Table S3: Subject-reported onset of effect in the 7-day diary (mITT population)

	Cumulative proportion of subjects reporting onset in the aboBoNT-A solution group (N=250)	**Cumulative proportion of subjects reporting onset in the responder subgroup[a] of the aboBoNT-A solution group (N=219)**
Day 1	23%	24%
Day 2	49%	53% (median onset)
Day 3	64% (median onset)	68%
Day 4	75%	80%
Day 5	82%	86%
Day 6	85%	88%
Day 7	86%	90%

[a] The responder subgroup included all subjects who were none-or-mild responders for ILA at maximum frown, at Month 1.

For placebo, no median values of time to onset could be calculated due to few subjects reporting an onset of effect in the diary, i.e. 19/121 subjects (16%), mITT population. Three placebo subjects were included in the ILA-responder subgroup at Month 1.