

Appendix G: Procedures and Guidelines for the Use of AOAC Voluntary Consensus Standards to Evaluate Characteristics of a Method of Analysis

Official Methods Board, Expert Review Panels, First and Final Action *Official Methods*SM

Since 2011, Expert Review Panels (ERPs) have been used to assess methods against Standard Method Performance Requirements (SMPR[®]). In 2021, after a decade of successful adoption of over one hundred Official Methods, the Official Methods Board (OMB) undertook a review to integrate and clarify the standards process, to ensure continued best practice, while recognizing and embracing flexibility to meet each community's needs as part of the Official Methods Program.

Definitions

Call for Methods: public announcement inviting method submissions for a given analyte and/or matrix.

Candidate Method: a method accepted into the Official Methods Program for possible adoption as First Action.

Evaluation Period: interval between adoption as First Action and consideration as Final Action during which further method validation or information gathering is undertaken.

First Action: AOAC First Action *Official Methods*.

Final Action: AOAC Final Action *Official Methods*.

Lead Reviewer: an ERP Voting Member charged with presenting in depth method reviews and making initial recommendations to the ERP.

Non-voting Observer: a contributing expert to scientific deliberations of ERP, however, is ineligible to vote during an ERP meeting.

Method Author: method developer or developer's representative who serves as primary contact throughout Official Method development.

Voting Member: scientific expert vetted by the OMB and selected to vote on motions as part of an ERP meeting.

OMB Oversight

The OMB serves the Association in a scientific and advisory capacity, including on the process of method adoption. As such the OMB is responsible for oversight of the Official Methods Program and ensures compliance to policies and procedures in the development of voluntary consensus standards.

See Figure 1 for process flowchart.

ERP Formation

An ERP is authorized to adopt candidate methods as First Action and to recommend subsequent adoption for Final Action status. Scientists are recruited to serve as ERP members or as ERP Chair through a public call or by recommendation by members of AOAC. Interested scientists

40 are invited to submit their curriculum vitae (CV) for initial assessment by the AOAC Science
41 Team, who then forward to the OMB evaluations and recommendations for formal review.
42 Both the Science Team and OMB strive to ensure that the composition of a proposed ERP is
43 both qualified and equitably representative of stakeholder groups. The OMB-approved ERP
44 candidates are reviewed and appointed as ERP Members by the AOAC President.

45 The Chair of an ERP serves as moderator for discussions, ensuring all relevant topics of a
46 method are adequately discussed prior to a call for a vote.

47 **ERP Requirements**

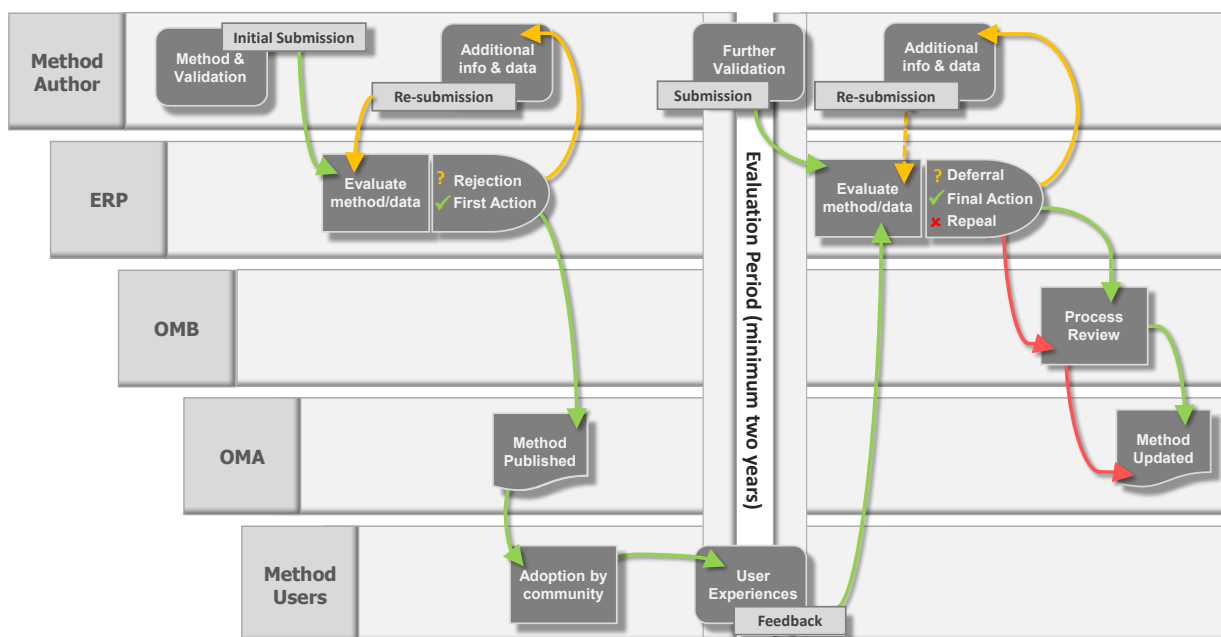
- 48 (1) When established, an ERP shall consist of a minimum of 7 Voting Members representing a
49 balance of stakeholders.
- 50 (2) A quorum is established by the presence of 7 Voting Members or 2/3 of total Voting
51 Members, whichever is greater.
- 52 (3) The ERP must hold transparent public meetings.

53 **Method Submission**

54 Methods may be submitted by the Method Author at any time, although typically during the
55 Call for Methods period. All submitted methods must be accompanied by validation data upon
56 which the ERP can undertake a comprehensive review. Various method performance
57 parameters may be required or expected by different ERPs depending upon the nature of the
58 analytes, matrices, and techniques pertinent to the method.

59 Each community will assess the necessary forms that this validation may include, such as:

- 60 • range of matrices tested
- 61 • repeatability
- 62 • reproducibility
- 63 • other inter-laboratory precision
- 64 • recovery
- 65 • comparison to reference material
- 66 • comparison to reference method
- 67 • ruggedness/robustness
- 68 • specificity/selectivity
- 69 • linearity and/or analytical range
- 70 • limits of detection and/or quantitation
- 71 • stability
- 72 • inclusivity/exclusivity
- 73 • uncertainty
- 74 • probability of detection.



75
76 **Figure 1. Process for development of voluntary consensus standards for methods of analysis**
77 **within the AOAC Official Methods Program**

78 The minimum necessary parameters may be specified by the relevant community as part of
79 SMPR development (*Official Methods of Analysis*, Appendix F: Guidelines for Standard
80 Method Performance Requirements). Acceptable experimental designs used to collect this
81 data may vary with the method protocol and the intended use of the method.

82 **ERP Voting**

83 Votes shall be cast by show of hands at in-person meetings and by roll call at virtual/remote
84 meetings.

85 **Abstentions**

86 At the beginning of an ERP meeting when the agenda is confirmed, Voting Members shall
87 declare any perceived or actual conflicts of interest to any agenda items on which a vote will
88 be called. Such a declaration need not preclude a Voting Member from voting, and at that
89 time, the ERP Chair will rule whether any Voting Member must abstain from voting on that
90 particular agenda item based upon this declaration. All Voting Members and Non-voting
91 Observers, whether they have a declared conflict or not, are freely able to share their
92 expertise during the discussion period prior to the First Ballot.

93 Voting Members may need to abstain on motions pertaining to: (i) methods they have
94 authored or co-authored; (ii) methods from entities with which they are affiliated; or (iii)
95 methods from other entities in which a conflict of interest has been identified.

96 Abstentions are not counted as a “yes” or “no” vote, but instead are a non-vote and
97 contribute only toward establishing a quorum.

98 **Duty to Vote**

99 Unless required by the Chair to abstain, as experts Voting Members are expected to vote on
100 all motions. Any Voting Member who abstains on grounds other than a declared conflict of

101 interest should delineate reasons. Where the number of abstentions exceeds 1/3 of the
102 Voting Members present, the vote is declared invalid and must be retaken at a later date.

103 ***First Ballot***

104 (1) A motion shall pass the First Ballot only by unanimous affirmative vote of the ERP.

105 (2) A motion shall fail if negative votes exceed 1/3 of the vote.

106 (3) If neither (1) or (2) is achieved, scientific reasons must be delineated for negative votes.

107 Following further discussion, a second ballot is taken.

108 ***Second Ballot***

109 After further discussion and consideration, and the motion shall pass in a Second Ballot by
110 2/3 or greater affirmative vote.

111 **Review of Methods for First Action**

112 Methods submitted to AOAC are collected and compiled by the Science Team and are
113 categorized as Candidate Methods and assigned a unique identifier.

114 An ERP meeting, open to all interested parties, is convened to review Candidate Method(s).
115 Two (or more) ERP members may be assigned by the ERP Chair as Lead Reviewers to provide
116 in depth, written reviews and to make a recommendation for First Action adoption, as
117 appropriate. The merits and deficiencies of the Candidate Method are reviewed and
118 discussed by the ERP, using the relevant SMPR (where applicable) as a guide.

119 ***Down-selection***

120 If the stakeholders have designated in the SMPR the need for a dispute resolution method,
121 the ERP may identify a single candidate method as dispute resolution method.

122 ***Requirements/Recommendations for Final Action***

123 After First Action adoption, the ERP may choose to make specific requirements or
124 recommendations to the Method Author. This information should be clearly delineated and
125 approved by the ERP as official recommendations and/or requirements, to be revisited upon
126 consideration for Final Action.

127 ***Candidate Method Resubmission***

128 When a Candidate Method is not adopted as First Action, the ERP shall document its concerns
129 with the methodology and/or associated validation data, the reasons for this decision, and
130 any expected remedies necessary as part of resubmission of the method. This information
131 should be clearly delineated and approved by the ERP as official recommendations and/or
132 requirements.

133 Upon subsequent review(s), the ERP should focus on whether the Method Author has
134 complied with the documented concerns from the initial ERP review. However, the ERP
135 reserves the right to raise any issue at any time that may materially impact upon method
136 fitness-for-purpose and/or ability to meet the requirements as defined in the applicable
137 SMPR(s), regardless of whether this was raised in a previous review.

138 **Publication of First Action Methods**

139 Candidate Methods are not required to be submitted for ERP review in AOAC Official Methods
140 format. However, subsequent to First Action adoption, AOAC Staff will support the Method
141 Author in ensuring proper formatting of the method for publication in *Official Methods of*
142 *Analysis*.

- 143 (1) A Candidate Method becomes First Action on the date when the ERP motion is passed.
144 (2) Methods must be drafted into AOAC format by the Method Author in collaboration with
145 AOAC staff.
146 (3) The Official Method status decision is reported concurrently with the method in
147 traditional AOAC publication venues.

148 As part of First Action publication in the *Official Methods of Analysis*, the method authors
149 must have an accepted manuscript or published paper in a reputable scientific journal,
150 preferably the *Journal of AOAC INTERNATIONAL*, containing relevant validation data.

151 **Evaluation Period**

152 Methods remain as First Action for a minimum period of two years. During this Evaluation
153 Period, the method undergoes further evaluation and validation studies. Users of First Action
154 methods are asked to provide feedback to AOAC or the Method Author on the performance
155 of the method during the Evaluation Period, to include positive and negative feedback, as well
156 as specific feedback about ruggedness. This feedback, as well as an assessment of future
157 availability of vital equipment, reference materials, and supplies should be documented in a
158 report by the Method Author for consideration by the ERP.

159 Any additional validation data obtained during this period is to be collated by the Method
160 Author and submitted to the ERP for review. Additionally, responses to requirements or
161 recommendations made by the ERP at the time of First Action adoption should be submitted
162 to the ERP for consideration.

163 For quantitative methods, data demonstrating reproducibility and uncertainty must be
164 present for Final Action consideration. Experimental designs to collect this data may vary with
165 the method protocol, available laboratories, and the intended use of the method (i.e.,
166 collaborative studies, proficiency testing, etc.). The ERP may consider other forms of
167 information in lieu of the traditional collaborative study to demonstrate method
168 reproducibility.

169 For qualitative methods, data demonstrating the probability of detection at specified
170 concentration levels as applicable must be present for Final Action consideration.
171 Experimental designs to collect this data may vary with the method protocol, available
172 laboratories, and the intended use of the method.

173 **Review of Methods for Final Action**

174 At the conclusion of the Evaluation Period, an ERP meeting is convened. Lead Reviewers will
175 report to the ERP on assigned First Action methods and should assess any additional

176 validation data or information provided during the Evaluation Period and make a
177 recommendation for deferral, repeal of First Action status, or adoption as Final Action.

178 ***Deferral***

179 If, at the end of the Evaluation Period, the feedback from method users or additional
180 validation data supplied by Method Author at the end of the Evaluation Period is deemed
181 inadequate or inconclusive, the ERP may choose to retain First Action status to allow time for
182 further information or validation data to be acquired. The ERP should discuss strategies to
183 obtain additional information to make an appropriate Final Action decision. Subsequent
184 reviews of a deferred First Action Method by the ERP must occur within two years.

185 ***Repeal***

186 At the end of the Evaluation Period, if the feedback from method users indicates that the
187 performance of a First Action method in other laboratories is unacceptable; or if no further
188 validation data is obtained, the ERP may vote to repeal the First Action status of a method.

189 ***Final Action***

190 A recommendation for a method as Final Action is forwarded to the OMB and the method
191 process undergoes a full procedural review and OMB approval.

192 ***OMB Review***

193 The OMB will review all recommendations for Final Action adoption, deferral, or repeal by
194 the ERP using applicable factors in their decision:

195 ***Procedural***

- 196 • ERP recommendations and improvements completed
- 197 • Draft Final Action method reviewed by ERP
- 198 • Reference materials used
- 199 • Verify Community validation protocols followed
- 200 • Verify SMPR criteria met
- 201 • Feedback from users of method considered
- 202 • Statistics Committee review
- 203 • Safety and Security Committee review

204 ***Documentation***

- 205 • Validation data
- 206 • Statistics Committee report
- 207 • Safety and Security Committee report
- 208 • User feedback
- 209 • External status
- 210 • ERP Report
- 211 • Impact statement from author
- 212 • Method in OMA format
- 213 • Manuscript(s) published or in press

214 The OMB may ask ERPs for further information on any potential points of concern.

215 *Publication of Final Action Methods*

216 As part of Final Action adoption, method authors must have an accepted manuscript or
217 published paper in a reputable scientific journal, preferably the *Journal of AOAC*
218 *INTERNATIONAL*, containing relevant validation data.

219 **Extensions of Scope**

220 For methods which have attained Official Method status for an analyte or analytes in one or
221 more matrices, an extension of scope may be sought such that the method would be
222 considered as an Official Method for additional matrices or additional analytes. The validation
223 required for an extension of method scope would typically be, as a minimum, the same as
224 that required by the ERP for a method to obtain First Action status. However, the ERP may
225 recommend alternative validation data to demonstrate that the extended method performs
226 in the same manner as the method under its original scope. Each ERP should develop method
227 extension guidelines to suit its needs.

228 **Method Modifications**

229 Modification to an Official Method may be editorial, minor, or major. Upon submission of a
230 method modification application, AOAC staff identify editorial modifications and process
231 changes through AOAC publications. The classification of minor and major modifications is
232 made by the ERP following a public comment period.

233 Minor changes should not be expected to affect the current validated performance nor
234 significantly affect measured results. Supporting information to justify the proposed
235 modification must accompany a request for ERP review of a minor modification to an Official
236 Method and equivalency data may be required to justify a method change.

237 Major modifications to a method will likely impact measured results or change method
238 performance. This level of modification will result in the creation of a new method, with a
239 new method number, and will follow the Official Methods Program guidance for voluntary
240 consensus standards in same manner as any other new method.

241 **Conclusion**

242 This universal pathway to Official Methods is deliberately designed to avoid creation of
243 elaborate review systems applicable only to each particular community within AOAC. The
244 intent of this universal pathway is to provide a single framework for experts in analytical
245 sciences to apply their scientific knowledge, experience, and judgment in an evidenced-based
246 manner to identify, review, and adopt the best methods currently available to meet the
247 analytical needs of each community within AOAC INTERNATIONAL.

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This guidance document was approved by the AOAC Board of Directors on MMM DD, YYYY