Summit Corporation plc
(‘Summit’ or ‘the Company’)

SUMMIT TO PRESENT CLINICAL AND BIOMARKER DEVELOPMENT PROGRAMMES FOR DMD AT THE 18th INTERNATIONAL WORLD MUSCLE SOCIETY CONGRESS

Oxford, UK, 30 September 2013 – Summit (AIM: SUMM), a drug discovery and development company advancing therapies for Duchenne Muscular Dystrophy (‘DMD’) and C. difficile infection, will present a summary of the clinical and biomarker programmes for its oral utrophin modulator SMT C1100 for the treatment of DMD at the 18th International World Muscle Society Congress meeting in Asilomar, California, USA from 1-5 October 2013.

The poster presentation entitled “Future clinical and biomarker development for SMT C1100, the first utrophin modulator to enter clinical trials for Duchenne Muscular Dystrophy” will outline the proposed clinical development plan to achieve proof of concept for SMT C1100 in DMD patients. Details will also be presented on the supporting biomarker programme that aims to develop new exploratory clinical endpoints to help evaluate the benefit of therapies such as SMT C1100 in future DMD patient trials.

The presentation will be part of the “Dystrophinopathies: Imaging and biomarkers” session that will be held at 16:30 PST on Friday, 4 October. A copy of the presentation will be available on the Company’s website, www.summitplc.com from Wednesday, 2 October 2013.

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Notes to Editors

About Summit
Summit is an Oxford, UK based drug discovery and development Company targeting high-value areas of unmet medical need including Duchenne Muscular Dystrophy and C. difficile infection. Summit is listed on the AIM market of the London Stock Exchange and trades under the ticker symbol SUMM. Further information is available at www.summitplc.com and follow Summit on Twitter (@summitplc).

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Forward Looking Statements

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