

Circle the letter of the correct answer choice.

1. List the 5 most abundant elements on earth?

Oxygen Silicon aluminum iron calcium

2. List the 5 most abundant elements in the human body?

Oxygen Carbon Hydrogen Nitrogen Calcium

3. State the Law of Constant Composition.

A compound always contains the same fixed ratio of elements.

4. State the Law of Conservation of Matter

Matter can NOT be created nor destroyed; only rearranged

5. State the Law of Conservation of Energy

Energy can NOT be created nor destroyed; only transferred

6. The law of constant composition applies to

A. heterogeneous mixtures

B. homogeneous mixtures

B. metalloids

D. compounds

E. metals

7. Define Compound.

a unique substance created by the chemical combination of 2 or more elements in a fixed ratio

8. Define Element

a substance that cannot be further broken down into a <sup>more simple substance</sup>

9. How many total hydrogen atoms are indicated by the formula  $H_2C_3H_4O_2$ ?

A. 4

B. 6

C. 12

D. 16

E. 20

10. The chemical formula  $Al_2O_3$  indicates

A. six atoms of each element

B. five atoms of each element

C. three atoms of aluminum and two atoms of oxygen

D. two atoms of aluminum and three atoms of oxygen

# Chapter 4 Modern Chemistry Test

**Robert C. Smoot, Smoot, Richard G.  
Smith, Jack Price**

## Chapter 4 Modern Chemistry Test:

**Modern Methods of Drug Discovery** Alexander Hillisch, Rolf Hilgenfeld, 2012-11-28 Research in the pharmaceutical industry today is in many respects quite different from what it used to be only fifteen years ago There have been dramatic changes in approaches for identifying new chemical entities with a desired biological activity While chemical modification of existing leads was the most important approach in the 1970s and 1980s high throughput screening and structure based design are now major players among a multitude of methods used in drug discovery Quite often companies favor one of these relatively new approaches over the other e.g screening over rational design or vice versa but we believe that an intelligent and concerted use of several or all methods currently available to drug discovery will be more successful in the medium term What has changed most significantly in the past few years is the time available for identifying new chemical entities Because of the high costs of drug discovery projects pressure for maximum success in the shortest possible time is higher than ever In addition the multidisciplinary character of the field is much more pronounced today than it used to be As a consequence researchers and project managers in the pharmaceutical industry should have a solid knowledge of the more important methods available to drug discovery because it is the rapidly and intelligently combined use of these which will determine the success or failure of preclinical projects

**Laboratory Experiments to Accompany "Modern Chemistry,"** Fredus Nelson Peters, 1902 **Lab Experiments for Modern Chemistry** Tzimopoulo, 1990 **Modern Chemistry** Nicholas D. Tzimopoulos, 1990 **Elementary Modern Chemistry** Wilhelm Ostwald, Harry Wheeler Morse, 1909 **Lab Experiments Modern Chemistry** Harold Clark Metcalfe, 1982 *Handbook of Isolation and Characterization of Impurities in Pharmaceuticals* Satinder Ahuja, Karen Mills Alsante, 2003-07-18 The United States Food and Drug Administration FDA and other regulatory bodies around the world require that impurities in drug substance and drug product levels recommended by the International Conference on Harmonisation ICH be isolated and characterized Identifying process related impurities and degradation products also helps us to understand the production of impurities and assists in defining degradation mechanisms When this process is performed at an early stage there is ample time to address various aspects of drug development to prevent or control the production of impurities and degradation products well before the regulatory filing and thus assure production of a high quality drug product This book therefore has been designed to meet the need for a reference text on the complex process of isolation and characterization of process related synthesis and formulation impurities and degradation products to meet critical regulatory requirements Its objective is to provide guidance on isolating and characterizing impurities of pharmaceuticals such as drug candidates drug substances and drug products The book outlines impurity identification processes and will be a key resource document for impurity analysis isolation synthesis and characterization Provides valuable information on isolation and characterization of impurities Gives a regulatory perspective on the subject Describes various considerations involved in meeting regulatory requirements Discusses various sources of

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wellbore combined with artificial lift techniques such as pumps which bring the oil to the surface But only about 10 percent  
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