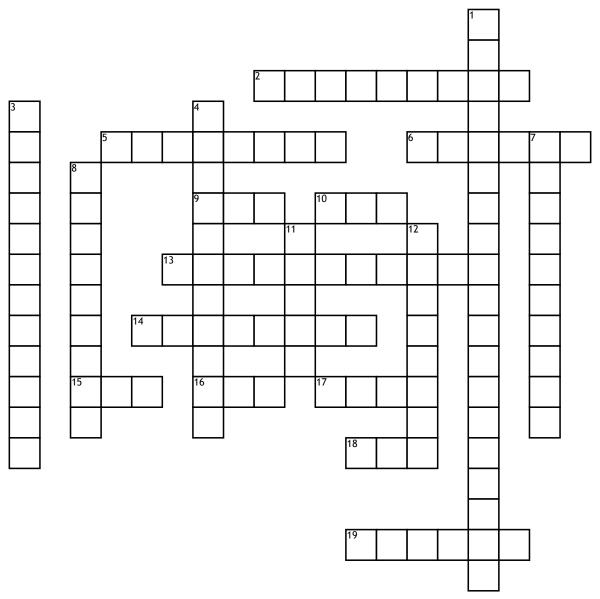
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## Chapter Two Crossword



## **Across**

- ${\bf 2.}\ {\bf A}\ {\bf change}\ {\bf in}\ {\bf an}\ {\bf original}\ {\bf act}\ {\bf or}\ {\bf law}$
- Federal- and state-operated insurance program that covers health care costs and prescription drugs for low-income children, adults, and elderly and those with disabilities
- ${\bf 6.}$  Standards concerning or relating to what is right or wrong in human behavior
- $\boldsymbol{9.}$  State board that regulates the practice of pharmacy within the state (abbreviation
- 10. The agency within the US Department of Health and Human Services that is responsible for ensuring the safety, efficacy, and security of human and veterinary drugs, biological products, medical devices, the national food supply, cosmetics, and radioactive products (abbreviation)
- 13. A drug derived from barbituric acid; a barbiturate acts as a central nervous system depressant. Barbiturates are often used in the treatment of seizures and as sedative and hypnotic agents
- 14. Federal- and state-managed insurance program that covers health care costs and prescription drugs for individuals older than 65, persons younger than 65 with long-term disabilities, and individuals with end-stage renal disease

- 15. A term used to describe a patient's personal health data. Under HIPAA, this information is protected from being shared or distributed without permission (abbreviation)
- **16.** A 10-digit number that indicates specifics of a prescription drug or an insulin product. The NDC specifies the drug manufacturer, the drug product (drug strength, dosage form, and formulation), and the package size
- **17.** An act that causes harm or injury to a person intentionally or because of negligence
- **18.** Describes medication that can be purchased without a prescription; nonlegend medications (abbreviation)
- 19. The values and morals used within a profession
- 1. Any drug or other substance that is scheduled I through V and regulated by the Drug Enforcement Administration
- 3. Drug warning that is placed in the prescribing information or package insert of the product and indicates a significant risk of potentially dangerous side effects

- 4. Labeling of a product that is false or misleading; label information must include directions for use; safe and/or unsafe dosages; manufacturer, packer, or distributor; quantity; and weight
- 7. Drug that requires a prescription for dispensing; these drugs carry the federal legend: "Federal law prohibits the dispensing of this medication without a prescription"
- 8. Comprehensive information on a medication's actions within that class of drugs. Also lists generic and trade names, ingredients, dosages, side effects, adverse effects, how the patient should take the medication, and foods or other drugs (eg, over-the-counter [OTC] medications, herbals) to avoid while taking the medication.
- 11. Federal act that protects patients' rights, establishes national standards for electronic health care communication, and ensures the security and privacy of health data
- 12. A nonspecific term used to describe a drug (such as opium) that in moderate doses dulls the senses, relieves pain, and induces profound sleep but in excessive doses causes stupor, coma, or convulsions and may lead to addiction.