Name \_\_\_\_\_ date: \_\_\_\_\_ hr: \_\_\_\_

Viewing Guide for The Medici: Godfathers of the Renaissance Program 1: "The Birth of a Dynasty"

Medici Gulfalan of the Romanno

Use the list below to fill in the blanks. Use each term only once.

Albizzi buildings arches classical artists manuscripts bribing columns bricks concrete Cosimo Donatello Brunelleschi Florence gears

General council heresy John XXIII Pater Patriae patronage perspective pope power & prestige republic

- The Renaissance begins when Cosimo de Medici and his friends search Europe for \_\_\_\_\_\_.
   Simply reading pagan authors like Socrates and Plato was punishable by excommunication from the church.
- Popes could excommunicate (throw from the church and damn forever) Christians guilty of \_\_\_\_\_\_, which was believing anything other than what the church preached.
- who built the dome of the Florence Cathedral, was both architect and engineer.
- Brunelleschi used as supports for first time in 1000 years, creating a revolution in architecture.
- Although Brunelleschi examined the construction of the dome of the Roman Pantheon, he couldn't use the same techniques because of the size of the dome and because the recipe for making \_\_\_\_\_\_had been lost.
- Brunelleschi also devised a way to alter the \_\_\_\_\_\_ on pulleys so the oxen could pull the 1700-pound sandstone beams 250 feet into the air and return them to the ground without changing direction.
- Brunelleschi personally laid some of the \_\_\_\_\_\_ on the dome because what he proposed was so revolutionary that the brick masons were afraid the technique would fail and they would die.
- 9. Cosimo's patronage of Brunelleschi helped the Medici family gain \_\_\_\_\_\_\_
- The banking family, who resented the power of the Medici, had arrested.
- Brunelleschi was jailed and forced to stop work on Il Duomo (the dome) when his patron was found guilty of treason against

# **Medici Viewing Guide Answers Program 4**

**Dimitris Tsoukalas, Christos Tsamis** 

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arrhythmia many are detected too late in non clinical studies to inform clinicians On the other hand negative findings in chronic toxicity studies in animals may turn out to be irrelevant for human beings New scientific approaches e g high throughput screening human pluripotent stem cells transgenic animals knock out animals in silico models pharmaco genomics and pharmaco proteomics as well as Artificial Intelligence AI methods offered new possibilities There are several examples that show that the druggability of compounds was considerably underestimated when the probability of success of a new project was assessed The success rate in the pharmaceutical industry and the introduction of new chemical entities to the market per year dropped dramatically whereas the development time for a new compound increased sometimes exceeding the patent protection Research and development scientists involving the following changes therefore adopted a change of strategy Parallel instead of sequential involvement of the various disciplines multidimensional compound optimization The term Safety Pharmacology was coined The International Conference on Harmonization ICH founded a Safety Pharmacology Working Group and the Safety Pharmacology Society SPS was launched The discipline provided for evaluation development and validation of a multitude of safety tests outlined in the Core Battery of Studies Characterizing the exposure profile of a drug by conducting pharmacokinetic studies that evaluates the absorption distribution metabolism and excretion should to be investigated at an early stage of development as results contribute to the selection of a compound for further development Advancements in Toxicology were achieved by the introduction of new methods e g in silico methods genetic toxicology computational toxicology and AI The book is a landmark in the continuously changing world of drug research and developments As such it is essential reading for many groups not only for all students of pharmacology and toxicology but also for industry scientists and physicians especially those involved in clinical trials of drugs and for pharmacists who must know the safety requirements of drugs The book is essential for scientists and managers in the pharmaceutical industry who are involved in drug discovery drug development and decision making in the development process In particular the book will be of use to government institutions and committees working on official guidelines for drug evaluation worldwide Catalog of Copyright Entries. Third Series Library of Congress. Copyright Office, 1958 Includes Part 1 Number 1 2 Books and Pamphlets Including Serials and Contributions to Periodicals January December

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