

### Four Key Criteria for Genetic Material:

<b>Information:</b>	Contains instructions to build an organism.
<b>Replication:</b>	Capable of accurate copying (DNA replication).
<b>Transmission:</b>	Passed from parent to offspring and between cells during division.
<b>Variation:</b>	Accounts for differences within and between species.

### Discovery of Genetic Material:

<b>Early Hypotheses (Late 1800s):</b>	August Weismann and Karl Nägeli proposed a biochemical basis for inheritance.
<b>Chromosome Insight:</b>	Chromosomes, composed of proteins and DNA, identified as carriers of genetic information.

### Griffith's Bacterial Transformation Experiments:

<b>Experiment Background:</b>	<input type="checkbox"/> Studied <i>Streptococcus pneumoniae</i> . <input checked="" type="checkbox"/> Type S (smooth, virulent) strains produce a polysaccharide capsule. <input checked="" type="checkbox"/> Type R (rough, non-virulent) strains lack this capsule.
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### Experimental Steps:

Step 1:	Injected live type R bacteria into a mouse → Mouse survived, no live bacteria found.
Step 2:	Injected live type S bacteria into a mouse → Mouse died, live type S bacteria found in blood.
Step 3:	Injected heat-killed type S bacteria into a mouse → Mouse survived, no live bacteria found.
Step 4:	Mixed heat-killed type S with live type R bacteria → Injected into a mouse → Mouse died, live type S bacteria found in blood.

### Conclusion:

- 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 1:	Mixed purified DNA from type S bacteria with type R bacteria. Allowed DNA uptake by type R bacteria, converting some to type S.
Step 2:	Enzyme Treatments : <input type="checkbox"/> DNase: Digests DNA. <input type="checkbox"/> RNase: Digests RNA. <input type="checkbox"/> Protease: Digests proteins.
Step 3:	Aggregated type R cells (non-transformed) removed by centrifugation.
Step 4:	Type S cells (transformed) remain in the supernatant.
Step 5:	Supernatant plated on growth media to observe bacterial colony formation.
Step 6:	Control plates (without DNA extract) showed no type S 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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### 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.

### Significance

**Impact:** The experiment provided convincing evidence that DNA, not protein, is the genetic material.

**Scientific Legacy:** This study was crucial in establishing DNA's role in heredity, greatly influencing molecular biology.

### 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  $^{32}\text{P}$  (radioactive phosphorus).  
 Protein labeled with  $^{35}\text{S}$  (radioactive sulfur).

**Infection Process:** *E. coli* cells are infected with either  $^{32}\text{P}$ -labeled phage or  $^{35}\text{S}$ -labeled phage.

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

**Centrifugation:** Separate heavier bacterial cells (pellet) from lighter phage coats (supernatant).

**Detection:** Measure the radioactivity in the pellet and supernatant using a Geiger counter.

### Results

$^{35}\text{S}$  (Protein): Majority found in the supernatant.

$^{32}\text{P}$  (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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