

**Table 1: Typical Processes for Managing Clinical Decision Support**

Processes	Description	Keys to Success
Agenda setting/ targets	<ul style="list-style-type: none"> <li>Individuals and committees request new application of CDS to support specific quality or safety objective.</li> <li>A steering committee reviews and prioritizes requests.</li> <li>Major changes to clinical policy or practice referred to Medical Executive Committee or other group for approval.</li> </ul>	<ul style="list-style-type: none"> <li>Accountability for CDS linked with governance of medical practice (assigned to chief medical officer or other physician leader such as medical director of patient safety).</li> <li>Review and approval by appropriate accountable clinical leader or group.</li> <li>Effective communication and coordination among all individuals and groups.</li> </ul>
Setup and testing	<ul style="list-style-type: none"> <li>Analysts in IS setup and test new CDS in development system.</li> <li>One or more physicians may test new CDS on a provisional basis.</li> </ul>	<ul style="list-style-type: none"> <li>Ability to set up and test new tools in other than the operational system.</li> <li>Status tracking (e.g., development, testing, release) and audit trail for CDS tools.</li> <li>Ability to release tools on a limited basis.</li> </ul>
Review	<ul style="list-style-type: none"> <li>Steering committee reviews and approves test.</li> <li>May require sign-off of Pharmacy and Therapeutics Committee or department chair.</li> <li>Some hospitals require physician sign-off on personal order sets.</li> </ul>	<ul style="list-style-type: none"> <li>Formal accountability for different targets of CDS tools (medications, disease state).</li> </ul>
Disseminate in operational system	<ul style="list-style-type: none"> <li>New order sets available immediately.</li> <li>Batches of new CDS released at regular system updates.</li> <li>CDS addressing major (dangerous or high risk) situations released immediately.</li> <li>Physician community notified of major new CDS in advance and necessary training provided.</li> <li>Collect metrics (baseline if needed) to measure effectiveness.</li> </ul>	<ul style="list-style-type: none"> <li>Effective processes for communicating with physicians about major updates (usually multiple modes are used).</li> </ul>
Evaluate and update	<ul style="list-style-type: none"> <li>Responsibility of committee authority.</li> <li>Review each application of CDS periodically to validate currency of clinical content or update as necessary.</li> <li>Monitoring of physician response to implemented CDS (acceptance, override).</li> <li>Physician feedback solicited.</li> <li>Collect metrics on targets of CDS and make changes as appropriate based on findings.</li> </ul>	<ul style="list-style-type: none"> <li>Automated tracking of ownership, clinical research base, and update schedule for each “rule” or type of CDS (e.g., medication checking).</li> <li>Easy mechanisms (two-way) for physicians to provide feedback on form and/or content of CDS.</li> <li>Commitment to respond to each physician suggestion or complaint.</li> </ul>