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Executive Summary

Acceptance Sampling by Variables (ASbV) is a statistical testing technique used in Personal Protective Equipment programs to determine the quality of the equipment in First Article and Lot Acceptance Tests. This article intends to remedy the lack of existing references that discuss the similarities between ASbV and certain techniques used in different sub-disciplines within statistics. Understanding ASbV from a statistical perspective allows us to provide DOT&E with customized test plans, beyond what is available in MIL-STD-414. We plan to submit this article to a statistics journal. This paper does not include any real test data, and does not mention any specific program by name.

Sample Size Determination Methods for Acceptance Sampling by Variables

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August 28, 2019

Abstract

The sample size for an Acceptance Sampling by Variables experiment is often determined using a standard such as MIL-STD-414 or one of its many derivatives. These standards specify sample sizes for a series of experiments to be used on a stream of incoming lots. They are not intended to be used to plan a stand-alone experiment, which requires an alternative approach. In this paper we focus on three alternatives: the operating characteristic (OC) curve approach, the Faulkenberry and Weeks approach, and a power analysis for a hypothesis test on a quantile. These three methods originate from different scientific sub-disciplines, but their concepts are quite similar and are easily confounded. To illustrate this, we review their development, highlight their similarities and differences, and present them in a consistent notation that demonstrates their mathematical equivalence.

Keywords: Acceptance Sampling by Variables, Design of Experiments, Sample Size Determination, Reliability, Operating Characteristic Curve, Tolerance Intervals, Power Analysis

1 Introduction

Determining the sample size for an Acceptance Sampling by Variables (ASbV) test can be challenging. ANSI/ASQ Z1.9-2003, a modern derivative of MIL-STD-414, is over 100 pages long and includes a “flow chart for use” that contains

26 over 50 arrows (see Figure 1 in ANSI/ASQ Z1.9-2003). Neubauer and Luko
27 (2013, 182) state, “MIL-STD-414 is complex”, while Horsnell opines, “It is my
28 view that there are probably more frustrated designers of acceptance sampling
29 schemes than in any other branch of applied statistics” (Gascoigne and Hill
30 1976, 312).

31 Wetherhill criticizes BS 6002, another MIL-STD-414 derivative, for its lack
32 of guidance involving the development of individual plans. He states, “[it] suf-
33 fers from the characteristic failing of Defence Sampling Schemes, of a lack of
34 guidance as to when the plans should be used, and of what to do if they are not
35 appropriate” (Gascoigne and Hill 1976, 308). One such example is when the
36 stream of lots is too short to provide an effective use of the switching rules. In
37 this case, a sampling plan that is independent of the overall scheme may be de-
38 sired, and Schilling (2017, 216) recommends the use of an OC curve to develop
39 this plan. BS 6002 provides the same recommendation as Schilling, and even
40 though Wetherhill states “...BS 6002 is a great improvement on MIL-STD-414”
41 he complains that “the brief reference to looking at the OC-curve [in BS 6002]
42 is totally unsatisfactory [for developing an individual plan]” (Gascoigne and Hill
43 1976, 308).

44 If one is accustomed to using a standard to develop a variables sampling
45 plan, then it can be challenging to develop an individual plan using a tradi-
46 tional OC curve approach. Textbooks that cover ASbV (e.g., Duncan [1959]
47 or Schilling [2017]) include an abundance of content on standards and their as-
48 sociated switching rules. As a result, a bit of searching is required to find the
49 theory that underpins the OC curve to develop an individual, stand-alone plan.

50 Another challenge involved with designing a stand-alone ASbV plan comes
51 from the overlapping of concepts between separate scientific sub-disciplines. It
52 is not uncommon to find nearly identical concepts nested within scientific sub-
53 disciplines that use different terminology to address the same problems (e.g.,
54 the overlap of statistical learning, machine learning, and data mining, as well
55 as the overlap of reliability analysis and survival analysis). This paper aims to
56 disentangle commonly confounded approaches for developing stand-alone ASbV
57 plans.

58 The first example of such confounding involves the methodology developed

59 both in ASbV literature and in statistical tolerance interval literature. Statis-
 60 tical tolerance intervals share a close history with ASbV. They both grew in
 61 popularity around the same time that look-up tables for the non-central t dis-
 62 tribution were becoming available (e.g., Owen [1963]). An approach for sizing
 63 tolerance intervals, called the Faulkenberry and Weeks approach (abbreviated
 64 FW; [1968]), was developed more than a decade after ASbV and is some times
 65 used instead of the OC curve approach to size an individual sampling plan. For
 66 example, Young (2016) uses the FW approach to design a sampling plan and
 67 uses historical data to inform the setting of the parameters involved in the FW
 68 computation.

69 Mixing of sub-disciplines also occurs between the OC curve approach in
 70 ASbV literature and a power analysis for a hypothesis test in statistics liter-
 71 ature. In particular, associated with the use of OC curves are the symbols α
 72 and β , which are respectively referred to as the producer and consumer risk
 73 (Montgomery 2009, 642). This implies the setup of a hypothesis test, yet this
 74 is absent in ASbV literature (for example, hypothesis tests are not included
 75 in Shilling’s textbook [2017], or Montgomery’s textbook [2009]). This raises
 76 the question: can the ASbV sample size determination problem be set up as
 77 a traditional power analysis? This concept was recently investigated for the
 78 Acceptance Sampling by Attributes problem (Samohyl 2017), but we have yet
 79 to see a similar exposition for ASbV.

80 Acceptance sampling has received less attention in literature in the last few
 81 decades. Jenson et al. (2018) showed that articles on acceptance sampling
 82 commonly appeared in the Journal of Quality Technology in the 70’s and 80’s,
 83 but in the 90’s the field shifted towards capability analysis, Stewart control
 84 charts, and classical Design of Experiments. Despite this trend, acceptance
 85 sampling remains a statistically defensible approach, and is still commonly used
 86 in defense tests.

87 In this paper we clarify the equivalence between three sample size determi-
 88 nation methodologies that can be used to plan a stand-alone ASbV test: the
 89 OC curve approach, the FW approach, and a power analysis for a hypothe-
 90 sis test on a quantile. We review the development of these methods, highlight
 91 their similarities and differences, and present them in a consistent notation that

102 demonstrates their mathematical equivalence.

103 Assumptions throughout this paper follow the classic setup of the Acceptance
 104 Sampling by Variables problem. Let X denote the response variable, where
 105 $X \sim \mathcal{N}(\mu, \sigma^2)$ and both μ and σ^2 are unknown. Let X_1, X_2, \dots, X_n be a sample
 106 from $\mathcal{N}(\mu, \sigma^2)$. The sample mean \bar{x} and variance s^2 are

$$\bar{x} = \frac{1}{n} \sum_{i=1}^n X_i \quad , \quad s^2 = \frac{1}{n-1} \sum_{i=1}^n (X_i - \bar{x})^2 \quad . \quad (1)$$

107 Let Z_p denote the p quantile of a standard normal distribution, where the p
 108 quantile of $\mathcal{N}(\mu, \sigma^2)$ is given by

$$Q_p = \mu + Z_p \sigma \quad . \quad (2)$$

109 Acceptance criteria for variables sampling plans can be based on an upper,
 100 lower, or double specification limit, but for the sake of simplicity, in this paper
 101 we focus solely on the upper specification limit, U .

102 2 Operating Characteristic Curve

103 Jennet and Welch (1939), Johnson and Welch (1940), the Statistical Research
 104 Group (abbreviated as SRG; see Eisenhardt et al. [1947]), Bowker and Goode
 105 (1952), and Liebermann and Resnikoff (1955) are often credited as the architects
 106 of ASbV. The following review primarily comes from *Techniques of Statistical*
 107 *Analysis* prepared by Eisenhardt et al. (1947) that reflects the work of the SRG.

108 The SRG states, “An ASbV procedure may be analyzed into three phases”
 109 (1947, 13). The first phase is “the *plan of action*, that is, the set of rules on
 110 the basis of which to accept or reject the lot.” The plan of action that we
 111 focus on in this paper for assessing an upper specification limit, U , is called the
 112 “Standard Deviation Method - Variability Unknown” by MIL-STD-414, and is
 113 also sometimes referred to as the “k-method” (Schilling and Neubauer 2017).
 114 The Standard Deviation Method acceptance criteria is

$$\frac{U - \bar{x}}{s} \geq k \quad , \quad (3)$$

115 where k is referred to as an “acceptability constant (e.g. MIL-STD-414)” that
 116 must be determined for a given plan.

117 The SRG notes that other measures of central tendency could be used instead
 118 of \bar{x} and s in Equation 3, such as the median or range. Measures such as these
 119 have found their way into MIL-STD-414 and subsequent derivative standards
 120 because they were thought to have been easier to implement (i.e., the range
 121 is easier to calculate than the standard deviation), but the SRG notes that
 122 “the gains in computational simplicity that may be afforded by other measures
 123 are likely to be unimportant in the situation for which variables inspection is
 124 appropriate.” For an additional discussion on the controversy involving the
 125 inclusion of “other” measures, see Acheson and Duncan (1975, 40).

126 The SRG defines the next two phases as follows. The second phase is “the
 127 *amount of inspection* required by the plan, that is, the number of items that
 128 must be inspected from each lot,” and the third and final phase is “the *operating*
 129 *characteristics* of the plan, that is, the proportion of submitted lots of various
 130 qualities that will be accepted and rejected if the plan is used.”

131 The amount of inspection and operating characteristics can be investigated
 132 using an OC curve. SRG states, “it is important to know what proportion of
 133 submitted lots will be accepted for each possible quality...as a plan is clearly
 134 unsuitable if it passes too many of the lots of unsatisfactory quality or rejects
 135 too many of the lots of acceptable quality that are submitted to it.” For a given
 136 sample size, an OC curve displays the probability of lot acceptance, denoted as
 137 P_A , versus the assumed, or anticipated, lot proportion defective, denoted as \tilde{p} ,
 138 where \tilde{p} is related to p (see Equation 2) by $\tilde{p} = 1 - p$.

139 OC curves were challenging to produce in the middle of the twentieth century
 140 because computing the probability of acceptance involved the evaluation of the
 141 non-central t distribution. In the paper that is credited with some of the first
 142 theory on ASbV (Jennett and Welch 1939), the authors state, “[the distribution
 143 of k] is a particular example of what is termed the non-central t distribution.
 144 Tables of this distribution, in a form suitable for the present problem, do not
 145 exist, but are in process of calculation.” This task was accomplished in a follow-
 146 up paper (Johnson and Welch 1940).

147 The derivation of the distribution of k in Equation 3 leads to the equation
 148 for the OC curve. Jennet and Welch (1939) note, “The true distribution of $[k]$
 149 is not difficult to find,” while Johnson and Welch (1940) and Liebermann and

150 Resnikoff (1955) seemed to have shared the same sentiment, as they showed no
 151 intermediate steps in re-expressing Equation 3 as

$$\left(\frac{\sqrt{n}(U - \mu)}{\sigma} - \frac{\sqrt{n}(\bar{x} - \mu)}{\sigma} \right) \frac{\sigma}{s} \geq \sqrt{nk} \quad . \quad (4)$$

152 For more details on obtaining Equation 4, one can consult Resnikoff and Lieber-
 153 man (1957) or Schilling and Neubauer (2017, 238).

154 In Equation 4, by definition of the non-central t distribution, the quantity
 155 on the left hand side of the equation is distributed as a non-central t random
 156 variable, with $n - 1$ degrees of freedom and non-centrality parameter equal to

$$\sqrt{n} \frac{U - \mu}{\sigma} \quad . \quad (5)$$

157 Thus, the probability of acceptance is

$$P_A = Pr \left(\frac{U - \bar{x}}{s} \geq k \right) = 1 - t \left(\sqrt{nk}, n - 1, \sqrt{n} \frac{U - \mu}{\sigma} \right) \quad , \quad (6)$$

158 where $t(\cdot, \cdot, \cdot)$ is the noncentral t cumulative distribution function with quantile
 159 \sqrt{nk} , $n - 1$ degrees of freedom, and noncentrality parameter $\sqrt{n} \frac{U - \mu}{\sigma}$. Equation
 160 6 is not yet the equation of the OC curve, since it requires one more assumption.

161 SRG stresses that the OC curve cannot predict the future, because the true
 162 population parameters are unknown. They state, “...the OC curve does *not*
 163 show the probability that an accepted lot will be of quality p ” (1947, 16). The
 164 OC curve simply shows, for a given sample size, the probability of accepting
 165 a lot of an assumed (or anticipated) quality. This means that the OC curve
 166 assumes that the fraction of the normal population exceeding U is equal to \bar{p} ,
 167 or equivalently

$$\frac{U - \mu}{\sigma} = Z_{1-\bar{p}} \quad . \quad (7)$$

168 Finally, substituting this assumption into Equation 6 gives the equation of the
 169 OC curve as

$$P_A = 1 - t \left(\sqrt{nk}, n - 1, \sqrt{n} Z_{1-\bar{p}} \right) \quad . \quad (8)$$

170 The equation of the OC curve can be uniquely defined in a number of differ-
 171 ent ways. One way is to specify two points that the OC curve passes through.

Let these two points be $(\tilde{p} = 1 - p_1, P_A = 1 - \alpha)$ and $(\tilde{p} = 1 - p_2, P_A = \beta)$.
 Substituting these points, respectively, into Equation 8 yields

$$1 - \alpha = 1 - t(\sqrt{n}k, n - 1, \sqrt{n}Z_{p_1}) \quad , \quad (9)$$

$$\beta = 1 - t(\sqrt{n}k, n - 1, \sqrt{n}Z_{p_2}) \quad . \quad (10)$$

which can be respectively expressed as

$$\sqrt{n}k = t'(\alpha, n - 1, \sqrt{n}Z_{p_1}) \quad , \quad (11)$$

$$\sqrt{n}k = t'(1 - \beta, n - 1, \sqrt{n}Z_{p_2}) \quad . \quad (12)$$

Here, $t'(\cdot, \cdot, \cdot)$ denotes the non-central t quantile function, where the first, second, and third arguments to this function are the cumulative density, degrees of freedom, and non-centrality parameter.

Theoretically, the sample size that causes the OC curve to pass through the two points can be found as the solution for n that satisfies

$$t'(\alpha, n - 1, \sqrt{n}Z_{p_1}) = t'(1 - \beta, n - 1, \sqrt{n}Z_{p_2}) \quad , \quad (13)$$

but in nearly all cases the discrete nature of n prevents the OC curve from passing through those exact two points. Instead, one can show that the OC curve passes through the two points $(\tilde{p} = 1 - p_1, P_A = 1 - \alpha^*)$ and $(\tilde{p} = 1 - p_2, P_A = \beta)$, where α^* is the actual value of α that the OC curve passes through. That is, the OC curve passes through three of the four coordinates of the intended two points, but the fourth coordinate, α , has a source of error that is introduced due to the discrete nature of n . Additionally, to *control the risk* associated with the parameter α , we constrain the solution for n such that $\alpha^* < \alpha$. That is, the sample size solution is the minimum value of n that satisfies

$$t'(\alpha, n - 1, \sqrt{n}Z_{p_1}) \geq t'(1 - \beta, n - 1, \sqrt{n}Z_{p_2}) \quad . \quad (14)$$

After numerically solving for n , we solve for k by substituting n and the coordinates of one of the points that the OC curve passes through into Equation

191 8. Then, using n and k , the OC curve displays the probability of acceptance,
 192 P_A , versus the anticipated fraction defective, \tilde{p} .

193 Before we move on we should note that ASbV plans typically assign names
 194 to the parameters of the OC curve. According to Montgomery (2009), α is
 195 the “producer’s risk”, $1 - p_1$ is the Acceptable Quality Limit (AQL), β is the
 196 “consumer’s risk”, and $1 - p_2$ is the Rejectable Quality Limit (RQL). That is,

$$P_A(\tilde{p} = 1 - p_1 = AQL) = 1 - \alpha \quad , \quad (15)$$

197 and

$$P_A(\tilde{p} = 1 - p_2 = RQL) = \beta \quad . \quad (16)$$

198 In terms of these variables, the OC curve takes the usual form (e.g., Juran and
 199 Godfrey [1999, 46.46]) as shown in Figure 1.

200 3 OC Curve Notional Example

201 The Army buys body armor from a supplier. The Army has established an upper
 202 specification on a bullet’s penetration depth into the body armor that is equal
 203 to 5 mm. If 1 percent or more of the bullets fired demonstrate a penetration
 204 depth above this limit, the Army wishes to accept the lot with probability 0.95
 205 ($p_1 = 1 - AQL = 0.99, 1 - \alpha = .95$), whereas if 6 percent or more of the bullets
 206 fired demonstrate a penetration depth above this limit, the Army would like to
 207 reject the lot with probability .90 ($p_2 = 1 - RQL = .94, \beta = .10$). Determine
 208 the sample size for this variables sampling plan.

209 Begin by finding the minimum value of n that satisfies Equation 14. A
 210 brute-force way of doing this is to simply plot the quantity on the left and right
 211 hand side of Equation 14 as a function of n , as shown in Figure 1, and visually
 212 locate the minimum value of n that satisfies 14. It is clear from this figure that
 213 the solution is $n = 42$.

214 Then, calculate k using Equation 10 with $\beta = 0.1$ and $p_2 = .94$ to obtain
 215 $k = 1.898$. Substituting this value of k , $n = 42$, and $p_1 = .99$ into Equation 9,
 216 we obtain $\alpha^* = 0.047$, which means we are properly *controlling the risk*, since

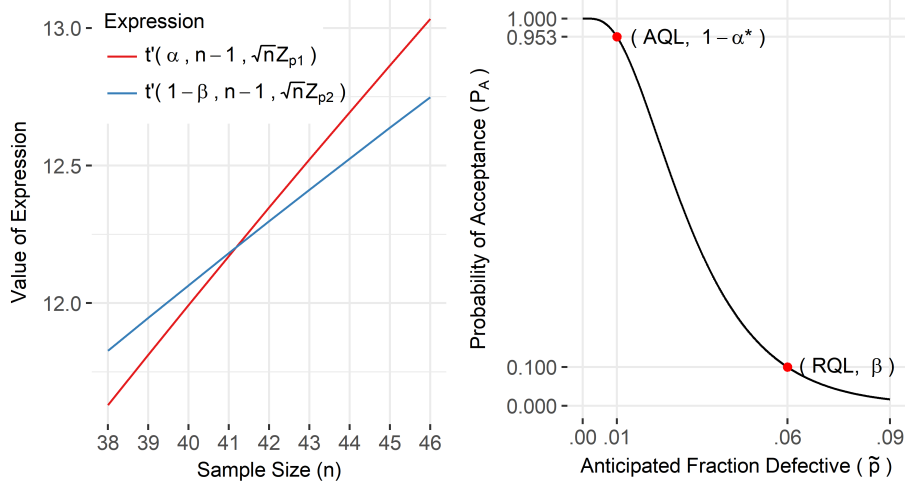


Figure 1: Left side of figure demonstrates the numerical solution of the body armor example. Right side of figure shows the OC curve.

217 $\alpha^* < \alpha$. The OC curve can be plotted using Equation 8 as shown on the right
 218 side of Figure 1.

219 4 Faulkenberry and Weeks Approach

220 The Faulkenberry and Weeks approach, a sample size determination methodol-
 221 ogy quite similar to the OC curve approach, comes from the field of statistical
 222 tolerance regions. The following review is taken from Krishnamoorthy (2009).
 223 A tolerance interval is constructed using the random sample X_1, X_2, \dots, X_n , and
 224 is required to contain a proportion p or more of the sampled population, with
 225 confidence level $1 - \Gamma$. Formally, a $(p, 1 - \Gamma)$ upper tolerance interval has the
 226 form $\bar{x} + ks$, where k is to be determined such that it satisfies the condition

$$Pr(\bar{x} + ks \geq \mu + Z_p \sigma) = 1 - \Gamma \quad . \quad (17)$$

227 Krishnamoorthy shows the derivation for the solution of k using a “classical
 228 approach” and a “generalized variable approach” that both result in

$$k = \frac{1}{\sqrt{n}} t'(1 - \Gamma, n - 1, \sqrt{n} Z_p) \quad . \quad (18)$$

229 Faulkenberry and Weeks (1968) developed a procedure for determining the
 230 sample size that produces a tolerance interval that meets a “goodness criterion.”
 231 We summarize their approach as follows.

232 Consider two different tolerance intervals that are constructed from the same
 233 random sample, X_1, X_2, \dots, X_n . For the first tolerance interval, let $p = p_1$ and
 234 $\Gamma = \alpha$, and for the second tolerance interval, let $p = p_2$ and $\Gamma = 1 - \beta$. That is,
 235 consider a (p_1, α) tolerance interval, and a $(p_2, 1 - \beta)$ tolerance interval, where
 236 $p_1 > p_2$, and α and β are typically values between .01 – .20, and p_1 and p_2 are
 237 typically values between .80 – .99.

238 Faulkenberry and Daly (1970) define the “goodness criterion” in a compact
 239 form, stating, “The criterion used for determining sample size is as follows:
 240 For a tolerance limit such that $Pr(\text{coverage} \geq P) = \gamma$, choose $P' > P$ and
 241 δ (small) and require $Pr(\text{coverage} \geq P') \leq \delta$.” For the sake of comparison
 242 between sample size determination methods in this paper, we use a different set
 243 of variables than FW. That is, let $\alpha = \delta$, $\beta = 1 - \gamma$, $p_1 = P'$, and $p_2 = P$.

244 Thus, for the $(p_2, 1 - \beta)$ tolerance interval that can be expressed as

$$t(\sqrt{n}k, n - 1, \sqrt{n}Z_{p_2}) = 1 - \beta \quad , \quad (19)$$

245 according to the “goodness criterion”, the sample size solution is the minimum
 246 value of n that satisfies

$$t(\sqrt{n}k, n - 1, \sqrt{n}Z_{p_1}) \leq \alpha \quad . \quad (20)$$

247 Or equivalently, the FW sample size solution is the minimum value of n that
 248 satisfies

$$t'(\alpha, n - 1, \sqrt{n}Z_{p_1}) \geq t'(1 - \beta, n - 1, \sqrt{n}Z_{p_2}) \quad . \quad (21)$$

249 Equation 21 is identical to the sample size solution using the OC curve
 250 approach (same as Equation 14).

251 4.1 FW Example

252 The following brief example is taken from Faulkenberry and Daly, (1970, 818).
 253 Suppose we are interested in $p_1 = .95$, $\alpha = .10$, $p_2 = .90$, and $\beta = .10$. Using

Equation 21, and solving for n in a similar manner as the OC curve example, we obtain $n = 104$, which is used to obtain $k = 1.466$. Substituting these values and $p_1 = .95$ into Equation 9, we obtain $\alpha = .099$. Thus, the goodness inequality, $Pr(\text{coverage} \geq p_1) \leq \alpha$, is satisfied.

5 Hypothesis Test on Quantile Approach

The last approach in this paper is a power analysis for the hypothesis that tests whether a population quantile is different from a constant. In developing the power function that we use to determine sample size, we follow a procedure similar to that used in the classical examples, such as the test involving $H_0 : \mu = \mu_0$, $H_a : \mu \neq \mu_0$ (e.g. Mathews [2010, 31]). Lenth (2001) outlines this general procedure as follows.

The procedure starts with the definition of the null and alternative hypotheses, and the definition of the test statistic (or as Lenth states, “the underlying probability model for the data”). This is followed by the definition of the effect size (what Casella and Berger [2002, 382] call “defining the rejection region”), the solution for the power function, and finally the sample size that provides a desired level of power.

We implement this general procedure for the hypothesis that tests whether a population quantile is different from a constant. To begin, define the null and alternative hypothesis as

$$H_0 : Q_{p_1} = U \implies \text{Accept Lot} \quad , \quad (22)$$

$$H_a : Q_{p_1} > U \implies \text{Reject Lot} \quad . \quad (23)$$

Here, Q_{p_1} denotes the p_1 quantile, and U , the upper specification limit, is treated as a constant. The null hypothesis implies the lot is accepted, while the alternative implies the lot is rejected.

The test statistic is conveniently obtained from the equation for the confidence interval on Q_{p_1} . Chakraborti and Li (2007) present a derivation of this equation (originally shown by Lawless [2002]) based on the biased estimator $\hat{Q}_{p_1} = \bar{x} + Z_{p_1} \hat{\sigma}$, where $\hat{\sigma}$ is the biased MLE of σ . It is straightforward to

perform a similar derivation, based on a slightly different estimator that is also biased, $\hat{Q}_{p_1} = \bar{x} + Z_{p_1}s$, to obtain an equation for a confidence interval on Q_{p_1} that satisfies

$$Pr\left(\bar{x} + t'(1 - \alpha, n - 1, \sqrt{n}Z_{p_1}) \frac{s}{\sqrt{n}} \geq Q_{p_1}\right) = 1 - \alpha \quad . \quad (24)$$

This equation also satisfies the definition of a $(p_1, 1 - \alpha)$ upper one-sided tolerance limit, as it is well known that a one-sided upper tolerance limit is equivalent to a one-sided upper confidence interval on a quantile (e.g. Krishnamoorthy [2009, 27]). We can rearrange Equation 24 to obtain

$$Pr\left(\frac{Q_{p_1} - \bar{x}}{s/\sqrt{n}} \leq t'(1 - \alpha, n - 1, \sqrt{n}Z_{p_1})\right) = 1 - \alpha \quad . \quad (25)$$

which implies that the pivotal quantity,

$$\frac{Q_{p_1} - \bar{x}}{s/\sqrt{n}} \quad , \quad (26)$$

is distributed as a non-central t random variable with $n - 1$ degrees of freedom and non-centrality parameter equal to $\sqrt{n}Z_{p_1}$.

Under H_0 we assume $Q_{p_1} = U$ and apply this assumption to Equation 26 to obtain the test statistic

$$T = \frac{U - \bar{x}}{s/\sqrt{n}} \quad . \quad (27)$$

In practice, once data collection is complete, we reject H_0 if $T < T_{crit}$, where the critical value is

$$T_{crit} = t'(\alpha, n - 1, \sqrt{n}Z_{p_1}) \quad . \quad (28)$$

Prior to conducting the experiment, and for the purpose of determining sample size, we need to assume a value for U assuming that we reject H_0 . That is, we need to define the effect size. Thus, when H_a is true, let $U = Q_{p_2}$. The difference between Q_{p_1} and Q_{p_2} can be interpreted as the effect size.

If in Equation 27 we substitute U with Q_{p_2} , then the test statistic assuming H_a is true is distributed as a non-central t random variable with $n - 1$ degrees of freedom and non-centrality parameter equal to $\sqrt{n}Z_{p_2}$.

Power is the probability that the test statistic (assuming H_a is true) is less than the critical value, and represents the probability of correctly rejecting the lot. The power function is

$$1 - \beta = t(T_{crit}, n - 1, \sqrt{n}Z_{p_2}) \quad . \quad (29)$$

Additionally, if we *control the risk* associated with α as we did in the previous sections, then we can constrain the sample size solution such that the integer solution for n yields an actual value, α^* , that is less than the intended value, α . This implies that the integer solution for n satisfies

$$1 - \beta \leq t(T_{crit}, n - 1, \sqrt{n}Z_{p_2}) \quad . \quad (30)$$

Applying the inverse t quantile function with $n - 1$ degrees of freedom and noncentrality parameter $\sqrt{n}Z_{p_2}$ to both sides of Equation 30, results in

$$t'(1 - \beta, n - 1, \sqrt{n}Z_{p_2}) \leq T_{crit} \quad , \quad (31)$$

which is identical to the sample size equations (14 and 21) from the previous sections.

6 Power Analysis Example

The Air Force buys guided missiles from a manufacturer. The Air Force has established an upper specification on the missile's radial miss distance that is equal to 5 feet. If 4 percent or more of the missiles fired demonstrate a radial miss distance above this limit, the Air Force wishes to accept the lot with probability 0.95 ($p_1 = 1 - AQL = 0.96, 1 - \alpha = .95$), whereas if 12 percent or more of the missiles fired demonstrate a radial miss distance above this limit, the Air Force would like to reject the lot with probability .90 ($p_2 = 1 - RQL = .88, \beta = .10$). Determine the sample size for this variables sampling plan.

As in the previous sections, the solution for n can be found numerically, which is $n = 53$. This results in $\alpha^* = .0499$, which satisfies $\alpha < \alpha^*$. It is instructive to plot the density of the test statistic assuming H_0 is true, and assuming H_a is true, as shown in Figure 2.

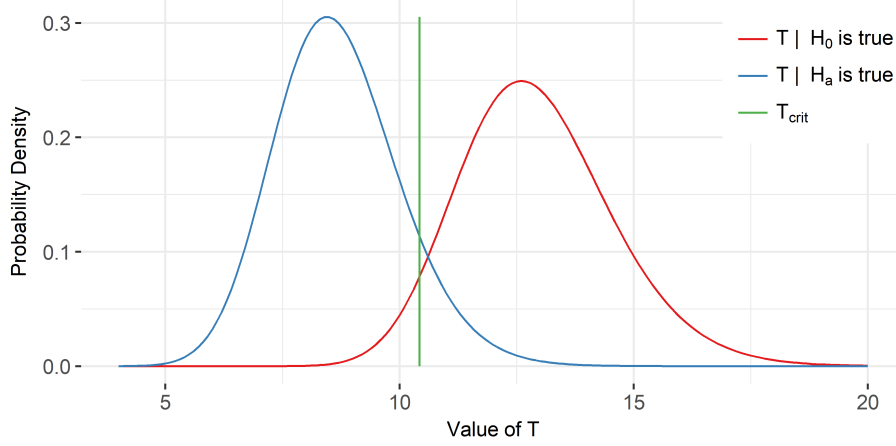


Figure 2: The distribution of the test statistic under H_0 and H_a for $n=53$.

Power, denoted as $1 - \beta$, is the probability of correctly rejecting H_0 , or equivalently the probability of correctly rejecting the lot. Power is the area under the blue curve to the left of T_{crit} . The complement of power is the Type II error, denoted as β , which represents the consumer risk, because it conveys the risk of accepting a bad lot.

Confidence level, denoted as $1 - \alpha$, is the probability of correctly accepting H_0 , or equivalently the probability of correctly accepting the lot. Confidence level is the area under the red curve to the right of T_{crit} . The complement of confidence level is the Type I error, denoted as α , which represents the producer's risk, because it conveys the risk of rejecting an acceptable lot.

7 Discussion

We highlighted three methods for sizing a stand-alone ASbV experiment, but other approaches can be found in literature. For instance, Lieberman and Resnikoff's (1955) landmark paper bases an acceptance criteria on a minimum variance unbiased estimator (MVUE). The power function for the MVUE in this case is quite complex and does not have the simple non-central t format that we saw from the other three methods in this paper, which may be why it is less commonly used in practice.

Similarly, Hamilton (1995) also bases the acceptance criteria on an estimator

of the fraction defective, but this time the estimator is biased. Their sample size determination equation is similar in form to those used by the other three approaches in this paper, but Hamilton relies on OC curves as opposed to a hypothesis test and power function to formulate the sample size determination problem.

Alternative Bayesian approaches are also available in literature. Average coverage criteria, average length criterion, utility theory, or Bayes factors (see Adcock [1997]), could potentially be adapted to the ASbV problem. One method in particular (Easterling and Weeks 1970) adapted the FW approach for use in a Bayesian setting. Surely, other sample size determination methodologies exist that would also be appropriate for the ASbV problem.

We compared the OC curve approach, the FW approach, and the power analysis for a test on a quantile because of their prevalence of use for sizing ASbV tests. In this paper we believe we helped clarify how nearly identical these methods really are by reviewing their origins and by demonstrating the equivalence of their sample size determination equations (Equations 14, 21, and 30).

To demonstrate this equivalence, we had to make a couple of minor assumptions involving the inequality sign in the sample size equations. For the FW approach, the inequality sign organically occurs due to the definition of the “goodness criterion.” For the OC curve approach, theoretically, the inequality sign should be an equals sign because the OC curve approach is typically defined by specifying two points that the curve passes through. Given that the integer solution for n prevents the curve from exactly passing through two points, it seemed reasonable that we impose the arbitrary constraint, $\alpha^* < \alpha$, for the sake of matching the OC curve sample size equation to the FW approach. For the power analysis, we used the same line of reasoning and imposed the same arbitrary constraint, $\alpha^* < \alpha$. We should note that Lenth’s guideline for power analyses suggests that the sample size solution be the minimum value of n that provides a target value of power that is greater than the intended value of power. Consequently, this would flip the inequality sign. However, this is quite trivial in practice, as the direction of the inequality can only change the sample size result by one.

378 The FW approach and the OC curve approach use a nearly identical deriva-
 379 tion that follows the “classical approach” (e.g., Krishnamoorthy [2009, 26]).
 380 That is, they start with the equation of a tolerance interval or the equation of
 381 the acceptance criteria, and re-express it in terms of the non-central t distribu-
 382 tion. The FW approach and OC curve approach even share the same symbol, k .
 383 For tolerance intervals, k is the “k-factor”, while in ASbV it is an “acceptability
 384 constant,” but they have the same mathematical interpretation. In contrast,
 385 the derivation of the test statistic in the power analysis (e.g. Chakraborti and
 386 Li [2007]) more closely follows the “generalized variable approach” (e.g. Krish-
 387 namoorthy [2009, 26]) and does not use k or any equivalent symbol.

388 Complex methodologies can be presented in simpler terms to gain traction
 389 within a community of practitioners, but it obscures the underlying theory.
 390 The architects of ASbV worked in an applied setting, and it seems plausible
 391 that they conceived ASbV with statistical terminology and language in mind
 392 (e.g., Neyman Pearson hypothesis testing), but then packaged ASbV for their
 393 engineering audience by defining names, such as “consumer risk,” and omitting
 394 statistical terms. It is challenging for a practitioner to customize a stand-alone
 395 ASbV plan when the underlying theory is obscured.

396 Framing the ASbV problem in terms of a power analysis can assist in creating
 397 customized test plans. For instance, the hypothesis test in this paper easily could
 398 have been changed to a regression model problem that focused on a hypothesis
 399 test on a conditional quantile corresponding to a particular point in a factorial
 400 experiment. Extensions like this could add to the body of ASbV theory, but we
 401 save such work for the future.

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