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In the initial edition of Williams Obstetrics, very few fetal disorders could be identified before delivery. Now, more than 100 years later, prenatal diagnosis has become a separate field of its own. Strictly speaking, prenatal diagnosis is the science of identifying congenital abnormalities, aneuploidies, and other genetic syndromes in the fetus. It encompasses the diagnosis of structural malformations with specialized sonography; routine screening tests for aneuploidy and neural-tube defects;

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Prenatal care undoubtedly contributed to the dramatic decline in this mortality rate from 690 deaths per 100,000 births in 1920 to 50 per 100,000 by 1955 (Loudon, 1992). And, the low current maternal mortality rate of 10 to 15 per 100,000 is likely associated with the high utilization of this care .

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Since the early 1990s, the largest gains in timely prenatal care have been among minority groups. As shown in Figure 9-1, however, disparity continues. Of the 27 states using the revised birth certificate, the percentage of non-Hispanic white, Hispanic, and African-American women who received no prenatal care in 2008 was 1.1, 2.7, and 3.3, respectively (Osterman, 2011).

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It encompasses routine screening tests for aneuploidy and neural-tube defects, invasive diagnostic tests such as chorionic villus sampling and amniocentesis, additional screening and diagnostic tests offered to those at risk for specific genetic disorders, and the diagnosis of structural malformations with specialized sonography and other fetal imaging techniques discussed in Chapter 10. The goal of prenatal diagnosis is to provide accurate information regarding short- and long-term ...

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Fetal imaging of congenital anomalies is discussed in Chapter 10, and pregnancy termination is discussed in Chapter 18. HISTORICAL PERSPECTIVE More than 40 years ago, Brock (1972, 1973) observed that pregnancies complicated by neural-tube defects had higher levels of alpha-fetoprotein (AFP) in maternal serum and amniotic fluid.

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