

DNA as the Genetic Material Cheat Sheet

by UmeshJagtap via cheatography.com/186232/cs/43907/

Four Key Criteria for Genetic Material:

Inform- Contains instructions to build an organism.

ation:

Replic- Capable of accurate copying (DNA replication).

ation:

Transm- Passed from parent to offspring and between cells

ission: during division.

Variation: Accounts for differences within and between species.

Discovery of Genetic Material:

Early August Weismann and Karl Nägeli proposed a Hypotheses biochemical basis for inheritance.

(Late 1800s):

Chromosome Chromosomes, composed of proteins and DNA,

Insight: identified as carriers of genetic information.

Griffith's Bacterial Transformation Experiments:

Background: ✓ Type S (smooth, virulent) strains produce a

polysaccharide capsule.

✓ Type R (rough, non-virulent) strains lack this capsule.

Experimental Steps:

Step Injected live type R bacteria into a mouse → Mouse survived,

1: no live bacteria found.

Step Injected live type S bacteria into a mouse → Mouse died, live

2: type S bacteria found in blood.

Step Injected heat-killed type S bacteria into a mouse → Mouse

3: survived, no live bacteria found.

Step Mixed heat-killed type S with live type R bacteria → Injected

into a mouse → Mouse died, live type S bacteria found in blood.

Conclusion:

4:

☐ Genetic material from heat-killed type S bacteria transformed live type R bacteria.

☐ This phenomenon was called **"transformation"** without knowing the biochemical nature of the transforming substance.

Transformation Concept:

Living type R bacteria transformed into type S, gaining the ability to produce a capsule.

This transformation indicated transfer of genetic material.

Avery, MacLeod, and McCarty

Focus:Investigated bacterial transformation, following up on Griffith's observations to identify the biochemical nature of the genetic material.

Experimental Approach:

Question: What substance from dead type S bacteria transforms live type R bacteria?

Purification Process: ☐ Purified macromolecules (proteins, DNA, RNA) from type S *Streptococcus pneumoniae*.

☐ Found only purified DNA could convert type R to type S bacteria initially.

Detailed Experiment:

Step Mixed purified DNA from type S bacteria with type R

1: bacteria.

Allowed DNA uptake by type R bacteria, converting some to type S.

Step Enzyme Treatments:

2: DNase: Digests DNA.

 \square RNase: Digests RNA.

☐ Protease: Digests proteins.

Step Aggregated type R cells (non-transformed) removed by

3: centrifugation.

Step Type S cells (transformed) remain in the supernatant.

4:

Step Supernatant plated on growth media to observe bacterial

5: colony formation.

Step Control plates (without DNA extract) showed no type S

6: colonies

Conclusion:

☐ DNA from type S bacteria alone could convert type R bacteria to type S, proving DNA as the genetic material.

☐ Elimination of transformation with DNase confirmed DNA's essential role.



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Significance

protein, is the genetic material.

Impact: The experiment provided convincing evidence that DNA, not

Scientific Legacy: This study was crucial in establishing DNA's role

in heredity, greatly influencing molecular biology.

Hershey and Chase Experiment

Researchers: Alfred Hershey and Martha Chase (1952)

Objective: To determine whether DNA or protein is the genetic material in the T2 bacteriophage, a virus that infects E. coli.

□ Virus Structure Components:

- ✓ Capsid (phage coat): Made entirely of protein, consisting of a head, sheath, tail fibers, and base plate.
- ✓ DNA: Found inside the head of the capsid. Simplicity: Composed of only DNA and proteins.

Experimental Design

Goal: To identify which component, DNA or protein, enters the bacterial cell and directs the synthesis of new viruses.

Key Insight: T2 phage injects its genetic material into the bacterial cell while the protein coat remains outside.

Methodology

Labeling: □ DNA labeled with 32P (radioactive phosphorus).

☐ Protein labeled with 35S (radioactive sulfur).

Infection E. coli cells are infected with either 32P-labeled phage

or 35S-labeled phage. Process:

Use a blender to detach phage coats from bacterial Shearing Force: cells after allowing the phages to inject their genetic

Centrifug-Separate heavier bacterial cells (pellet) from lighter

ation: phage coats (supernatant).

Detection: Measure the radioactivity in the pellet and supernatant

using a Geiger counter.

☐ 35S (Protein): Majority found in the supernatant. ☐ 32P (DNA): Majority found in the bacterial pellet.

Conclusion:

DNA enters the bacterial cell, not protein. This indicates that DNA is the genetic material responsible for the production of new viruses.

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