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The clinical decision-making skills essential for effective respiratory care practice across the continuum of health care. Features actual clinical cases for critical analysis, patient assessment, clinical application, decision-making, and discussion.

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Also addresses
managed care,
health care costs,
ethics, and
alternative site
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Implementation,
analysis,
interpretation, and
quality evaluation of
registries created
to increase
understanding of
patient outcomes.
For the purposes of
this guide, a patient
registry is an
organized system
that uses
observational study

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Methods to collect uniform data (clinical and other) to evaluate specified outcomes for a population defined by a particular disease, condition, or exposure, and that serves one or more predetermined scientific, clinical, or policy purposes.

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A registry database is a file (or files) derived from the registry. Although registries can serve many purposes, this guide focuses on registries created for one or more of the following purposes: to describe the natural history of disease, to determine clinical

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effectiveness or
cost-effectiveness
of health care
products and
services, to
measure or monitor
safety and harm,
and/or to measure
quality of care.
Registries are
classified according
to how their
populations are
defined. For

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Example, product registries include patients who have been exposed to biopharmaceutical products or medical devices. Health services registries consist of patients who have had a common procedure, clinical encounter, or hospitalization. Disease or

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Condition registries are defined by patients having the same diagnosis, such as cystic fibrosis or heart failure. The User ' s Guide was created by researchers affiliated with AHRQ ' s Effective Health Care Program, particularly those

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