## ePRO Information

- The CAT™ has been tested and used on a number of electronic data collection devices. A list of the devices on which CAT™ has been tested is under development and will be provided below.

- Migration of the CAT™ to a new ePRO device needs to be supported by evidence demonstrating the comparability, or measurement equivalence of the ePRO to the paper-based CAT™. Published reports and guidance are available which provide support and general frameworks for this development.[1,2] Further information on the requirements for migration can be found in the relevant section of the HCP User Guide on this website.

- If you want further ePRO information or want to use a new ePRO and would like to understand about gaining accreditation please contact us at: (CATmailbox@gsk.com )

- The CAT™ governance board will accredit new presentations of the CAT™ on e-PRO which will then be listed on the website in due course.

- Development and/or use of an accredited ePRO presentation is requested as a condition of the terms of use.

1. Coons SJ, Gwaltney CJ, Hays RD et al. Recommendations on evidence needed to support measurement equivalence between electronic and paper-based Patient-Reported Outcome (PRO) measures: ISPOR ePRO Good Research Practices Task Force Report. Value in Health 2009: 12:419-429

2. Coons SJ, Eremenco S, Lundy JJ. Capturing Patient-Reported Outcome (PRO) data electronically: the past, present, and promise of ePRO measurement in clinical trials. Patient 2014: Patient :DOI 10.1007/s40271-014-0090-z