BSUG RESEARCH GRANT CHECK LIST

Eligibility

□ BSUG Member

Topic

- Original research question and rational for study clearly explained
- Important topic, compatible with BSUG aims and strategy
- □ Relevant to clinical practice
- Research involving BSUG Database

Method

- \Box Clear research hypothesis
- \Box Specific aims
- □ Specific objectives
- Study methodology appropriate for hypothesis
- Appropriate design of study and use of control/placebo arm
- Appropriate Inclusion and Exclusion criteria
- □ ICS/IUGA definitions used throughout
- □ ICS/IUGA guidelines followed where appropriate
- Appropriate power calculation
- Primary and Secondary outcome measures specified and appropriate
- □ Primary Outcome appropriate for aims
- Secondary Outcome measures relevant to aims
- \Box Details of interventions to be used in the study
- \Box Study should be completed within the 12 months duration of the grant
- Details regarding appropriate statistical analysis

Feasibility

- Realistic recruitment estimate and consideration of feasibility
- \Box Number of sites
- Duration of study
- \Box Facilities for the study are available at all study sites
- Patient assessment, recruitment, intervention and follow up can be completed within the time span allotted to the project

Ethics/Registration

- Confirmation of ethical approval if required
- □ Registration with ISRCTN
- Suitable for NIHR Portfolio registration
- Evidence of GCP training

Centre

- Adequate patient numbers to achieve recruitment target
- Adequate experience to met the study requirements

- Track record in completing studies and presentation/publication
- □ Clinical trial experience
- CV of Principal and Co-investigators documenting research experience
- Involvement of Research Fellow and registration for higher degree

Documentation

- CV of Principle Investigator an Co-investigators
- Centre information demonstrating ability to recruit patients and complete the study
- Ethical approval documentation
- Trial registration documentation
- Trial protocol
- □ Patient consent form
- □ Patient information form
- □ Data collection forms
- Questionnaires/Dairies used in the study

Cost

- □ Inclusion of detailed budget for study
- □ Reasonable and realistic cost for materials and staff

Presentation

- Clear succinct and methodological abstract
- Clear justification of the study, study design and aims
- \Box Clear methodology
- Clear plans for statistical analysis
- Plans for presentation and distribution of the results
- Appropriate and contemporary references

Name:

Signature:

Date: