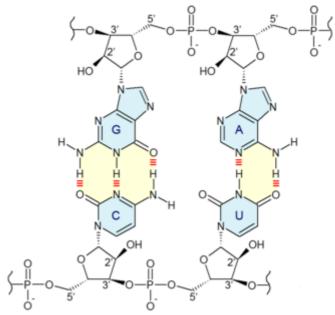
Nucleobase

Nucleobases, also known as *nitrogenous bases* or often simply *bases*, are nitrogen-containing biological compounds that form <u>nucleosides</u>, which, in turn, are components of <u>nucleotides</u>, with all of these <u>monomers</u> constituting the basic building blocks of <u>nucleic acids</u>. The ability of nucleobases to form <u>base pairs</u> and to stack one upon another leads directly to long-chain helical structures such as <u>ribonucleic acid</u> (RNA) and deoxyribonucleic acid (DNA).

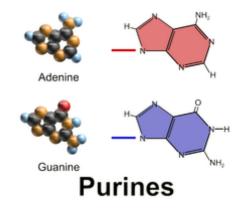
Five nucleobases—<u>adenine</u> (A), <u>cytosine</u> (C), <u>guanine</u> (G), <u>thymine</u> (T), and <u>uracil</u> (U)—are called *primary* or *canonical*. They function as the fundamental units of the <u>genetic code</u>, with the bases A, G, C, and T being found in DNA while A, G, C, and U are found in RNA. Thymine and uracil are distinguished by merely the presence or absence of a methyl group on the fifth carbon (C5) of these heterocyclic six-membered rings. [1]

Adenine and guanine have a <u>fused-ring</u> skeletal structure derived of <u>purine</u>, hence they are called **purine bases**. The purine nitrogenous bases are

characterized by their single <u>amino group</u> (NH2), at the C6 carbon in adenine and C2 in guanine. Similarly, the simple-ring structure of cytosine, uracil, and thymine is derived of <u>pyrimidine</u>, so those three bases are called the **pyrimidine bases**. Each of the base pairs in a typical double-<u>helix</u> DNA comprises a purine and a pyrimidine: either an A paired with a T or a C paired with a G. These purine-pyrimidine pairs, which are called <u>base complements</u>, connect the two strands of the helix and are often compared to the rungs of a ladder. The pairing of purines and pyrimidines may result, in part, from dimensional constraints, as this combination enables a geometry of constant width for the DNA spiral helix. The A-T and C-G pairings function to form double or triple <u>hydrogen bonds</u> between the <u>amine</u> and <u>carbonyl</u> groups on the complementary bases.



Base pairing: Two <u>base pairs</u> are produced by four nucleotide monomers, nucleobases are *in blue*. Guanine (G) is paired with cytosine (C) via *three* <u>hydrogen bonds</u>, *in red*. Adenine (A) is paired with uracil (U) via *two* hydrogen bonds, *in red*.



Purine nucleobases are fused-ring molecules.

Nucleobases such as adenine, guanine, <u>xanthine</u>, <u>hypoxanthine</u>, purine, <u>2,6-diaminopurine</u>, and 6,8-diaminopurine may have formed in outer space as well as on earth. [3][4][5]

The origin of the term <u>base</u> reflects these compounds' chemical properties in <u>acid–base reactions</u>, but those properties are not especially important for understanding most of the biological functions of nucleobases.

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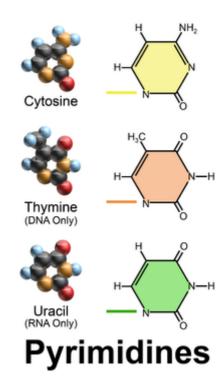
See also

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Structure

At the sides of nucleic acid structure, phosphate molecules successively connect the two sugar-rings of two adjacent nucleotide monomers, thereby creating a long chain biomolecule. These chain-joins of phosphates with sugars (ribose or deoxyribose) create the "backbone" strands for a single- or double helix biomolecule. In the double helix of DNA, the two strands are oriented chemically in opposite directions, which permits base pairing by providing complementarity between the two bases, and which is essential for replication of or transcription of the encoded information found in DNA.



Pyrimidine nucleobases are simple ring molecules.

Modified nucleobases

DNA and RNA also contain other (non-primary) bases that have been modified after the nucleic acid chain has been formed. In DNA, the most common modified base is $\underline{5\text{-methylcytosine}}$ (m⁵C). In RNA, there are many modified bases, including those contained in the nucleosides $\underline{pseudouridine}$ ($\underline{\Psi}$), $\underline{dihydrouridine}$ (D), inosine (I), and 7-methylguanosine (m⁷G). $\underline{[6][7]}$

<u>Hypoxanthine</u> and <u>xanthine</u> are two of the many bases created through <u>mutagen</u> presence, both of them through <u>deamination</u> (replacement of the amine-group with a carbonyl-group). Hypoxanthine is produced from adenine, xanthine from guanine, [8] and uracil results from deamination of cytosine.

Modified purine nucleobases

These are examples of modified adenosine or guanosine.

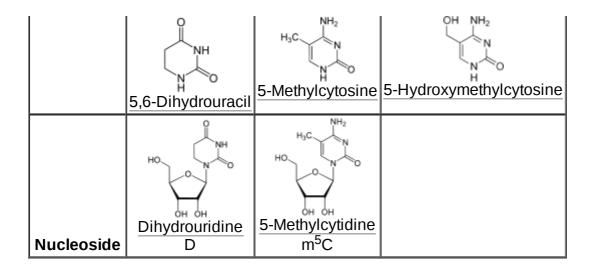
Chemical structure of DNA, showing four nucleobase pairs produced by eight nucleotides: adenine (A) is joined to thymine (T), and guanine (G) is joined to cytosine (C). + This structure also shows the <u>directionality</u> of each of the two phosphate-deoxyribose backbones, or strands. The 5' to 3' (*read* "5 prime to 3 prime") directions are: *down* the strand on the left, and *up* the strand on the right. The strands twist around each other to form a double helix structure.

	S Z Z	0 Z Z Z Z Z Z Z Z Z Z Z Z Z Z Z Z Z Z Z	N NH NH ₂
Nucleobase	Hypoxanthine	<u>Xanthine</u>	7-Methylguanine
	CH20H NH	CH ₂ OH NH OH OH Xanthosine	Ho NH ₂ NH ₂ NH ₂ 7-Methylguanosine
Nucleoside		Xantilosine	m ⁷ G

Modified pyrimidine nucleobases

These are examples of modified cytosine, thymine or uridine.

Nucleobase



Artificial nucleobases

A vast number of nucleobase analogues exist. The most common applications are used as fluorescent probes, either directly or indirectly, such as <u>aminoallyl nucleotide</u>, which are used to label cRNA or cDNA in <u>microarrays</u>. Several groups are working on alternative "extra" base pairs to extend the genetic code, such as <u>isoguanine</u> and <u>isocytosine</u> or the fluorescent <u>2-amino-6-(2-thienyl)purine</u> and <u>pyrrole-2-carbaldehyde</u>. [9][10]

In medicine, several <u>nucleoside analogues</u> are used as anticancer and antiviral agents. The viral polymerase incorporates these compounds with non-canonical bases. These compounds are activated in the cells by being converted into nucleotides; they are administered as <u>nucleosides</u> as charged nucleotides cannot easily cross cell membranes. [11] At least one set of new base pairs has been announced as of May 2014. [12]

Prebiotic condensation of nucleobases with ribose

In order to understand how <u>life</u> arose knowledge is required of chemical pathways that permit formation of the key building blocks of life under plausible <u>prebiotic conditions</u>. According to the <u>RNA world</u> hypothesis free-floating <u>ribonucleotides</u> were present in the primitive soup. These were the fundamental molecules that combined in series to form <u>RNA</u>. Molecules as complex as RNA must have arisen form small molecules whose reactivity was governed by physico-chemical processes. RNA is composed of <u>purine</u> and <u>pyrimidine</u> nucleotides, both of which are necessary for reliable information transfer, and thus Darwinian <u>evolution</u>. Nam et al. [13] demonstrated the direct condensation of nucleobases with ribose to give ribonucleosides in aqueous microdroplets, a key step leading to RNA formation. Similar results were obtained by Becker et al. [14]

See also

- Nitrogenous base
- Nucleoside
- Nucleotide
- Nucleic acid notation
- Nucleic acid sequence

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External links

■ Base pairing in DNA Double Helix (shows specific hydrogen bonds) (https://web.archive.org/web/20060621234655/http://www.elmhurst.edu/~chm/vchembook/582dnadoublehelix.html)

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