based searches, analyse and discuss results in the light of molecular biology knowledge;	
• explain major steps in pairwise and multiple sequence alignment, explain its principles and execute pairwise	
sequence alignment by dynamic programming; • predict secondary and tertiary structures of protein	
sequences;	
• perform and analyse various statistical tools available to analyse the data.	

Programme: M. Sc. Biotechnology

Course Code: GBO-186 Title of the Course: Field Trip and Report

Number of Credits: 1

Effective from AY: 2019-2020

Programme: M. Sc. Biotechnology

Course Code: GBO-187 Title of the Course: IPR, Biosafety And Bioethics

Number of Credits: 2

Effective from AY: 2019-2020

Prerequisites for the course:	No prerequisites required.	
Objective:	To provide basic knowledge onintellectual property rights and theirimplications in biological research and product development; • To become familiar with India's IPR Policy; • To learn biosafety and risk assessment of products derived from biotechnology and regulation of such products; • To become familiar with ethical issues in biological research. This coursewill focus on consequences	

ofbiomedical research technologies suchas cloning of whole organisms, genetic modifications, DNA testing.

Content:

12 hours

Introduction to intellectual property; types of IP: patents, trademarks, copyright & related rights, industrial design, traditional knowledge, geographical indications, protection of new GMOs; International framework for the protection of IP; IP as a factor in R&D; IPs of relevance to biotechnology and few case studies; introduction to

history of GATT, WTO, WIPO and TRIPS; plant variety protection and farmers rights act; concept of 'prior art': invention in context of "prior art"; patent databases - country-wise patent searches (USPTO, EPO, India);

analysis and report formation.

MODULE I

Basics of patents: types of patents; Indian Patent Act 1970; recent amendments; WIPO Treaties; Budapest Cooperation Treaty; Patent Treaty (PCT) implications; procedure for filing a PCT application; role of a Country Patent Office; filing of a patent application; precautions before patenting-disclosure/non-disclosure patent application- forms and guidelines including those of National Bio-diversity Authority (NBA) and other regulatory bodies, fee structure, time frames; types of patent applications: provisional and complete specifications; PCT and conventional patent applications; international patenting-requirement, procedures and costs; financial assistance for patenting-introduction to existing schemes; publication of patents-gazette of India, status in Europe and US; patent infringement- meaning, scope, litigation, case studies and examples; commercialization of patented innovations; licensing - outright sale, licensing, royalty; patenting by research students and scientists-university/organizational rules in India and abroad, collaborative research - backward and forward IP; benefit/credit sharing among parties/community, commercial (financial) and non-commercial incentives.

MODULE II

12 hours

Biosafety and Biosecurity - introduction; historical background; introduction to biological safety cabinets; primary containment for biohazards; biosafety levels; **GRAS** organisms, biosafety levels of specific microorganisms; recommended biosafety levels for infectious agents and infected animals; definition of GMOs & LMOs; principles of safety assessment of transgenic plants - sequential steps in risk assessment; concepts of familiarity and substantial equivalence; risk – environmental risk assessment and food and feed safety assessment; problem formulation – protection goals, compilation of relevant information, risk characterization and development of analysis plan; risk assessment of transgenic crops vs cisgenic plants or products derived from RNAi, genome

International regulations — Cartagena protocol, OECD consensus documents and Codex Alimentarius; Indian regulations — EPA act and rules, guidance documents, regulatory framework — RCGM, GEAC, IBSC and other regulatory bodies; Draft bill of Biotechnology Regulatory authority of India - containments — biosafety levels and category of rDNA experiments; field trails — biosafety research trials — standard operating procedures — guidelines of state governments; GM labeling — Food Safety and Standards Authority of India (FSSAI).

Introduction, ethical conflicts in biological sciences - interference with nature, bioethics in health care - patient confidentiality, informed consent, euthanasia, artificial reproductive technologies, prenatal diagnosis, genetic screening, gene therapy, transplantation. Bioethics in research – cloning and stem cell research, Human and animal experimentation, animal rights/welfare, Agricultural biotechnology - Genetically engineered food, environmental risk, labeling and public opinion. Sharing benefits and protecting future generations - Protection of environment and biodiversity – biopiracy.

Pedagogy:

lectures/ tutorials/assignments/self-study

References/Readings

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- 2. Patents for Chemicals, Pharmaceuticals a nd Biotechnology:Fundamentals of Global Law, Practice and Strategy (2010) Grubb P. W. Grubb, P. L. Thomsen, P. R. Oxford University Press.
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- 9. A User's Guide to Patents (2007) Trevor M. Cook. Tottel Publishing.
- 10. Biotechnology and Patent laws:patenting living beings (2008) Sreenivasulu, N.S. and Raju C.B. Manupatra Publishers.
- 11. Complete Reference to Intellectual Property Rights Laws. (2007). Snow White Publication Oct.
- 12. Craig, W., Tepfer, M., Degrassi, G., &Ripandelli, D. (2008). An Overview of General divisions/csurv/geac/annex-5.pdf F. (2009). Problem Formulation in the Environmental Risk Assessment for Genetically Modified Plants. Transgenic Research, 19(3), 425-436. doi:10.1007/s11248-009-9321-9
- 13. Features of Risk Assessments of Genetically Modified Crops. Euphytica
- 14. Ganguli, P. (2001). *Intellectual Property Rights: Unleashing the Knowledge Economy*. New Delhi: Tata McGraw-Hill Pub.
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- 19. National Biodiversity Authority.

http://www.nbaindia.org 20. National IPR Policy, Department of Industrial Policy & Promotion, Ministry of Commerce, GoI, National Portal of India. http://www.archive.india.gov.in 21. Office of the Controller General of Patents, Design & Trademarks; Department of Industrial Policy & Promotion; Ministry of Commerce & Industry; Government of India. http://www.ipindia.nic.in/ 22. Patents for Chemicals. Pharmaceuticals a nd Biotechnology:Fundamentals of Global Law, Practice and Strategy (2010) Grubb P. W. Grubb, P. L. Thomsen, P. R. Oxford University Press. 23. Recombinant DNA Safety Guidelines, 1990 Department of Biotechnology, Ministry of Science and Technology, Govt. of India. Retrieved from http://www.envfor.nic.in/ 24. Wolt, J. D., Keese, P., Raybould, A., Fitzpatrick, J. W., Burachik, M., Gray, A., Wu, World Intellectual Property Organisation. http://www.wipo.int 25. World Trade Organisation. http://www.wto.org **Learning Outcomes** On completion of this course, students should be able to: • understand the rationale for andagainst IPR and especially patents; • understand why India has adopted n IPR Policy and be familiar withbroad outline of patent regulations; • understand different types ofintellectual property rights in generaland protection of products derivedfrom biotechnology research andissues related to application and obtaining patents; • gain knowledge of biosafety andrisk assessment of products derived from recombinant DNA researchand environmental release modified ofgenetically

organisms, national and international regulations.