbiology
Food Mic	robiology
Pathology	T.
Epidemio	logy
Systematic Biology Molecular	r Systematics
Phylogene	etics
Population	n Genetics
Bioinform	natics
Evolution	ary Systematics
Biotechnology Health Bio	otechnology
Agricultur	ral Biotechnology
Industrial	Biotechnology
	technology
Molecular	r Genetics
Bioproces	ssing
Tissue Cu	
Bioinform	
	and Proteomics
Genetics Cytogenet	
Molecular	
Human G	
Microbial	
Population	n and Quantitative
Genetics	
	nental Genetics
Biosocial	Genetics

Table 4 list of biotechnology-related seminars conducted at the University of the Philippines in 2018-2019 (Selected)

Biotechnology

CRISPR-Cas and Genomics.

Institute of Plant Breeding, UPLB (Jan 23, 2018)

Establishment and Management of BRCs/Biobanks: a case of KNRCC.

Philippine Genome Center, UP Diliman (Jul 17, 2018)

Molecular and Phenotypic Characterization of Mechanically-Stimulated Transgenerational Arabidopsis.

Institute of Biology, UP Diliman, and Philippine Genome Center (Sep 17, 2018)

Genomics on the road: From Agriculture to Health and Beyond.

PCARI-SGCL, NUAL-PH, and Philippine Genome Center, UP Diliman (Dec 8, 2018)

CRISPR-CAS9: Principles, Applications, & Related Issues.

Institute of Biological Sciences, UPLB (Mar 25, 2019)

Cracking the genomes of marine & non-model organisms: novel approaches for correcting, assembling, and scaffolding genomic data using Brujingraphs and Hi-C contact maps.

Philippine Genome Center, UP Diliman (Apr 8, 2019)

Biosafety Seminar.

Philippine Genome Center-Mindanao Satellite Facility, UP Mindanao (Jul 30, 2019)

NGS Data Management and Bioinformatics Challenges, Basic Concepts and Considerations in Omics Analysis.

Philippine Genome Center-Visayas Satellite Facility (Aug 30, 2019)

Gene editing and Biotechnology trends.

Institute of Biology, UP Diliman, and Rautaki Solutions (Sep 9, 2019)

1st National Genomics Conference.

Philippine Genome Center, UP Diliman (Oct 10, 2019)

Mindanao Wide Genomics Seminar: Accelerating Omics Research.

Philippine Genome Center-Mindanao Satellite Facility, UP Mindanao (Oct 29, 2019)

Structural Biology of Scaffolds: from phages to RNA.

Institute of Biology, UP Diliman, and University of Manitoba (Nov 20, 2019)

Applying Emerging Trends: Developmental Biology in Health, Agriculture and Species Conservation.

Philippine Society for Developmental Biology (Nov 29, 2019)

Bioscience

Novel proteins in the oxidative stress response.

Institute of Biology, UP Diliman, and Vanderbilt University (Nov 26, 2018)

Development of a functional screen for bacterial gene clusters involved in the biosynthesis of fungicides.

Philippine Genome Center, UP Diliman (Aug 3, 2019)

Newton Agham Seminar on Bioactive Natural Products.

National Institute of Molecular Biology and Biotechnology, UPLB (Aug 20, 2019)

Enzyme Engineering: Directed Evolution and Rational Protein Design for the generation of efficient and highly selective biocatalysts.

Institute of Chemistry, UP Diliman (Sep 16, 2019)

PSCB's 10th Annual Meeting and Scientific Convention: Unboxing the Cell: Organelles and Sub-cellular Structures.

Philippine Society for Cell Biology (Oct 7, 2019)

Technology

Molecular delimitation of species using haplowebs and conspecificity matrices: Examples from amphipods, corals and plants.

Philippine Genome Center-Visayas Satellite Facility (Apr 10, 2019)

Assembling genomes into complete chromosomes using Bwise and chromosome conformation capture.

Philippine Genome Center-Visayas Satellite Facility (Apr 10, 2019)

Medical Science

Illuminating Chikungunya Virus proteins through Microscopy.

Institute of Molecular Biology and Biotechnology-National Institutes of Health, UP Manila (Jan 11, 2018)

Medical Records in the Genomic Era.

Philippine Genome Center, UP Diliman (May 29, 2018)

Integrating Microbial Genomics in Public Health and Environment, From Antibiotic Production to Antibiotic Resistance.

National Institute of Molecular Biology and Biotechnology, UPLB (Jul 31, 2018)

Lessons from the Eye and the Race to Finding a Cure for Alzheimer's Disease.

Philippine Genome Center, UP Diliman (Dec 5, 2018)

Molecular Genetic Approaches to the Studies of Freshwater Biodiversity and Dengue Eco-epidemiology.

Institute of Biology, UP Diliman, and Ehime University (Jan 18, 2019)

Cell signaling pathways during thyroid follicle development.

Institute of Biology, UP Diliman, and Mindanao State University-IIT (Apr 29, 2019)

Institute of Biology, UP Diliman, and Orentreich Foundation for the Advancement of Science (OFAS) (May 24, 2019)

Mesenchymal actomyosin contractility is required for androgen-driven urethral masculinization in mice. Institute of Biology, UP Diliman, and Wakayama Medical University (Aug 13, 2019)

Pleiotropic Effects of Methionine Restriction.

Agriculture

Forum on Locally Developed Genetically-Modified Products.

Institute of Biology, UP Diliman (Apr 29, 2019)

Genomics and Proteomics of Onion Armyworm.

National Crop Protection Center, UPLB (Jun 4, 2019)

Application of CRISPR-Cpf1 System in Increasing Yield Components of Rice (var Samba Mashuki).

Institute of Chemistry, UPLB (Oct 28, 2019)

The Genomics of Date Palms.

Philippine Genome Center, UP Diliman (Nov 11, 2019)

"Oh, my Genes": Lessons from Plant Genetics and Genomics for Trait Development.

Philippine Genome Center, UP Diliman (Nov 18, 2019)

Cytogenetics in Plant Breeding: Conservation and Evolution in the Post-Genomic era.

Central Mindanao University, and Philippine Genome Center-Mindanao Satellite Facility, UP Mindanao (Dec 16, 2019)

Table 5 list of laboratories in the Philippines that have large equipment needed for biotechnological research (selected)

Philippine Genome Center: UP Diliman, UP Visayas (Satellite Facility), UP Mindanao (Satellite Facility) (pgc.up.edu.ph)

Institute of Biology UP Diliman (https://biology.science.upd.edu.ph/)

Marine Science Institute UP Diliman(http://www.msi.upd.edu.ph/)

National Institute of Molecular Biology and Biotechnology - UP Diliman (nimbb.science.upd.edu.ph)

Institute of Biological Sciences UPLB (cas.uplb.edu.ph)

National Institute of Molecular Biology and Biotechnology (BIOTECH) - UPLB (biotech.uplb.edu.ph)

National Institutes of Health - Institute of Molecular Biology and Biotechnology UP Manila (http://nih.upm.edu.ph/institute/institute-molecular-biology-and-biotechnology)

College of Fisheries and Ocean Sciences UP Visayas (upvcfos.wordpress.com)

Institute of Environmental and Marine Sciences Siliman University, Dumagete City, Negros Oriental (http://su.edu.ph/schools-colleges/institute-of-environmental-and-marine-sciences/)

National Science Research Institute UP Diliman (nsri.upd.edu.ph)

4. Setting-up of the Biotechnology Laboratory at BUTC and Suggestions for Operational Improvements

BUTC was established in 1949 as the Bicol School of Fisheries. In 1969, it was integrated into Bicol University and became an important college of fisheries representing the Bicol region. It now offers six courses: Bachelor of Secondary Education and Science in "Secondary Education," "Entrepreneurship," "Nursing," "Social Work," "Food Technology," and "Fisheries". When Satoshi Kubota (first author) visited BUTC for the first time in March 2017, Plutomeo M. Nieves (Last author) has just submitted an application titled "Upgrading of Research Facilities/Laboratories of the Bicol University Tabaco Campus PCARRD Building" to DOST-PCAARRD. It was to elucidate the species composition of glass eel by DNA barcoding

for the research project, "The Eel Fishery in Tributaries along the Lagonoy Gulf: Implication to Management and Conservation (Eel Project)" funded by the DOST-PCAARRD. We shared information on the requirements for the establishment of a biotechnology laboratory at the campus and agreed to set up a functional biotechnology laboratory not only for the development of research and education at BUTC but also for the enhancement of the Kuroshio Science Network. In addition, Satoshi Kubota presented "Analysis of

Stony Coral with Advanced Life Technologies," where we discussed the methods used in biotechnology (e.g., PCR, Restriction Fragment Length Polymorphism (RFLP), and amino acid sequencing by LC-MS/MS) on March 7, 2017. After the application was officially accepted by DOST-PCAARRD, Satoshi Kubota presented the results of our research titled "New Approach for Scleractinian Coral Analysis (DNA sequencing and phylogenetic analysis)" at the "Research Proposal Write-Shop and RDE Management Forum" and "Multiple clades of zooxanthellae" at the "Bicol Region Aquatic Resource Management Forum (Metagenomic analysis)" held at BUTC and BU Legazpi East Campus on August 31 and October 25 and 26, 2017, respectively. In November 2017, we received a notification regarding the delivery of expected equipment and reagents for DNA analysis, and hence, we planned a Conference on Biotechnology Research [ConBio] (Organizer: Assistant Professor Alex P. Camaya, graduated from Kuroshio Science Program, Kochi University) on December 7 and 8, 2017. We organized lectures and hands-on seminars for young researchers and graduate students not only from Bicol University but also from other state universities. the renovation of the biotechnology laboratory in the BUTC-PCAARRD building was still in progress, and the equipment and reagents necessary for DNA extraction and subsequent PCR amplification, except for micropipettes, thermal cycler, and image analyzer (Photos 1 and 2), did not arrive as of the first week of December. Therefore, Satoshi Kubota made a video of the experimental procedures, including sample pulverization by liquid N₂, DNA extraction, and DNA amplification by PCR, before leaving Japan; the movie was shown and explained in the lecture "Virtual Demo: Molecular Phylogenetic Analysis for Marine" (Photo 3).



Photo 1 Biotechnology Laboratory in PCAARRD building of BUTC (Dec. 7, 2017)



Photo 2 Introduced equipment and tools in PCAARRD building of BUTC (Dec. 7, 2017)



Photo 3 Virtual Technical Seminar in BUTC (Dec. 7, 2017)

To promote research projects using biotechnological experiments, it is necessary to develop human resources that can devote themselves to research. After Katrina Canon (5th author) applied for the MEXT Special Program in January 2018, we discussed the target organisms to be analyzed for her doctoral course research; we also discussed the research plan, including DNA extraction and PCR amplification, to be carried out in the BUTC biotechnology laboratory in February 2018. On August 30, 2018, Katrina Canon and Satoshi Kubota instructed about the research on DNA extraction, quantification, and computer analysis of DNA sequences to the research assistant of the Eel Project; however, they could not analyze their major samples because some equipment and kits had not been delivered even though it had been more than one year since the acceptance of the research project. Since the DNA extraction kit was delivered in December 2018, we were able to extract glass eel samples that needed to be analyzed in the Eel Project for the first time in this biotechnology laboratory on January 4, 2019. Subsequently, the research assistants prepared DNA extracts for DNA amplification by PCR from many glass eel samples. A hands-on seminar on PCR amplification was conducted at the BUTC biotechnology laboratory by Katrina Canon on September 4, 2019, however, the PCR amplification reaction stopped due to the failure of the new thermal cycler despite that was the first time it was used. It took more than a month to discover that the failure was caused by the program due to insufficient support from the supplier. Ultimately, it took more than half a year to introduce the updated program. Therefore, the DNA extraction was carried out at BUTC, and with the permission of the Bureau of Fisheries and Aquatic Resources, Department of Agriculture, the samples were sent to Kochi University for PCR amplification, RFLP analysis, and DNA sequence analysis¹. As described here, it was not easy to carry out even the basic biotechnological experiments involving DNA extraction from biological samples and the amplification of DNA fragments by PCR. However, there are few facilities in the Bicol region where such experiments can be performed at present. Therefore, the sample preparation for the degree research of Teresa Avila (4th author), who is working on the dried fish "Abo" in the neighboring state of Camarines Sur, was also performed at BUTC³⁷. Thus, it would be of great benefit to other campuses and universities if a biotechnology laboratory could be established in a local state university. Unfortunately, on November 1, 2020, Super Typhoon Goni hit the Bicol region and damaged many buildings in BUTC, leaving not only the biotechnology laboratory, but also many other facilities still unusable.

Through the ongoing trials pertaining to the setup of our biotechnology laboratory since 2017, some problems have become apparent. Three possible improvements are suggested below. The first is shortening the approval period. In our experience, there have been cases where research projects have been conducted and orders for equipment and reagents have been submitted, but orders to the supplier have been delayed. The reagents required for biotechnological experiments are expensive, and many have short expiration dates. While waiting for the reagents to be delivered, reagents that have already been purchased may become unusable rendering it as a waste of government resources. To avoid this, the existing procurement system must be improved to enable prompt approval, processing, purchase and delivery, at least for reagents that have been requested for the project use. The second suggestion is to improve the infrastructure of the university, BUTC in particular. The frequent arrival of large typhoons and unstable energy supplies have caused frequent power outages on the campus, as such, its mandated functions and delivery of services greatly disrupted to the detriment of its target clients it serves. This kind of investment may be huge but the benefits are of paramount importance to the vision, mission and goals of the University. Therefore, it is essential that the government should invest heavily in State Colleges and Universities (SUCs) in Bicol to create a more responsive, relevant and innovative R&D in the region. The third suggestion pertains to systematizing the human resource

development. It is sufficient to attend short-term courses or technical seminars offered by universities in Metro Manila, such as UP, UST, DSLU to learn specific experimental techniques. However, to support the smooth progress of biotechnological research at the university, it is necessary to acquire the ability to create a comprehensive staff development plan and biotech roadmap in the region. It may be difficult for a single university to implement these three suggestions. We hope that the national and local government support and assistance for higher education, research, and development will be given utmost attention and priority.

References

- 1 Canon, K.L. et al., 2021. Occurrence of *Anguilla luzonensis* in the Tributaries along the Lagonoy Gulf, Philippines. J. Fish. Sci., 3, 8-16.
- 2 Research And Markets com's..2020. Global Next Generation Battlefield Technology Market: Focus on Technology, Application, and Component Analysis and Forecast, 2019-2024.
- Avery, O.T., 1944. Studies on the Chemical nature of the Substance Inducing Transformation of Pneumococcal Types Induction of Transformation by a Deoxyribonucleic Acid Fraction Isolated from Pneumococcus Type III. J. Exp. Med. 79, 137-158.
- Watson, J.D., Crick, F.H., 1953. Molecular Structure of Nucleic Acids: A Structure for Deoxyribose Nucleic Acid. Nature, 171, 737-738.
- 5 Crick, F.H., 1958. On Protein Synthesis. Symp. Soc. Exp. Biol., 12, 138-63.
- 6 Crick, F.H. et al., 1961. General Nature of the Genetic Code for Protein. Nature, 192, 1227-1232.
- Nirenberg, MW, Matthaei, H., 1961. The Dependence of Cell-Free Protein Synthesis in *E. coli* upon Naturally Occurring or Synthetic Polyribonucleotides. Proc. Natl. Acad. Sci. USA. 47, 1588-1602.
- 8 Sanger, F., Tuppy, H., 1951. The Amino-Acid Sequence in the Phenylalanyl Chain of Insulin -1. The Identification of Lower Peptides from Partial Hydrolysates-. Biochem. J., 49, 463-481.
- 9 Sanger, F., Tuppy, H., 1951. The Amino-Acid Sequence in the Phenylalanyl Chain of Insulin -2. The Investigation of Peptides from Enzymatic Hydrolysates-. Biochem. J., 49, 481-490.
- 10 Sanger F., Thompson, E.O.P., 1953. The Amino-Acid Sequence in the Glycyl Chain of Insulin -I. The Identification of Lower Peptides from Partial Hydrolysates-. Biochem. J., 53, 353-366.
- anger F., Thompson, E.O.P., 1953. The Amino-acid Sequence in the Glycyl Chain of Insulin -II. The Investigation of Peptides from Enzymatic Hydrolysates -. Biochem. J., 49, 366-374.
- 12 Weiss B., Richardson C.C., 1967. Richardson, Enzymatic Breakage and Joining of Deoxyribonucleic Acid, I. Repair of Single-Strand Breaks in DNA by an Enzyme System from *Escherichia coli* Infected with T4 Bacteriophage. Proc. Natl. Acad. Sci. USA. 57, 1021-1028.
- 13 Smith H.O., Wilcox K.W. 1970. A Restriction Enzyme from *Hemophilus influenzae*. I. Purification and general properties. J. Mol. Biol., 51, 379-91.
- 14 Kelly T.J., Jr, Smith H.O., 1970. A Restriction Enzyme from *Hemophilus influenzae*. II. J. Mol. Biol., 51, 393-409.
- 15 Jackson D.A., 1972. Biochemical Method for Inserting New Genetic Information into DNA of Simian Virus 40: Circular SV40 DNA Molecules Containing Lambda Phage Genes and the Galactose Operon of Escherichia coli. Proc. Natl. Acad. Sci. USA. 69, 2904-2909.
- 16 Mandel, M., Higa A., 1970. Calcium-Dependent Bacteriophage DNA Infection. J. Mol. Biol., 53, 159-162.

- 17 Cohen, S.N. et al., 1972. Nonchromosomal Antibiotic Resistance in Bacteria: Genetic Transformation of *Escherichia coli* by R-Factor DNA. Proc. Natl. Acad. Sci. USA. 69, 2110-2114.
- 18 Cohen, S.N. et al., 1973. Construction of Biologically Functional Bacterial Plasmids *in vitro*. Proc. Natl. Acad. Sci. USA. 70, 3240-3244.
- 19 Chang, A.C. et al., 1978. Phenotypic Expression in *E. coli* of a DNA Sequence Coding for Mouse Dihydrofolate Reductase. Nature, 275, 617-24.
- 20 Goeddel, D.V. et al., 1979. Direct Expression in *Escherichia coli* of a DNA Sequence Coding for Human Growth Hormone. Nature, 281, 544-548.
- 21 National Institute of Health (NIH), 1976. DNA Recombinant Research. NIH Guide for Grants and Contracts, 5, 11-13.
- 22 Johnson, J.A., 1982. The NIH Recombinant DNA Guidelines: Brief History and Current Status., The Library of Congress Congressional Research Service Major Issues System, Issue Bried Number, IB82057.
- 23 Cold Spring Harbor, 1982. Molecular Cloning: A Laboratory Manual (Eds: Fritsch and Joe Sambrook).
- 24 Creage, N.H., 2020. Recipes for recombining DNA: A History of Molecular Cloning: A Laboratory Manual. Angela. BJHS Themes, 5, 225-243.
- 25 Saiki, RK., 1985. Enzymatic Amplification of Globin Genomic Sequences and Restriction Site Analysis for Diagnosis of Sickle Cell Anemia. Science, 230, 1350-1354.
- 26 Saiki, RK., 1988. Primer-Directed Enzymatic Amplification of DNA with a Thermostable DNA Polymerase. Science, 239, 487-491.
- Sangar, F. et al., 1977. DNA Sequencing with Chain-Terminating Inhibitors. Proc. Natl. Acad. Sci. USA, 74, 5463-5467.
- Wrischnik, L. A., 1987. Length Mutations in Human Mitochondrial DNA: Direct Sequencing of Enzymatically Amplified DNA. Nucl. Acid Res, 15, 529-542.
- Wang, A.M., 1989. Quantitation of mRNA by the Polymerase Chain Reaction. Proc. Natl. Acad. Sci. USA. 86, 9717-9721 (1989).
- 30 Heid, C.A., 1996. Real Time Quantitative PCR. Genome Research, 6, 986-994.
- 31 Vogelstein, B., Kinzler, B.V., 1999. Digital PCR. Proc. Natl. Acad. Sci. USA. 96, 9236-9241.
- 32 International Human Genome Sequencing Consortium. 2001. Initial Sequencing and Analysis of the Human Genome. Nature, 409, 860-921.
- Ronaghi, M. et al., 1998. A Sequencing Method Based on Real-Time Pyrophosphate. Science, 281, 363-365.
- 34 Brenner, S. 2000. Gene Expression Analysis by Massively Parallel Signature Sequencing (MPSS) on Microbead Arrays. Nature Biotechnology, 18, 630-634.
- 35 Commission on Higher Education. 2012. Policy-Standard to Enhance Quality Assurance (QA) in Philippine Higher Education Through an Outcomes-Based and Typology-Based QA. Commission on Higher Education Memorandum Order, No. 46, Series of 2012.
- 36 Commission on Higher Education. 2017. Policies, Standards and Guidelines for the Bachelor of Science inf Biology (BS BIO) Program. Commission on Higher Education Memorandum Order, No. 49, Series of 2017.
- 37 Avila et al., 2022. Proximate composition and changes in muscle proteins of dried salted abo (*Otolithes ruber*). Food Research, 6, 178-187.