



## Other Benefits and Contextual Considerations

5. Does treating patients with one of the new FDA approved drugs offer one or more of the following “other benefits”? (select all that apply)

<input type="checkbox"/> This intervention provides significant direct patient health benefits that are not adequately captured by the QALY.
<input type="checkbox"/> This intervention offers reduced complexity that will significantly improve patient outcomes.
<input type="checkbox"/> This intervention will reduce important health disparities across racial, ethnic, gender, socioeconomic, or regional categories.
<input type="checkbox"/> This intervention will significantly reduce caregiver or broader family burden.
<input type="checkbox"/> This intervention offers a novel mechanism of action or approach that will allow successful treatment of many patients who have failed other available treatments.
<input type="checkbox"/> This intervention will have a significant impact on improving return to work and/or overall productivity.

6. Are any of the following contextual considerations important in assessing the new FDA approved drug’s long-term value for money in patients with tardive dyskinesia? (select all that apply)

<input type="checkbox"/> This intervention is intended for the care of individuals with a condition of particularly high severity in terms of impact on length of life and/or quality of life.
<input type="checkbox"/> This intervention is intended for the care of individuals with a condition that represents a particularly high lifetime burden of illness.
<input type="checkbox"/> This intervention is the first to offer any improvement for patients with this condition.
<input type="checkbox"/> Compared to standard therapy there is significant uncertainty about the long-term risk of serious side effects of this intervention.
<input type="checkbox"/> Compared to standard therapy, there is significant uncertainty about the magnitude or durability of the long-term benefits of this intervention.