July 27, 2018

TO: IAS-ACCREDITED FABRICATOR INSPECTION PROGRAMS FOR STRUCTURAL STEEL, INSPECTION AGENCIES AND OTHER INTERESTED PARTIES

SUBJECT: Proposed Revisions to the Accreditation Criteria for Fabricator Inspection Programs for Structural Steel, Subject AC172-0918-0918-R1 (DM/SM)

Hearing Information:
IAS Accreditation Committee
Thursday, September 27, 2018
8:00 a.m.
Fullerton Marriott at California State University
2701 Nutwood Avenue
Fullerton, CA 92831
(714) 738-7800

Dear Madam or Sir:

The proposed IAS Accreditation Criteria for Fabricator Inspection Programs for Structural Steel, AC172, has been placed on the agenda for committee consideration at the above-noted meeting. The changes are being requested to accommodate a recent request for changes and for changes to clarify specific areas of the criteria.

The changes proposed are summarized below:

1. A universal change throughout the criteria to change “quality system” to “management system.”

2. Section 1.2 Overview, was added to stay in line with the other criteria for steel fabrication and manufacturing. Two items to take notice of are as follows:
   a. Move the requirements of the inspection agency to the Rules of Procedure and Annex A of these criteria.
   b. Add verbiage that matches with the requirements in Section 1704.2.5.1 of the International Building Code (IBC).
3. Add the following standards to Section 1.3 Normative and Reference Documents and make editorial revisions:
   c. American Iron and Steel Institute: AISI S100: North American Specification for the Design of Cold-Formed Steel Structural Members.

4. Add the definition for Nondestructive Testing (NDT) to Section 2.8 of the criteria.

5. Remove “in concert with an IAS accredited inspection agency” in Section 4.2.1.2 and all of Section 4.2.1.4. The requirements of these sections are being redefined in the Rules of Procedure and Annex A of these criteria.

6. Remove EN-287-1 in Section 4.2.4.1 and Section 4.3.8. This standard has been replaced by ISO 9606-1.

7. Add “inspectors” in the list of items that need to be traced on the finished product in Section 4.5.4.

8. Add to the Note in Section 4.5.4 to clarify the need for heat numbers in the traceability of the main member’s web and flanges.

9. Add Annex A to define the requirement of the inspection agency.

You are cordially invited to submit written comments, or to attend the committee hearing and present verbal comments. Written comments will be forwarded to the committee, prior to the hearing, if received by September 6, 2018. Please use the comment form link found on the Accreditation Committee meeting page on the IAS website, www.iasonline.org. Comments may be postal mailed to the address above, or emailed to iasinfo@iasonline.org.

Any written material submitted for committee consideration will be available for public distribution as set forth in Section 4.0 of the Rules of Procedure for Accreditation Committee Meetings (copy enclosed).

Visual aids (including, but not limited to, charts, slides, videos, or presentation software) for viewing at meetings will be permitted only if the presenter provides to IAS, before the presentation, a copy of the visual aid(s) in a medium that can be retained by IAS with its record of the meeting, and that can also be provided to interested parties.

Your cooperation is requested in forwarding to the Brea office, as noted above, all material directed to the committee. Prior to the hearing, parties interested in the deliberations of the committee should refrain from communicating, whether in writing or
verbally, with committee members regarding agenda items. The committee reserves the right to refuse communications that do not comply with this request.

If you have any questions, please contact Sandi McCracken, senior program manager, at 562-364-8201, extension 3442, or the undersigned at 562-364-8201. You may also reach us by e-mail at iasinfo@iasonline.org.

Yours very truly,

[Signature]

Raj Nathan
President

RN/nl

Enclosures

cc: Accreditation Committee
RULES OF PROCEDURE FOR ACCREDITATION COMMITTEE MEETINGS

1.0 PURPOSE

The purpose of the Accreditation Committee and its meetings is to safeguard IAS’ impartiality to monitor the work of and to approve accreditation criteria for International Accreditation Service, Inc. (IAS).

The committee meetings, which are open public hearings, provide an opportunity for effective involvement by all interested parties.

2.0 MEETINGS

2.1 The Accreditation Committee shall schedule meetings that are open to the public in discharging its duties under Section 1, subject to Section 5.0 of these rules.

2.2 To properly discharge its responsibilities with respect to monitoring of IAS accreditation activities, the committee shall have a standing item on its meeting agenda for a presentation by staff on the status of its accredited programs and information on any pending appeals.

2.3 All scheduled meetings shall be publicly announced.

2.4 A majority of the voting Accreditation Committee members shall constitute a quorum. A majority vote of members present is required on any action.

2.5 If a specific interest group is not represented, votes by the committee on subjects related to that interest group will be held in abeyance. IAS staff shall make pertinent information available to absentee committee members, and ballot the members at a later stage. Records of such ballots shall be made available upon request.

2.6 In the absence of the nonvoting Chair-Moderator, Accreditation Committee members present shall elect an alternate Chairman from the committee for that meeting. The alternate Chairman shall be counted as a voting committee member for purposes of maintaining a committee quorum and to cast a tie-breaking vote of the committee.

2.7 Minutes of the meetings shall be kept.

3.0 MEMBER COMPETENCE CRITERIA

Members of the Accreditation Committee shall be familiar with conformity assessment and the implementation of regulatory requirements within their industry sector. They shall possess:

• A Baccalaureate degree from an accredited institution or a minimum of ten years equivalent experience as determined by IAS;

• Current employment within the conformity assessment, regulatory field, academia, industry, or IAS accredited CAB; and

• Demonstrated expertise in one or more accreditation programs offered by IAS.
4.0 MEETING RECORDS

An electronic record of meetings shall be made by IAS; no other audio, video, electronic or stenographic recordings of the meetings will be permitted. Visual aids (including, but not limited to, charts, slides, videos, or presentation software) viewed at meetings shall be permitted only if the presenter provides IAS before presentation with a copy of the visual aid in a medium which can be retained by IAS with its record of the meeting and which can also be provided to interested parties requesting a copy. A copy of the IAS recording of the meeting and such visual aids, if any, will be available to interested parties upon written request made to IAS together with a payment as required by IAS to cover costs of preparation and duplication of the copy. These materials will be available shortly after the conclusion of the meeting but will no longer be available after 60 days have elapsed from the conclusion of the meeting.

5.0 WRITTEN COMMUNICATIONS AND SUBMISSIONS

Parties interested in the deliberations of the committee should refrain from communicating, whether in writing or verbally, with committee members regarding agenda items. All written communications and submissions regarding agenda items should be delivered to IAS. All such written communications and submissions shall be considered nonconfidential and available for discussion in open session of an Accreditation Committee meeting, and shall be delivered at least twenty days before the scheduled Accreditation Committee meeting if they are to be forwarded to the Committee. Correspondence received by IAS will not be released to any party, except to the Accreditation Committee, prior to the meeting without permission of the author. The committee reserves the right to refuse recognition of communications which do not comply with the provisions of this section. All such communications and submissions will be available from IAS upon written request and payment of costs associated with duplication. The materials will be available shortly after the conclusion of the meeting but will no longer be available after 60 days have elapsed from the conclusion of the meeting.

6.0 CLOSED SESSIONS

Meetings shall be open except that the chairman may call for a closed session to seek advice of counsel.

7.0 ACCREDITATION CRITERIA

Criteria are established by the committee to provide a basis for International Accreditation Service, Inc., accreditations. Consideration of accreditation criteria must be in conjunction with a current and valid application for an IAS accreditation listing or as otherwise determined by the Accreditation Committee.

7.1 Procedure

7.1.1 New Criteria

7.1.1.1 Proposed accreditation criteria may be submitted by interested parties to IAS, and/or shall be developed by the IAS staff and discussed in open session with the Accreditation Committee during a scheduled meeting

7.1.1.2 Proposed accreditation criteria shall be available to interested parties approximately 60 days before discussion at the committee meeting, unless determined by IAS management that extraordinary consideration and electronic balloting are needed.

7.1.1.3 The committee shall be informed of all pertinent written communications received by IAS. Parties interested in proposed new criteria may deliver communications and submissions regarding such proposed criteria to IAS within 40 days of the posting of the public notice on the IAS website. Such communications and submissions will otherwise be subject to the provisions of Section 4.0 of these rules.

7.1.1.4 Attendees at Accreditation Committee meetings shall have the opportunity to speak on accreditation criteria listed on the meeting agenda, to provide information to committee members.
7.1.2 Existing Criteria

7.1.2.1 Changes to existing accreditation criteria may be submitted by interested parties to IAS, and/or shall be
changed by the IAS staff. Existing accreditation criteria may be revised by the committee either (i) at a public meeting pursuant
to the procedures set forth herein, or (ii) by electronic ballot, provided public notice is provided as stipulated in Section 7.1.1.2.

7.1.2.2 The committee shall be informed of all pertinent written communications received by IAS. Parties interested in
the proposed revisions to accreditation criteria may deliver communications and submissions regarding such proposed
revisions to IAS within the following timelines:

<table>
<thead>
<tr>
<th>Type</th>
<th>Dates</th>
</tr>
</thead>
<tbody>
<tr>
<td>Public Meeting</td>
<td>40 Days after posting of proposed criteria</td>
</tr>
<tr>
<td>Electronic Balloting Process</td>
<td>30 Days after posting of proposed criteria</td>
</tr>
</tbody>
</table>

Such communications and submissions will otherwise be subject to the provisions of Section 4.0 of these rules.

7.1.3 ELECTRONIC BALLOTING

7.1.3.1 IAS management shall provide written rationale and seek permission and documented approval from the IAS
Accreditation Committee chair to propose new criteria or to revise existing criteria for extraordinary consideration and
electronic balloting by the committee.

7.1.3.2 Proposed accreditation criteria shall be available to interested parties approximately 30 days before
consideration by the committee. All pertinent written communications received by IAS relating to the proposed criteria shall be
received no later than 30 days after the posting of the criteria. Ballots, along with comments received and staff
recommendations, will be submitted to the committee for consideration. The committee shall return their ballots with their
recommendations within 10 days from the date ballots are sent. The results of the balloting will be compiled and forwarded to
the chair of the committee for validation and decision.

7.1.3.3 The electronically balloted criteria shall be brought back to the next regularly scheduled accreditation
committee hearing as per Section 7.1.2 of these rules,

7.1.4 Effective Date of Published Criteria

7.1.4.1 The effective date of approved accreditation criteria or approved revisions to existing accreditation criteria shall
be no earlier than 30 days following the public meeting.

7.1.4.2 Approved criteria using electronic balloting shall be effective the date of posting of the criteria on the IAS
website.

7.2 Approval

Approval of accreditation criteria shall be as specified in Section 2.4 of these rules.

8.0 ACCREDITATION COMMITTEE MEMBERS
8.1 The IAS Accreditation Committee members are appointed or reappointed annually by the IAS Board of Directors in consultation with the IAS President.

8.2 Committee members are selected from senior management positions within accredited organizations, users of accreditation, industry groups and governmental or regulatory organizations. The individuals appointed to the committee shall have knowledge of regulatory codes within their industry sector and international conformity assessment process and practices.
PROPOSED REVISIONS TO THE ACCREDITATION CRITERIA FOR FABRICATOR INSPECTION PROGRAMS FOR STRUCTURAL STEEL

AC172

Proposed September 2018

PREFACE

The attached accreditation criteria have been proposed to provide all interested parties with an opportunity to comment. These criteria may be further revised as needed. The criteria are developed and adopted following public hearings conducted by the International Accreditation Service, Inc. (IAS), Accreditation Committee and are effective on the first of the month following approval by the Accreditation Committee, but no earlier than 30 days following the approval.
1. INTRODUCTION

1.1. Scope: These criteria set forth the requirements for obtaining and maintaining International Accreditation Service, Inc. (IAS), Fabricator Inspection Programs for Structural Steel accreditation. These criteria supplement the IAS Rules of Procedure for Accreditation of Fabricator Inspection Programs.

1.2. Overview: Accredited entities complying with these criteria will have demonstrated that they have the personnel, organization, experience, knowledge, quality procedures and commitment to fabricate in accordance with specified requirements. IAS-accredited inspection programs for manufacturers of metal building systems operate under a documented management system developed comply with these criteria. The management system includes the manufacturer's written fabrication procedures and quality control manuals that provide a basis for control of materials and workmanship, with periodic inspections of fabrication and quality control practices by an IAS-accredited inspection agency. Responsibilities and requirements for inspection agencies are documented in Annex A. Although accredited entities are evaluated on their performance measures to consistently produce products of the required quality mandated by specified requirements, these criteria do not cover the products or the design or performance characteristics of the products.

1.2.1.3. Normative and Reference Documents: Publications listed below refer to current editions (unless otherwise stated).

1.2.1.3.2. IAS Rules of Procedure for Accreditation of Fabricator Inspection Programs.
1.2.1.3.3. American Welding Society: AWS D1.1, AWS D1.3, AWS D1.4, AASHTO/AWS D1.5 and AWS D1.8 Structural Welding Code.
1.2.1.3.6. American Welding Society: AWS QC1, Standard for AWS Certification of Welding Inspectors.
1.2.1.3.7. Canadian Standards Association: CSA W178.2, Certification of welding inspectors.
1.2.1.3.8. The Society for Protective Coatings (SSPC):
2. DEFINITIONS

For the purposes of these accreditation criteria, the definitions given in the ISO/IEC Standard 17000 series, and the definitions that follow, apply.

2.1. **Approved Fabricator**: An established and qualified person, firm or corporation approved by the building official pursuant to the *International Building Code®*, published by the International Code Council.

2.2. **Contract Documents**: Documents that describe the fabricator's responsibilities for a given project. These documents include work orders, drawings, and project specifications.

2.3. **Corrective Action**: Implemented action of solutions necessary to eliminate or reduce the root cause of an identified problem.

2.4. **DAR (Designated Accreditation Representative)**: A quality professional, designated by the fabricator who has demonstrated competence in managing and implementing a quality management system with consistent results.

2.5. **DARD (Designated Accreditation Representative Deputy)**: An employee designated by the fabricator who has demonstrated competence in managing and implementing the fabricator’s quality management system during a temporary absence of the DAR.

   **Note**: Reference Appendix A of AC172 for the requirements of the Designated Accreditation Representative.

2.6. **Management System**: A set of interrelated or interacting elements that organizations use to direct, control and coordinate how policies are implemented and objectives are achieved.

2.7. **Nonconformance**: An action employed that renders a member or component unacceptable for the intended use as specified in contract specifications or these criteria.
2.8. Nondestructive Testing (NDT): The process of inspecting, testing, or evaluating materials, components or assemblies for discontinuities, or differences in characteristics without destroying the serviceability of the part or system.

2.7.2.9. PQR: Procedure Qualification Record in accordance with AWS or AASHTO/AWS Standards, as applicable.

2.8.2.10. Procedure: An implemented and written document that describes who does what, when, where, why and how.

2.9.2.11. Product: Result of activities or processes.

2.10.2.12. Project: A process consisting of a set of coordinated and controlled activities undertaken to achieve customer requirements.

2.11.2.13. Quality Assurance: Measurable systematic actions to assure confidence that the implementation of planned activities result in meeting objectives, goals and project specifications.

2.12.2.14. Quality Control: The act of examination, testing or measurement that verifies processes, services or that documents conform to specified criteria.

2.13.2.15. Quality Plan: A written document prepared by the designated accreditation representative that describes the procedures and policies implemented to assure product quality meets specific contract documents. As a minimum, quality plans must meet the requirements of AC172.

2.14.2.16. Repair: Action taken to render a member or component acceptable for the intended use.

2.15.2.17. Scope of Accreditation: Specific conformity assessment services for which accreditation is sought or has been granted.

2.16.2.18. Specification: A document that states the obligatory requirements the product must conform to.

2.17.2.19. Steel Construction, Cold-formed: That type of construction made up entirely or in part of steel structural members cold formed to shape from sheet or strip steel such as roof deck, floor and wall panels, studs, floor joists, roof joists and other structural elements.

2.18.2.20. Steel Element, Structural: Any steel structural member of a building or structure consisting of rolled shapes, pipe, hollow structural sections, plates, bars, sheets, rods or steel castings other than cold-formed steel or steel joist members.

2.19.2.21. Steel Joist: Any steel structural member of a building or structure made of hot-rolled or cold-formed solid or open-web sections, or riveted or welded bard strip or sheet steel members, or slotted and expanded, or otherwise deformed rolled sections.

2.20.2.22. WPS: Welding Procedure Specification in accordance with American Welding Society AWS D1.1, AWS-D1.3, AWS-D1.4, or AASHTO/AWS D1.5, and AWS-D1.8 as applicable.
3. ELIGIBILITY

Accreditation services are available to structural steel fabrication inspection program facilities that meet the requirements of these criteria.

4. REQUIRED BASIC INFORMATION

4.1. Fabricator inspection programs for structural steel must demonstrate compliance with the following requirements:

4.1.1. The requirements of these accreditation criteria;

4.1.2. IAS Rules of Procedure for Accreditation of Fabricator Inspection Programs.

4.2. General Requirements

4.2.1. Quality System

4.2.1.1. The fabricator shall establish and implement a management system that is fully documented. This documented management system must describe the fabricator’s procedures and quality activities for ensuring that fabricated products meet the specified requirements of this criteria.

4.2.1.2. The fabricator in concert with an IAS-accredited inspection agency, shall prepare and submit to IAS its documented management system, including a cross-reference matrix ensuring that the general requirements in Section 4.2, data in Section 4.3, the statements in Section 4.4, and the written procedures noted in Section 4.5 of these accreditation criteria have been included.

4.2.1.3. The submitted management system document must be signed and dated by the highest level of authority within the organization.

4.2.1.4. The submitted management system document must be signed and dated by an authorized representative of an IAS-accredited inspection agency, attesting that the inspection agency has reviewed the fabricator’s documented management system and that the fabricator’s documented management system is sufficient to schedule an onsite joint assessment with IAS.

4.2.2. Designated Accreditation Representative: The fabricator shall designate a Designated Accreditation Representative who has the necessary training and experience to complete the tasks listed in Sections 4.2.2.1. through 4.2.2.5. The Designated Accreditation Representative shall report directly to the highest level of authority within the organization. The Designated Accreditation Representative shall have the following responsibilities:

Note: Responsibilities noted in Sections 4.2.2.1. through 4.2.2.5. may be delegated to individuals such as a quality manager, where appropriate.
4.2.2.1. Maintaining the fabricator’s documented management system in accordance with these criteria.

4.2.2.2. Monitoring the effective implementation of the fabricator’s documented management system and reporting the results to the highest level of authority annually.

4.2.2.3. Assuring that, as a minimum, annual internal audits are conducted and documented, and that corrective actions are effectively implemented.

4.2.2.4. Assuring that annual management reviews are conducted and documented to assure the adequacy and effectiveness of the management system. Annual management reviews must include a summary and a documented plan of action for improvement. Documents to be considered during the annual management review must include, but are not limited to, customer complaints, back charges, internal audit results and corrective actions.

4.2.2.5. Developing quality plans that meet project specifications, and having knowledge of and access to the appropriate documents to meet this requirement.

4.2.3. **In-house Quality Control (QC) Inspector**: The fabricator shall designate an in-house quality control inspector(s) who, as a minimum, must meet the following requirements:

4.2.3.1. Be a Certified Welding Inspector (CWI) in accordance with the provisions of AWS QC1 or the equivalent requirements of the Canadian Standards Association (CSA) Standard W178.2 or ICC Structural Steel and Bolting Special Inspector, or Structural Welding Special Inspector.

4.2.3.2. Be familiar with and demonstrate knowledge of codes and specifications, as appropriate, for the scope of work specified in the contract documents.

4.2.3.3. Be responsible for assuring that only qualified and certified welders are used, as specified by contract documents for the welding process and procedures permitted for use.

4.2.3.4. Be responsible for assuring continuity of the welders’ qualifications as required by American Welding Society [AWS] D1.1.

4.2.3.5. Be responsible for overall workmanship and for making sure that all weldments are 100% visually inspected. Although inspections may be delegated to qualified personnel during the receipt and in-process stages of assembly, it is the responsibility of the quality manager to ensure that inspections are performed and that the product meets project requirements.

4.2.3.6. Be responsible for ensuring that incoming raw materials are properly identified and inspected for compliance with quality plans and specifications.

4.2.3.7. Be responsible for ensuring and documenting that the final fabrication assembly can be traced back to the incoming materials, the quality assurance inspection records and the individual welder.
4.2.3.8. Be responsible for reviewing all Welding Procedure Specifications (WPSs) and Procedure Qualification Records (PQRs) and ensuring they are adequate before they are used in production welding operations.

**Note:** Approval of welding procedures must be obtained by the customer when specified by contract documents.

4.2.4. **Welding Personnel:** The fabricator shall ensure that the following conditions are met:

4.2.4.1. All welding personnel shall be qualified by the test as described in AWS D1.1 or D1.3, or other accepted country-specific test standard, as appropriate, by a qualified independent third-party agency. Third-party qualification shall be by certification as an AWS Certified Welding Inspector (CWI) in accordance with the provisions of AWS QC1, *Standard for AWS Certification of Welding Inspectors*, or current qualification by the appropriate Canadian Welding Bureau (CWB) to the requirements of the Canadian Standards Association Standard W178.2, *Certification of Welding Inspectors*; or current qualification by approved third-party agencies, such as those accredited by an accreditation body that is an IAS Mutual Recognition Arrangement (MRA) partner, per ISO 9606-1 or EN-287-1; or by the International Code Council as an ICC Structural Welding Special Inspector (S2). The in-house CWI, CWB, or ICC Structural Welding Special Inspector (S2) may administer the welding tests; however, the qualification coupon shall be evaluated by the third party CWI, CWB or ICC Structural Welding Special Inspector (S2). If tensile testing is required for qualification of welding personnel, the test, or test sample, must be sent to an IAS-accredited testing laboratory for examination. Such laboratories must be accredited by IAS or by an accreditation body that is a partner with IAS in an MRA.

4.2.4.2. All welding personnel shall have and use an identifying number, letter or symbol for the purpose of traceability.

4.2.5. **Bolting:** Procedures shall be developed as required in the project documents and shall address the following: Fitting, snug-tight, pre-tensioning, and faying surfaces.

**Note:** Fabricators that include high-strength bolting using ASTM A325 or ASTM A490 bolts as a fabrication practice will receive recognition on the accreditation certificate. As a minimum, there must be an ICC certified Structural Steel and Bolting Special Inspector (S1) on staff.

4.2.6. **Nondestructive Testing:** Procedures shall be developed as required in the project documents.

**Note:** Fabricators that include nondestructive testing as a fabrication practice will receive recognition on the certificate of accreditation certificate.

4.3. **Required Data**
The following information shall be included in the management system submittal:

4.3.1. The name of the fabrication facility, the physical street address, mailing address (if different), information of the person serving as the IAS contact (including the telephone number and e-mail address), and the telephone number of the fabrication facility.

4.3.2. A floor plan of the fabrication facility. The floor plan need not be to scale.

4.3.3. A list of major production equipment, including welding, burning, lifting and inspection equipment.

4.3.4. A list of typical items fabricated (e.g., beams, trusses, towers, signs, girders, etc.).

4.3.5. A copy of all WPSs for production welding. The WPSs shall be written to include essential and nonessential variables, in accordance with AWS D1.1, AWS D1.3, AASHTO/AWS D1.5, or AWS D1.8, as appropriate for the type of fabrication performed at the facility.

4.3.6. A copy of all PQRs for WPSs qualified by testing, when required. PQRs pertaining to AASHTO/AWS D1.5 must be current within the last five years. PQRs for the welding of fracture-critical members must be current within the last three years and must include the submerged arc welding process.

4.3.7. A list of qualified welding personnel, including their approved welding process, limitations to their qualifications and their identification marks.

4.3.8. Evidence that welding personnel are qualified by the test as described in AWS D1.1 or D1.3, or other accepted country-specific test standard, as appropriate, by a qualified independent third-party agency. Third-party qualification shall be by certification as an AWS Certified Welding Inspector (CWI) in accordance with the provisions of AWS QC1, Standard for AWS Certification of Welding Inspectors, or current qualification by the Canadian Welding Bureau (CWB) to the requirements of the Canadian Standards Association Standard W178.2, Certification of Welding Inspectors, or current qualification by approved third-party agencies, such as those accredited by an accreditation body that is a partner with IAS in an MRA, per ISO 9606-1 or EN-287-1; or by the International Code Council as a Structural Welding Special Inspector. The in-house CWI, CBW, or ICC Structural Welding Special Inspector may administer the welding tests; however, the qualification coupon shall be evaluated by the third party CWI, CBW or ICC Structural Welding Special Inspector. If tensile testing is required for qualification of welding personnel, the test, or test sample, must be sent to an IAS-accredited testing laboratory for examination. Such laboratories must be accredited by IAS or by an accreditation body that is a partner with IAS in an MRA.

4.3.9. The name and identifying number, letter or symbol of the in-house quality control inspector, for the purpose of traceability.
4.3.10. The name(s) of the deputy in-house QC inspector who assumes the position in the absence of the primary in-house QC person.

4.3.11. An organizational chart of the fabricator, including the names of the responsible quality manager/Designated Accreditation Representative. This chart must show the relationships among the CEO, project manager, quality manager, in-house quality control inspector, deputy in-house inspector, production manager and welding personnel.

4.3.12. A list of approved vendors, including any testing agencies employed to verify a WPS.

4.3.13. A list of test and measuring equipment.

Note: Test and measuring equipment must be calibrated and traceable to a national standard. The equipment list must include sufficient testing instruments to assure quality compliance as appropriate for the items being fabricated.

4.4. **Required Statements**

The following statements shall be provided in the management system submittal:

4.4.1. A quality policy statement that includes the following elements:

4.4.1.1. All activities of the organization shall be directed in such a manner as to ensure that the quality requirements of AC172 will be met.

4.4.1.2. The elements of the quality assurance program will be disseminated to all personnel assigned activities that affect the quality of the product.

4.4.2. The manual shall, at a minimum, be reviewed annually.

4.4.3. IAS will be notified, in writing, prior to any cancellation of the inspection agreement with the accredited inspection agency.

4.4.4. Copies of reports of inspections conducted by the inspection agency, if they note major quality control variations, will be forwarded by the fabricator to IAS within 10 days of the major deficiency being reported.

4.4.5. The fabricator will notify the inspection agency when the fabrication facility is to be closed for extended time periods other than for normally scheduled periods for maintenance or vacations or two or more weeks regardless of the circumstances of the closure. IAS and the inspection agency will be notified 10 days prior to resumption of operations.

4.4.6. IAS will be notified in writing by the fabricator and the inspection agency if unannounced, follow-up inspections have not been conducted by the inspection agency.

4.4.7. The fabricator will promptly investigate and respond to IAS or a building official when informed of complaints regarding the noncompliance of finished product with stated specifications.
4.4.8. IAS and the accredited inspection agency must be notified within 30 days of any changes in management personnel. As a minimum, this would include the president, general manager, project manager, purchasing manager, production manager, Designated Accreditation Representative, quality manager or principal engineer.

4.5. Required Written Procedures

The fabricator shall submit written procedures for the following:

4.5.1. **Contract Review:** Review of contract documents to ensure that the needed resources exist to fulfill the contract requirements. The contract review procedure must include provisions that assure the review is appropriate, that the product and service will meet the specifications and must include a provision for the approval of exceptions or change requests. Reviews shall be performed by personnel who have access to the appropriate information and have adequate knowledge of the requirements and must be approved by the quality manager/Designated Accreditation Representative.

Reference Appendix A of AC172 for the requirements of the Designated Accreditation Representative.

4.5.2. **Document Control:** Control of documents and data relating to the quality functions of the fabricator. This control must include the following:

4.5.2.1. A document approval procedure.

4.5.2.2. A procedure to ensure that only current, approved documents are used.

4.5.2.3. A procedure to ensure that documents are available at all locations where necessary for the proper functioning of the quality management system.

4.5.2.4. Information on how detail drawings are prepared and how revisions to contract documents and change orders are approved.

4.5.3. **Purchasing**

4.5.3.1. Determining that purchased products will conform to specified requirements. The procedure must include a requirement that the type and grade of material be documented on the purchase order agreement.

4.5.3.2. Evaluation of subcontractors for their ability to meet subcontract requirements. Evaluations may contain summaries or logs, but must include a means of quantifying and measuring the ability of the subcontractor or supplier to provide quality products or services consistent with the required contract documents. For projects requiring IAS accreditation, subcontract fabrication may be subcontracted only to fabrication facilities that are currently IAS-accredited.
Note: While IAS understands some organizations use the term “subcontractor” synonymously with “supplier,” there is a difference, and both suppliers and subcontractors are required to be evaluated on an annual basis.

4.5.4. **Product Traceability**: The traceability procedure must describe the method used to ensure items are traceable as specified in the contract documents. Items that typically require traceability are materials and consumables that are incorporated into the final product. The project documents will determine if full materials traceability is required, however, the fabricator must have a procedure to meet the project needs for the type of fabrication performed. In addition to project requirement needs, the fabricator, as a minimum, must have in their control traceability of the finished product to incoming materials, certified welders, inspector, plans and specifications. The procedure must make provision for documentation of this traceability on inspection forms or on a controlled copy of the detail drawing.

Note: Material traceability, unless otherwise required by contract documents, is limited to main members and does not include items such as stiffeners. All main member webs and flanges must be traced to a heat number. All secondary members such as stiffeners, clips, and bolted end plates must be traceable to the type and grade of material.

4.5.5. **Process Control**: There must be a procedure that identifies how process control is communicated to appropriate personnel. Process control includes procedures such as cutting or saw operations, fitting and welding of the material, cambering and coating. Examples of forms used in the process control procedure are cut lists, standard drawings or detail drawings. The procedure must describe the fabricator’s method of communicating and establishing priorities of such operations.

4.5.6. **Inspection and Testing**: The inspection procedure shall include provisions for receipt, in-process and final inspections as appropriate to provide a level of assurance that products are manufactured in accordance with contract documents by qualified personnel. Final inspections shall include a record of the results and resolution of nonconformances identified by subsequent inspections. As a minimum, inspection procedures include the following:

4.5.6.1. Receiving inspection of incoming materials to the required specification, including review of mill test reports and certificates of conformance to ensure compliance with contract documents.

4.5.6.2. In-process inspection for workmanship that can affect subsequent operations.
(Examples of in-process inspections are nondestructive testing of welds that will be hidden or out of reach during the final inspection, visual examination of fit-up tolerances that will not be visible after welding, areas requiring coatings that will not
be accessible during final inspection, monitoring of welding and bolting operations, as appropriate.) Welding process inspections on multiple pass welds must ensure that proper preheat and interpass temperatures are maintained, and that the finished welds are of the proper size, without flaws, undercuts, inclusions or porosity.

4.5.6.3. Final inspection includes documented acceptance of all workmanship performed, including materials, welding, bolting, fitting operations, and coatings.

All final welds are to be accepted under the direction of the in-house CWI, CWB or ICC Structural Welding Special Inspector.

4.5.7. **Control of Inspection, Measuring and Test Equipment:** There must be a maintenance schedule, including calibration procedures for testing equipment. Wherever possible, calibration services shall be provided by a calibration laboratory accredited by IAS or by an accreditation body that is a partner with IAS in an MRA. **Note:** It is recognized there may not be nationally recognized standards available for unique testing equipment. When such instances exist, calibration procedures must be in compliance with manufacturer's recommendations to the extent that such testing equipment is calibrated to ensure consistency with the required measuring capabilities. It is the fabricator's responsibility to ensure that such testing equipment is approved prior to use.

4.5.8. **Control of Nonconforming Workmanship:** Procedures shall be established for identifying, documenting and assigning the disposition of nonconforming items.

4.5.9. **Corrective Action:** Procedure for corrective action shall include investigating, documenting and correcting nonconformances. The procedure must include a provision to preclude repetition.

4.5.10. **Handling, Storage and Delivery Procedure:** Procedure shall include identifying and storing of incoming materials and finished products as appropriate to minimize damage and deterioration.

4.5.11. **Internal Audits:** The fabricator shall identify the frequency, method of documentation and the content of internal audits to determine the effectiveness of the management system. Audits shall include a summary that compares the most recent audit to the previous audit and include the elements of AC172.

4.5.12. **Control of Quality Records:** The fabricator must determine methods for storing, maintaining and accessing quality records for a minimum of two years. Quality records must include the following:


4.5.12.2. Completed in-house quality inspection reports, forms, and checklists.
4.5.12.3. Manufacturer test reports and certificates of compliance from vendors, for incoming materials and consumables.

4.5.12.4. Copies of inspection reports by the inspection agency.

4.5.12.5. Records of internal audits.

4.5.12.6. Training records.

4.5.12.7. Evaluations of vendors and subcontractors.

4.5.13. **Training:** There must be a procedure for the training of personnel who have an effect on the quality of the finished product. The procedure must include provision for maintaining current personnel qualifications. As a minimum, there must be training requirements established for project managers, detailers, inspectors, welders, fitters and painters.

**Appendix A — Qualifications for Designated Accreditation Representative**

4.6. **Scope**

International Accreditation Service, Inc. (IAS), has established a Designated Accreditation Representative (DAR) and a Designated Accreditation Representative Deputy (DARD) requirement for quality assurance and quality control (QA/QC) personnel. It is the responsibility of the fabricator to designate a DAR and a DARD as described in Sections 2.4 and 2.5 to carry out the responsibilities under Section 4.8 below.

4.7. **Introduction**

Evaluations of DAR and DARD candidates are performed during an on-site joint review of a fabricator inspection program by IAS and the fabricator’s accredited inspection agency.

4.8. **General Requirements for Designated Accreditation Representative**

4.8.1. The DAR/DARD must successfully demonstrate his/her knowledge of the quality management system and technical operations of the fabricator, including an assessment of his/her general, practical and specific knowledge pertinent to the fabricator’s current project documents.

4.8.2. The DAR must report directly to the highest level of management within the organization and must have stop-work authority.

4.8.3. The DARD will report to the DAR. In the absence of the DAR, the DARD must report directly to the highest level of management within the organization and must have stop-work authority.

4.8.4. The DAR must be able to conduct effective internal audits, identify performance indicators and recommend corrective actions. The purpose of these activities is to
evaluate the overall effectiveness of the documented management system. At a minimum, the DAR must be able to perform the duties outlined in Sections 4.8.4.1, 4.8.4.2 and 4.8.4.3.

4.8.4.1. The ability to understand trend analysis measurements. Trend analyses must clearly show the direction that an activity is taking over time, to decide if corrective action is required. For example, trend analyses may be plotted to show whether costs are increasing or decreasing, if errors are declining or increasing, or if any number of factors being measured and plotted are meeting desired quality levels.

4.8.4.2. The ability to develop, implement and document staff training.

4.8.4.3. The ability to develop and implement quality plans, including generation of appropriate documentation.

Note: Although specific assignments may be delegated to a DARD, it will be the responsibility of the DAR to determine that a fabricator’s quality-management system has been successfully executed in accordance with contract documents.

4.8.5. The DAR must demonstrate competent knowledge of structural steel fabrication and inspection practices that are pertinent to products that are manufactured by the fabricator. Mandatory knowledge may include, but is not limited to: developing and implementing procedures for detailing, procurement, bolting, welding, inspection and nondestructive testing; operational procedures that include sawing, shearing, drilling and fitting practices, coatings, packaging, handling, and shipping of structural steel and/or their components. The submitted procedures must include inspection requirements as appropriate to assure compliance and implementation.

4.8.6. Fabricators must notify IAS within 10 days of the termination of employment of the DAR. Termination of the DAR may affect the fabricator’s accreditation status with IAS until IAS has evaluated and approved the company’s DAR replacement.

4.8.7. DAR status is not transferable from one company to another. It may be suspended upon extended leave of absence or other circumstances that prevent the DAR from performing his/her duties.

4.9. Specific Requirements for Designated Accreditation Representative

The DAR must demonstrate knowledge through a combination of education, training and experience of the latest editions of established codes and standards as appropriate to the fabrication of structural steel members and their components. Applicable documents may include, but are not limited to, the following:

4.9.2. AWS D1.1, AWS D1.3 or AWS D1.8 Standards as applicable for the type of fabrication performed at the facility.

4.9.3. AWS A2.4, Symbols.

4.9.4. AWS A3.0, Terms and Definitions.

4.9.5. AISC Code of Standard Practice.

4.9.6. SSPC Painting Manual, Volume 1, Good Painting Practice.


4.9.8. AISC Detailing for Steel Construction.


4.9.10. ASTM International (relevant standards).


4.9.12. Project specifications/contract documents for the current fabrication performed at the facility.


4.10. Control of Required Procedures

4.10.1. Contract Review: The DAR must ensure that contract quality requirements are met. The DAR will be responsible for reviewing any instructions and/or procedures relative to activities affecting quality to determine if they are properly understood and implemented.

As a minimum, the following elements must be documented to ensure that contract reviews are managed, controlled, and successfully implemented and communicated to appropriate personnel:

4.10.1.1. Quality plans to ensure that fabrication conforms to the most recent project specifications. Quality plans shall include proprietary buy-out items and subcontract fabrication. Project specifications include design drawings, detail drawings, and other related documents.

4.10.1.2. At a minimum, quality plans shall address the following:

4.10.1.2.1. Material: ASTM grade and type, AWS filler metal classification

4.10.1.2.1.1. Origin of materials

4.10.1.2.1.2. Substitution requirements

4.10.1.2.1.3. Material test report requirements

4.10.1.2.2. Workmanship
4.10.1.2.2. Cutting of plates or shapes
4.10.1.2.2. Drilling or punching of holes:
  4.10.1.2.2.1. Edge distance
4.10.1.2.2.2. Repair of mislocated holes
4.10.1.2.2.3. Welding requirements:
  4.10.1.2.2.3.1. Welding procedure specifications
  4.10.1.2.2.3.2. Control consumables
4.10.1.2.2.4. Cambering, bending, straightening
4.10.1.2.2.5. Dimensional tolerances
4.10.1.2.3. Coating/painting/galvanizing:
  4.10.1.2.3.1. Surface preparation
  4.10.1.2.3.2. Manufacture and type of coating
  4.10.1.2.3.3. Application of coating
4.10.1.2.4. Required inspections and sequence of inspections to verify conformance of an item or activity to specified requirements.
  4.10.1.2.4.1. Procedures:
    4.10.1.2.4.1.1. Receiving inspection procedures
    4.10.1.2.4.1.2. In-process inspection procedures
    4.10.1.2.4.1.3. Final inspection procedures
    4.10.1.2.4.1.4. Records and reports
4.10.1.2.4.2. Nondestructive testing requirements
4.10.1.2.5. Acceptance criteria for inspections required in the contract documents for the scope of the project.
4.10.1.2.6. Shipping, packaging and handling requirements.

4.10.2. **Document Control**: The Designated Accreditation Representative shall be responsible to ensure that only current, approved documents are used and to ensure that appropriate documents are available at all locations where necessary for the proper functioning of the management system. Document control must encompass the following elements:

  4.10.2.1. Controlled receipt of bid documents, specifications and revisions.
  4.10.2.2. Approval of working (detail) drawings prior to issuing to persons using them as work instructions.
  4.10.2.3. Approval of revisions, including a method for revision control to assure the latest revision is available and used by appropriate personnel.
  4.10.2.4. Approval of change orders.
  4.10.2.5. Documentation of back charges, including the root cause of the problem.
  4.10.2.6. Records of complaints.
4.11. **Education and Experience: Designated Accreditation Representative**
Personnel shall be qualified on the basis of appropriate education, training and experience. Education and training must be such that the DAR is competent to take full charge of his/her responsibilities under the IAS DAR program. Training requirements are based on the standards referenced in Section 4.9 and Table I.

4.12. **Education and Experience: Designated Accreditation Representative Deputy**
Personnel shall be qualified on the basis of appropriate education, training and experience. Education and training must be such that the DARD is competent to take full charge of his/her responsibilities under this program. Training requirements are based on the standards referenced in Section 4.9 and Table I.

5. **ADDITIONAL INFORMATION (AS APPLICABLE)**

5.1. AWS B5.1, Specification for Qualification of Welding Inspectors.
5.2. AWS B5.17, Specification for the Qualification of Welding Fabricators.
5.3. ANSI/AISC 341, Seismic Provisions for Structural Steel Buildings.
5.4. ANSI/AISC 360, Specification for Structural Steel Buildings.
5.5. CSA W47.1 Certification of companies for fusion welding of steel.

6. **LINKS TO ADDITIONAL REFERENCES**

6.1. IAS – [www.iasonline.org](http://www.iasonline.org)
### Table 1

<table>
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<th>DAR</th>
<th>DARD</th>
<th>Topic of Training Required</th>
<th>Credits</th>
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<td>x</td>
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<td>1. Total Quality Concepts(^1)</td>
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<td>x</td>
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<td>2. Customer Satisfaction(^1,2)</td>
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<td>3. Strategic Quality Planning(^1)</td>
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<td>4. Management and Leadership(^1,2)</td>
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<td>5. Personal Communications and Interrelationship Skills(^1)</td>
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<td>6. Quality Planning and Setting Objectives(^1)</td>
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**Note:** To qualify for DAR status, an individual must accrue twenty-five (25) credits. DARD education and experience must have a minimum accumulation of fifteen (15) credits.

\(^1\) Via seminars, videos, books, self-study correspondence courses
\(^2\) Customer feedback/information benchmarking
\(^3\) Via professional activities
\(^4\) Based on shop experience
\(^5\) Hands-on inspection experience
\(^6\) Up to two (2) credits may be earned for other performance factors not explicitly called out in this matrix, such as proven leadership, sound judgment, analytical ability, tenacity and past performance.
\(^7\) From an accredited institution
\(^8\) Familiarity with AC172
\(^9\) Based on ASNT examination
The inspection agency is limited to the review and implementation of the following shop procedures:

1. Product traceability
2. Process Control
3. Inspection and Testing
4. Control of Inspection, Measuring and Test Equipment
5. Control of Nonconforming Workmanship
6. Corrective Action
7. Handling and Storage
8. Training of Shop Personnel, Welder Qualification and Inspectors