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T e FDA def nes pati M M e

tools mailed directly to the patient or transmitted through a patient portal for those with the electronic medical record capacity to do so. On the date of the visit, PROs may be assessed in the waiting room via paper surveys, tablet computers, or computer kiosks. The tools may also be incorporated during the rooming process, administered by medical assistants or nurses, as an additional “vital sign” [6]. Once in the exam room, clinicians may administer PROs as paper or web-based tools or may enter them directly into the EMR. Immediately after the visit, patients may be given a survey to return at the next visit.

Additionally, PRO assessments may be measured longitudinally, to

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