TRAINING BROCHURE

How We Work & How You Will Be Trained

At our center, we provide comprehensive training on **DNA Profiling**, the gold standard in forensic DNA analysis. This brochure outlines our step-by-step process and how you will be trained to become proficient in DNA profiling.

Our program covers theoretical foundations, hands-on laboratory experience, and forensic case study applications to ensure you are well-equipped for real-world forensic DNA analysis.

TRAINING MODULES & WORKFLOW

1. SAMPLE COLLECTION & PRESERVATION

৺ Understanding Biological Samples:

- Blood, saliva (buccal swabs), semen, hair, bone, and tissue samples
- Sample handling techniques to avoid contamination

Collection Methods:

- Post-Mortem sample collection and storage
- Sample Collection from living subjects

♥ Preservation & Storage:

- Use of sterile collection kits
- Temperature-controlled storage methods
- Sample labeling and documentation for traceability

Q *Training Focus:* Hands-on practice with collection kits, labeling, and documentation



2. DNA EXTRACTION

Extraction Methods:

- Organic (Phenol-Chloroform)
- Automatic DNA Extraction
- Silica column-based extraction
- Magnetic bead-based purification

♦ Steps of DNA Extraction:

- 1. Lysis: Breaking open cells to release DNA
- 2. **Protein Removal:** Eliminating proteins using organic solvents or protease treatment
- 3. **Purification:** Using alcohol precipitation or column-based methods
- 4. **Storage:** Storing extracted DNA at -20°C to -80°C

Q *Training Focus:* Hands-on extraction of DNA from biological samples

3. DNA QUANTIFICATION

V Purpose of Quantification:

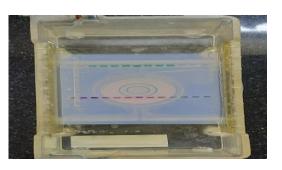
- Assessing DNA quality and quantity before PCR
- Ensuring optimal DNA input for amplification

Techniques Used:

- **Spectrophotometry** (NanoDrop): Measures DNA concentration based on absorbance at 260/280 nm
- Agarose Gel Electrophoresis: Uses EtBr to quantify DNA
- **Quantitative PCR (qPCR):** Determines human-specific DNA quantity for STR profiling

Q *Training Focus:* Practical experience using NanoDrop, Gel Electrophoresis, and qPCR for DNA quantification





4. POLYMERASE CHAIN REACTION (PCR) AMPLIFICATION

Principles of PCR:

- Denaturation, Annealing, and Extension
- Multiplex PCR for amplifying multiple STR loci

♥ STR Kits Used:

• 27 STR Marker Kit

PCR Setup:

- DNA template, primers, buffer, dNTPs, polymerase
- Thermal cycling conditions for STR amplification
- PCR contamination control

Q *Training Focus:* PCR setup, reagent preparation, and thermal cycling

5. CAPILLARY ELECTROPHORESIS (CE) & STR ANALYSIS

✓ Introduction to Capillary Electrophoresis:

- Separation of DNA fragments based on size
- Use of fluorescent dyes for STR detection

Equipment Used:

- ABI 3500/3500xL Genetic Analyzer
- Sample preparation and loading

⊘ Data Analysis:

- Interpretation of electropherograms
- Identifying peaks, stutter, artifacts, and allelic dropout

Q *Training Focus:* Operating CE instruments and analyzing STR profiles using GeneMapper software





6. STR DATA INTERPRETATION & QUALITY CONTROL

Vunderstanding Electropherograms:

- Peak height ratios, allelic ladders, and stutter analysis
- Interpretation of homozygous vs. heterozygous peaks

Ensuring Data Accuracy:

- Internal positive and negative controls
- Addressing contamination and drop-in/drop-out artifacts



V DNA Database:

• UMID-AIIMS DNA Database

Q *Training Focus:* Comparative analysis of known and unknown STR profiles

HANDS-ON TRAINING EXPERIENCE

Our training includes practical exercises such as:

- \checkmark DNA extraction from real biological samples
- $\operatorname{\mathscr{D}}$ STR amplification and analysis using CE
- **V** Contamination control techniques
- \checkmark Data interpretation and reporting
- **Simulated case studies**

≤ Join us and become an expert in DNA analysis!