

STRUCTURAL GUIDELINES FOR CIRSE QUALITY ASSURANCE DOCUMENTS AND STANDARDS

<u>Please note: all CIRSE Quality Assurance Documents and Standards released after 1st April, 2011 must adhere to the following structural guidelines.</u>

Introduction

This section should include a historical evolution of IR treatment/methods over the past years.

Definitions

This section should include relevant definitions regarding anatomy, clinical symptoms and signs, treatment methods, etc.

Pre-treatment Imaging

Indications for Treatment (if applicable also divided into absolute and relative) and Contraindications

Patient Preparation

Equipment Specifications

Recommendations for specific companies/products should be avoided.

Procedural Features (it is important to describe the technique with most accumulated evidence) **and Variations of the Technique(s)**

The level of evidence for each variation should be reported.

Medication and Peri-procedural Care

Post-procedural Follow-up Care (Including Imaging)

Outcome

Recommended thresholds for technical success, clinical success, and complications should be provided.

a. **Effectiveness** (include *clinical* as well as *technical* success)

Randomised clinical comparisons with competing surgical or conservative treatment should be included. If randomised study is not available, at least controlled trials should be mentioned.

b. Complications (immediate and long-term) and their Management

Conclusions

References

APPENDIX (Classification of complications by outcome)

Minor Complications

- a. No therapy, no consequence.
- b. Nominal therapy, no consequence; includes overnight admission for observation only.

Major Complications

- c. Require therapy, minor hospitalisation (<48 hours).
- d. Require major therapy, unplanned increase in level of care, prolonged hospitalisation(>48 hours).
- e. Permanent adverse.
- f. Death.