

# **SYLLABUS FOR THE BATCH FROM YEAR 2025 TO 2026**

**FOR**

## **Certificate Course in Artificial Intelligence in Human Genetics**

### **(Credit Based Evaluation and Grading System)**

**Semester: I**

**EXAMINATIONS: 2025-2026**

**Certificate Course in Artificial Intelligence in Human Genetics (6 Months duration)**



#### **Program Outcomes:**

- AI Applications in Genomics** – Learn how AI enhances genetic screening, diagnosis, biomarker discovery, and gene therapy.
- Technical Skills in AI-Driven Drug Discovery** – Gain insights into AI's role in pharmacogenomics, drug repurposing, and clinical decision support systems.
- Ethical and Regulatory Considerations** – Explore bioethical principles, genetic privacy regulations, and the implications of AI in genomic medicine.
- Genetic Counseling and AI** – Develop knowledge of genetic risk assessment, AI-assisted counseling, and ethical challenges in AI-driven genetic diagnosis.

**Name of the Department: Human Genetics**

**In collaboration with**

**Directorate of Open & Distance Learning and Online Studies**

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**GURU NANAK DEV UNIVERSITY**  
**AMRITSAR**

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**Programme Eligibility**

- +2 biology with atleast 45%marks in aggregate (40% for SC/ST candidates)
- Any student doing Bachelor Degree, Master Degree, Ph.D. from GNDU.
- Working Professionals in Health Sectors/Pharma/IT etc.

**SEMESTER-I**

<b>Paper Code</b>	<b>Subject</b>	<b>Marks</b>			<b>Credits</b>
		<b>Internal Assessment</b>	<b>End Term</b>	<b>Total</b>	
ODAIHG111T	Artificial Intelligence in Human Health and Diseases	30	70	100	4
ODAIHG112T	Artificial Intelligence in Genetic Screening, Diagnosis and Drug Development	30	70	100	4
ODAIHG113T	Artificial Intelligence in Gene Therapy and Gene Editing	30	70	100	4
ODAIHG114T	Artificial Intelligence and Ethical Considerations in Human Genetics	30	70	100	4
<b>Total Marks &amp; Credits</b>		<b>120</b>	<b>280</b>	<b>400</b>	<b>16</b>

**Subject Name: Artificial Intelligence in Human Health and Diseases**

**Subject Code: ODAIHG111T**

(Semester-I)

**Time: 03 Hours**

**Max.Marks:100Marks**

**Internal Assessment: 30 Marks**

**End Term: 70 Marks**

**Instructions for the Paper-Setter/examiner:**

1. Question paper shall consist of **Four sections**.
2. Paper setter shall set **Eight questions** in all by selecting **Two questions** of equal marks from each section. However, a question may have sub-parts (not exceeding four sub-parts) and appropriate allocation of marks should be done for each sub-part.
3. Candidates shall attempt **Five questions** in all, by at least selecting **One question** from each section and the **5<sup>th</sup> question** may be attempted from any of the **Four sections**.
4. The question paper should be strictly according to the instructions mentioned above. In no case a question should be asked outside the syllabus.

**Section A**

Structure of Nucleic acids and Protein; Gene expression and regulation, Genetic variations: SNPs, Mutations, copy number variations.

**Section B**

DNA Sequencing technologies: traditional and Next generation sequencing (NGS), Role of Artificial Intelligence (AI) in NGS, Role of AI in functional genomics and multi-omics.

**Section C**

Artificial Intelligence based diagnosis/screening for different diseases, biomarkers discovery and prediction of disease susceptibility.

**Section D**

Artificial Intelligence in prophylaxis, management and treatment of diseases, vaccine development: conventional to Artificial Intelligence approach.

**Recommended Books:**

1. Gardner, E.J. (2011). Human Genetics. Viva Books Pvt. Ltd., India.
2. Lodish, H., Baltimore, D., Berk, A., Zipursky, S.L., Matsudaira, P. and Daniell, J. (2007). Molecular Cell Biology. W.H. Freeman and Co., San Francisco.
3. Strachan, T. and Read, A. (2010). Human Molecular Genetics. Garland Publishers, London.

**Web Links:**

[10.1007/s44163-023-00049-5](https://doi.org/10.1007/s44163-023-00049-5)

[10.1007/s12652-021-03612-z](https://doi.org/10.1007/s12652-021-03612-z)

[10.1016/j.annonc.2023.10.125](https://doi.org/10.1016/j.annonc.2023.10.125)

[10.1016/j.cels.2021.06.006](https://doi.org/10.1016/j.cels.2021.06.006)

[10.3390/cells10040787](https://doi.org/10.3390/cells10040787)

[10.1016/j.trecan.2024.12.001](https://doi.org/10.1016/j.trecan.2024.12.001)

[10.1016/j.mimet.2024.106998](https://doi.org/10.1016/j.mimet.2024.106998)

[10.3390/biom14050568](https://doi.org/10.3390/biom14050568)

[10.1002/cpmb.59](https://doi.org/10.1002/cpmb.59)

[10.1517/17460441.2012.660145](https://doi.org/10.1517/17460441.2012.660145)

**Subject Name: Artificial Intelligence in Genetic Screening, Diagnosis and Drug Development**

**Subject Code: ODAIHG112T**

(Semester-I)

**Time: 03 Hours**

**Max.Marks:100Marks**

**Internal Assessment: 30 Marks**

**End Term: 70 Marks**

**Instructions for the Paper-Setter/examiner:**

1. Question paper shall consist of **Four sections**.
2. Paper setter shall set **Eight questions** in all by selecting **Two questions** of equal marks from each section. However, a question may have sub-parts (not exceeding four sub-parts) and appropriate allocation of marks should be done for each sub-part.
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### **Section A**

Types of genetic screening: Reproductive, Predictive, Diagnostic, Carrier, Direct-to-Consumer, AI in Newborn Screening, Prenatal Testing & Carrier Screening

### **Section B**

Overview of Drug discovery pipeline, Basics of Pharmacogenomics, Pharmacogenomic testing.

### **Section C**

AI in Drug Target Identification & Lead Optimization, AI-Based Drug Repurposing, Datasets for AI-driven Drug Discovery

### **Section D**

Clinical Decision Support Systems (CDSS) for Pharmacogenomics, AI in Clinical Trials, Regulatory guidelines for AI-driven Pharmacogenomics

### **Web Links**

doi: 10.1016/j.drudis.2020.10.010

doi: 10.1208/s12249-024-02901-y

doi: 10.1002/cpt.2387

doi.org/10.1016/j.ejphar.2024.177103

<https://doi.org/10.1186/s13073-019-0689-8>

<https://doi.org/10.1016/j.ipha.2024.08.005>

**Subject Name: Artificial Intelligence in Gene Therapy and Gene Editing**

**Subject Code: ODAIHG113T**

(Semester-I)

**Time: 03 Hours**

**Max. Marks:100 Marks**

**Internal Assessment: 30 Marks**

**End Term: 70 Marks**

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2. Paper setter shall set **Eight questions** in all by selecting **Two questions** of equal marks from each section. However, a question may have sub-parts (not exceeding four sub-parts) and appropriate allocation of marks should be done for each sub-part.
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#### **Section A**

Basics of Gene Therapy, Somatic and Germline Gene Editing, Gene Editing Tools, Role of AI in Gene Editing

#### **Section B**

Stem Cells, Cancer Stem Cells, B stem cell therapy, Basics of Regenerative Medicine and Role of AI.

#### **Section C**

Basics of Cancer genomics, Cancer risk prediction and screening, Role of AI in cancer genomics and medicine

#### **Section D**

AI and emerging concepts in Human Genetics, Digital Twins, AI-Powered Rare Disease Genomics

#### **Web Links:**

Human genome editing: Science, ethics, and governance. National Academies

Press.<https://nap.nationalacademies.org/catalog/24623/human-genome-editing-science-ethics-and-governance>

<https://pmc.ncbi.nlm.nih.gov/articles/PMC10800897>

<https://pmc.ncbi.nlm.nih.gov/articles/PMC6373233/>

<https://www.sciencedirect.com/science/article/pii/S2949820124000249>

<https://www.sciencedirect.com/science/article/abs/pii/S1359644623001216>

**Subject Name: Artificial Intelligence and Ethical Considerations in Human Genetics**

**Subject Code: ODAIHG114T**

(Semester-I)

**Time: 03 Hours**

**Max. Marks:100 Marks**

**Internal Assessment: 30 Marks**

**End Term: 70 Marks**

**Instructions for the Paper-Setter/examiner:**

1. Question paper shall consist of **Four sections**.
2. Paper setter shall set **Eight questions** in all by selecting **Two questions** of equal marks from each section. However, a question may have sub-parts (not exceeding four sub-parts) and appropriate allocation of marks should be done for each sub-part.
3. Candidates shall attempt **Five questions** in all, by at least selecting **One question** from each section and the **5<sup>th</sup> question** may be attempted from any of the **Four sections**.
4. The question paper should be strictly according to the instructions mentioned above. In no case a question should be asked outside the syllabus.

### **Section A**

Principles of Bioethics: Autonomy, Beneficence, Justice, Informed Consent, Privacy and Confidentiality

### **Section B**

Introduction to Genetic Privacy, Legal frameworks governing genetic privacy (e.g., GDPR, HIPAA, GINA), Genetic discrimination in employment and insurance settings

### **Section C**

Eugenics; Old versus Liberal, AI and Human Enhancement through liberal eugenics, Ethical Concerns

### **Section D**

Genetic Counseling, Principles, Role of AI in genetic counseling, Ethical concerns with AI and disease risk estimation and genomic analysis

### **Web Links:**

<https://pubmed.ncbi.nlm.nih.gov/38344556/>

<https://pmc.ncbi.nlm.nih.gov/articles/PMC7923912/>

<https://pmc.ncbi.nlm.nih.gov/articles/PMC4233176/>

<https://www.tandfonline.com/doi/full/10.1080/14636778.2021.2007064>

<https://www.nature.com/articles/s41431-024-01782-w>