

# Planning an EQA Scheme



The following list ("Minimum requirement list") has been prepared by the IFCC Committee on Analytical Quality to assist countries to plan and design an external quality assurance scheme. The C-AQ can provide assistance in this area. Contact details can be found at <http://www.ifcc.org/ifcc-education-division/emd->

|                           |  |  |
|---------------------------|--|--|
| <b>Organisation</b>       | <b>Responsible organisation</b>                                  |  |
|                           | <b>Name of EQA Scheme</b>  |  |
|                           | <b>Managerial Staff</b>  |  |
|                           | <b>Technical Staff</b>   |  |
|                           | <b>Administrative Staff</b>                                      |  |
|                           | <b>Accommodation</b>   |  |
|                           | <b>Laboratory Space</b>  |  |
|                           | <b>Equipment Required</b>  |  |
|                           | <b>Participation costs</b>                                       |  |
| <b>Planning</b>           | <b>What discipline/s?</b>  |  |
|                           | <b>What measurands (analytes)?</b>                               |  |
|                           | <b>How many participants?</b>                                    |  |
|                           | <b>What frequency of testing?</b>                                |  |
| <b>Material</b>           | <b>What is the specimen matrix ( eg human serum, urine)?</b>     |  |
|                           | <b>What is the specimen type (eg fresh frozen, lyophilised)?</b> |  |
|                           | <b>How will the material be sourced?</b>                         |  |
|                           | <b>What concentration ranges?</b>                                |  |
|                           | <b>Volume?</b>   |  |
|                           | <b>Reference values?</b>   |  |
|                           | <b>Homogeneity &amp; Stability</b>                               |  |
|                           | <b>Vials &amp; labels</b>  |  |
| <b>Specimen Packaging</b> |  |  |
| <b>Paperwork</b>          | <b>Design &amp; print paperwork</b>                              |  |
|                           | <b>Instructions for participants</b>                             |  |
|                           | <b>Set closing dates</b>   |  |
| <b>Results</b>            | <b>Electronic or paper result submission</b>                     |  |
|                           | <b>Database</b>  |  |
|                           | <b>Statistical design</b>  |  |
|                           | <b>Acceptance limits</b>   |  |
|                           | <b>Criteria for evaluation</b>                                   |  |
| <b>Reports</b>            | <b>Design report</b>   |  |
|                           | <b>Paper or electronic</b>                                       |  |
| <b>Quality System</b>     | <b>Document all aspects</b>                                      |  |
| <b>Recommendation</b>     | <b>Accreditation to ISO/IEC 17043:2010 in the long term</b>      |  |