

## Chapitre 5: les tests d'équivalence

Lorsqu'on applique un test d'hypothèse, l'hypothèse nulle la plus couramment définie est celle d'absence d'effet ou de différence entre les groupes (???). Il arrive également parfois que les chercheurs définissent un intervalle de valeur comme hypothèse nulle, mais le plus souvent, cet intervalle est borné par la valeur 0 (???), on parle alors d'hypothèse unilatérale. Avec cette stratégie, le rejet de l'hypothèse nulle constitue un soutien en faveur de la présence d'un effet non nul. Par contre, le non rejet de l'hypothèse nulle ne peut être interprété comme un soutien en faveur de l'absence d'effet, or il arrive souvent que des chercheurs l'interprètent de la sorte. (???), par exemple, ont reporté que parmi 150 articles publiés entre 1940 et 1999 dans le *JAP (Journal of Applied Psychology)*, 38% interprétaient un résultat non significatif comme une acceptation de l'hypothèse nulle. Plus récemment, (???) a noté que l'expression "pas d'effet" a été utilisée dans 108 articles publiés dans *Social Psychological and Personality Science* avant août 2016 et que dans presque tous les cas, c'était sur base du non rejet de l'hypothèse nulle que cette conclusion était tirée.

A travers ce chapitre, notre premier objectif sera d'expliquer pourquoi interpréter le non rejet de l'hypothèse d'absence d'effet comme un soutien en faveur d'une absence d'effet n'est pas une bonne stratégie. Nous introduirons ensuite les tests d'équivalence qui permettent d'obtenir un soutien en faveur d'un effet jugé non pertinent, et plus particulièrement le TOST (Two One-sided test). Pour finir, nous présenterons un article dans lequel nous comparons le TOST à la *SGPV* (Second Generation P-value), une stratégie récemment développée par (???).

### Limites de l'approche traditionnelle

Lorsqu'on teste une hypothèse nulle, il y a deux conclusions possibles: soit on la rejette soit on ne la rejette pas. Si rejeter l'hypothèse nulle amène à conclure en faveur de l'hypothèse alternative, ne pas la rejeter ne permet pas de conclure en faveur de l'hypothèse nulle. Au

mieux, cela nous montre que les données ne sont pas incompatibles avec l'hypothèse nulle, mais cela ne veut en aucun cas dire qu'elles ne sont compatibles avec aucune autre hypothèse. Afin de l'illustrer, la Table 1 résume les résultats de simulations Monte Carlo pour un ensemble de 70 scénarios qui varient en fonction de la taille des échantillons ( $n_j$ ) et de la différence entre les moyennes des deux populations dont sont extraits les échantillons ( $\mu_1 - \mu_2$ ). Pour chaque scénario, à 5000 reprises, nous avons généré une paire d'échantillons, réalisé un test  $t$  de Student pour échantillons indépendants et extrait la  $p$ -valeur du test. Ensuite, nous avons calculé la proportion de  $p$ -valeurs supérieures à .05 (le risque alpha communément accepté, ???), nous amenant à ne pas rejeter l'hypothèse nulle. Cette proportion correspond au taux d'erreur de type II (communément appelé  $\beta$ ).

Table 1.

*Proportion de  $p$ -valeurs supérieures à .05 ( $\beta$ ) en fonction de la taille des échantillons ( $n_j$ ) et de la différence entre les moyennes de chaque population ( $\mu_1 - \mu_2$ ).*

$n_j$	Différence de moyennes dans la population ( $\mu_1 - \mu_2$ )									
	.1	.3	.5	.7	.9	1.1	1.3	1.5	1.7	1.9
<b>10</b>	.943	.903	.818	.691	.513	.358	.206	.119	.058	.020
<b>15</b>	.940	.881	.732	.542	.348	.169	.066	.023	.005	.001
<b>20</b>	.939	.856	.660	.423	.208	.076	.020	.005	< .001	< .001
<b>25</b>	.932	.820	.587	.320	.119	.034	.006	< .001	< .001	< .001
<b>50</b>	.925	.667	.297	.061	.006	< .001	< .001	< .001	< .001	< .001
<b>100</b>	.890	.437	.063	.002	< .001	< .001	< .001	< .001	< .001	< .001
<b>200</b>	.828	.150	.001	< .001	< .001	< .001	< .001	< .001	< .001	< .001

*Note.* Pour chaque scénario, les deux échantillons sont toujours de même taille ( $n_j = n_1 = n_2$ ) et sont extraits de populations se distribuant normalement et ayant la même variance ( $\sigma_1 = \sigma_2 = \sigma$ ). La moyenne de la première population ( $\mu_1$ ) vaut systématiquement 0, et celle de la deuxième population ( $\mu_2$ ) varie de sorte à obtenir la différence de moyenne  $\mu_1 - \mu_2$  désirée. Par ailleurs,  $\sigma$  vaut systématiquement 1, si bien que la différence de moyenne brute est égale au  $\delta$  de Cohen.

Dans la mesure où une vraie différence entre les moyennes de population existe pour l'ensemble des scénarios envisagés, nous espérons être en mesure de rejeter l'hypothèse nulle le plus souvent possible. Pourtant, pour plusieurs scénarios, le nombre d'itérations amenant à conclure au non rejet de l'hypothèse nul est bien supérieur au nombre d'itérations amenant à conclure au rejet de l'hypothèse nulle, comme on peut le voir à travers les valeurs

$\beta$  très élevées dans certains cas. Par exemple, avec 10 sujets par groupes et considérant  $\sigma_1 = \sigma_2 = 1$ , on ne détectera pas une différence de moyenne de .1 dans près de 95% des cas. Avec 200 sujets par groupe, on ne détectera toujours pas cette différence dans 80% des cas. En présence d'un effet non nul, cela se justifie par un manque de puissance des tests réalisés, ce qui démontre bien qu'un non rejet de l'hypothèse nulle peut en fait signifier deux choses: soit qu'il n'y a vraiment pas de différence entre les moyennes des populations (ou autrement dit, que les différences observées sont dûes au hasard), soit que le test n'est pas suffisamment puissant pour détecter la différence. Or, le manque de puissance des tests est récurrent dans la littérature, comme tendent à le montrer diverses méta-analyses (voir par exemple ???; ???; ???).

Pour éviter d'interpréter un test peu puissant comme un soutien en faveur de l'hypothèse nulle, dans les années 80, l'approche de la puissance est devenue l'approche par défaut pour tester l'équivalence (???). A travers cette approche qui est restée très populaire (???), dans un premier temps, on définit ce qu'on considère comme étant la plus petite valeur d'intérêt (en anglais, le "SESOI" pour "Smaller Effect Size of Interest"), c'est-à-dire la taille d'effet minimale requise pour considérer qu'un effet est pertinent. Ensuite, on estime la puissance de notre test à détecter un effet de cette taille<sup>1</sup>, et si cette estimation atteint une valeur jugée satisfaisante (en général, 80%), alors on considère que l'on peut interpréter le non rejet de l'hypothèse nulle d'absence d'effet comme soutien en faveur de l'équivalence (???; ???; ???). L'idée sous-jacente est que si l'effet est au moins aussi grand que les limites de la zone d'équivalence, on devrait rejeter l'hypothèse nulle dans un pourcentage raisonnable de cas. Par conséquent, un non rejet de l'hypothèse nulle devrait vraisemblablement signifier que l'effet n'atteint pas le SESOI et donc, que l'effet observé n'est pas pertinent.

Bien que ce raisonnement puisse sembler tentant, de prime abord, il présente d'importantes

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<sup>1</sup> On parle d'estimation et non de mesure, car la puissance dépend de paramètres de population qu'on ne connaît pas (Schuirmann,1987).

limites. Via cette approche, la probabilité de détecter une absence d'effet va augmenter non seulement quand la taille des échantillons va diminuer (à condition de ne pas passer en deçà de la puissance minimale requise), mais également quand l'erreur (la variabilité des scores au sein de chaque groupe) va augmenter (???; ???). Ce dernier point est illustré au sein de la Figure 1. A travers cette figure, on représente les situations dans lesquelles l'approche de la puissance nous amènera à conclure à l'équivalence, lorsqu'on travaille avec des échantillons de taille 50, en acceptant un risque  $\alpha$  de 5% et en exigeant une puissance minimale de 80% pour détecter une différence de 20 unités ( $|\theta_j| = 20$  ( $j = 1, 2$ )). Sur l'axe des abscisses, on représente les estimations de la différence de moyenne ( $\bar{X}_1 - \bar{X}_2$ ) et sur l'axe des ordonnées, la précision des estimations ( $S\sqrt{\frac{2}{n}}$  correspond à l'estimation l'erreur standard de  $\bar{X}_1 - \bar{X}_2$ , avec  $S$  étant l'écart-type poolé et  $n$  étant la taille de chaque échantillon, lorsque les échantillons ont tous les deux la même taille et sont extraits de population ayant la même variance)<sup>2</sup>. La zone grisée représente l'ensemble des combinaisons estimation/précision qui vont amener à conclure à l'équivalence. On constate que plus notre estimation de  $\bar{X}_1 - \bar{X}_2$  est précise (lorsqu'on se déplace du haut vers le bas, sur l'axe des ordonnées), plus l'estimation doit être proche de 0 pour pouvoir conclure à l'équivalence. Ce n'est bien entendu pas une propriété désirable.

## Les tests d'équivalence

Avec les tests d'équivalence, il n'est pas possible de démontrer qu'un effet vaille exactement zéro (???). Il est par contre possible de montrer que l'effet observé est suffisamment petit pour être jugé non pertinent. Le point de départ des tests d'équivalence est de définir  $\theta_1$  et  $\theta_2$ , les limites inférieures et supérieures de la zone d'équivalence, cette dernière contenant

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<sup>2</sup> C'est pas pure facilité que l'on envisage le cas où les échantillons sont de même taille et que l'on suppose que la condition d'homogénéité des variances est respectée. Notons cependant que le raisonnement peut être généralisé aux scénarios où les deux échantillons n'ont pas la même taille et sont extraits de population n'ayant pas la même variance, d'après Schuirman (2011).

l'ensemble des valeurs jugées trop petites pour être susceptibles de nous intéresser. Ces limites peuvent être exprimées soit dans l'unité des données brutes, soit en terme standardisé.

Il existe ensuite plusieurs approches pour démontrer que l'effet observé se situe ans la zone d'équivalence (*voir Meyners, par exemple*). Parmi celles-ci, une approche très simple est celle du *TOST* (pour “Two one-sided tests”, (???) ; (???)<sup>3</sup>). Le principe est de définir deux hypothèses nulles: la première est que l'effet observé est inférieur à la limite inférieure de la zone d'équivalence:

$$H0_1 : \theta < \theta_1$$

La deuxième est que l'effet observé est supérieur à la limite supérieure de la zone d'équivalence:

$$H0_2 : \theta > \theta_2$$

Lorsque les deux hypothèses nulle peuvent être simultanément rejetées, on peut conclure à l'équivalence (Seaman & Serlin, 1998, cités par lakens\_\_equivalence\_2017). Il est à noter qu'il n'est pas nécessaire de réaliser deux tests pour rejeter les deux hypothèse nulles. En effet, si l'effet observé se situe à équidistance des deux limites de la zone d'équivalence, les tests contre chacune des limites mèneront à une conclusion identique, car ils seront associés à la même valeur de statistique (en termes absolu) et par conséquent seront associés à la même *p*-valeur. Si l'effet observé ne se situe *pas* à égale distance des deux limites de la zone d'équivalence, alors il suffira de tester l'hypothèse nulle qui se rapporte à la limite la plus proche de l'effet observé. En effet, le test associé à cette limite sera nécessairement celui qui est associé à la plus petite valeur de statistique et donc, à la plus grande *p*-valeur (???).

A titre d'illustration, la Figure 2 montre. . . (expliquer la figure: le carré = diff de moyenne observée, etc.)

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<sup>3</sup> il existe des alternatives au TOST qui sont très légèrement plus puissantes, mais le gain marginal en termes de puissance est contrebalancé par un niveau de complexité beaucoup plus élevé (Meyners, 2012).

Statistiquement parlant, montrer que les deux hypothèses nulles peuvent être rejetées, en considérant un risque  $\alpha$  donné, équivaut à montrer que l'intervalle de confiance à  $(1 - 2 \times \alpha)\%$  est entièrement inclus dans la zone d'équivalence (???). L'avantage de l'approche par intervalles de confiance est que ces derniers peuvent généralement être calculés dans la plupart des logiciels statistiques, contrairement aux tests d'équivalence qui ne sont pas encore disponibles partout (???)

## Avantage des tests d'équivalence

On ne peut falsifier une théorie si elle ne prédit pas une taille d'effet. Si elle ne le fait pas, même si on ne parvient pas à répliquer un effet significatif, on pourrait encore argumenter qu'il y a bien un effet mais plus petit que ce que l'étude était capable de détecter (Morey & Lakens, 2017, cités par ???).

## Conclusions possibles

- 1) statistical indeterminacy (???)
- 2) equivalence (???)

Lorsqu'ils sont combinés avec un test traditionnel pour lequel l'hypothèse nulle est l'absence d'effet, les TOST peuvent amener à 4 conclusions distinctes: 1) Equivalent et non statistiquement différent de 0: on conclura à l'absence de différence ayant un intérêt pratique (???). 2) Statistiquement différent de 0, mais équivalent: il existe un effet non nul, mais trop petit pour avoir un intérêt. C'est ce qui peut typiquement arriver avec des échantillons trop larges (résultant en des tests "trop" puissants, (???))

3) Statistiquement différent de 0, et on ne peut conclure à l'équivalence: on conclura alors à la présence d'un effet (???) 4) Ni statistiquement différent de 0 ni équivalent (quand les données sont si imprécises qu'on ne peut rien tirer comme conclusion)

→ A illustrer quand Jamovi le voudra!

## Définir les limites de la zone d'équivalence:LE sesoi

L'aspect le plus compliqué dans la réalisation du TOST est la définition des bornes d'équivalence. Il existe plusieurs stratégies:

- 1) définir comme limites la plus petite taille d'effet pour laquelle on peut atteindre une puissance de détection suffisante (déterminé par les ressources disponibles pour étudier l'effet, (???) → voir la section "Setting equivalence bounds" p. 359 mais je crois que j'en parle aussi dans la vidéo SOCLAB.
- 2) le SESOI peut parfois être fixé objectivement 3) Idéalement basé sur une analyse coût-bénéfice). Attention: bien sûr une dimension subjective dans la définition des coûts et des bénéfices. Attention: le SESOI doit être déterminé AVANT et INDEPENDAMMENT des données.

## Les 4 conclusions possibles quand combiné au NHST classique

Voir (???) p. 357. J'en parle ou po?

## Morceau qui servira peut-être (ou pas)

D'après (???), un test d'hypothèse (selon l'approche de Neyman-Pearson) vaut la peine à 2 conditions:

- 1) que l'hypothèse nulle soit assez plausible pour que son rejet puisse surprendre au moins certains;
- 2) le chercheur veut appliquer une procédure méthodol qui l'autorise à prendre des décisions quant à la manière d'agir, tout en contrôlant le taux d'erreur. Agir peut vouloir dire: adopter un traitement, une politique, une intervention, ou abandonner un domaine de recherche, modifier une manipulation, ou de faire un certain type de déclaration ou revendication.

\*One of the most widely suggested improvements of the use of p values is to replace null-hypothesis tests (where the goal is to reject an effect of exactly 0) with tests of range

predictions (where the goal is to reject effects that fall outside of the range of effects that is predicted or considered practically important) (???).

Un autre argument en défaveur de la p-valeur est la tendance des chercheurs à interpréter un effet NS comme l'acceptation de l'hypothèse nulle (Schmidt, 1996, cité par Harris, 1997).

Une fois encore, l'usage des tailles d'effet peut aider à cette fin, non pas en “remplaçant” les tests d'hypothèses, mais en les complétant. -> Tests d'équivalence.

## Remarques diverses

D'après Meyners, il existe pleins d'approches mais la différence entre elles est souvent négligeable (???).

## Quand veut-on montrer une absence d'effet?

Il arrive que des chercheurs souhaitent obtenir un soutien en faveur d'une absence d'effet. A titre d'illustration, (???) évoquent la situation où un chercheur désire écarter de potentielles covariables de leur analyse, en démontrant préalablement que ces variables ne sont pas corrélées à la variable dépendante. ESSAYER DE TROUVER ENCORE UN EXEMPLE. Or, on constate que très souvent, - Pour justifier la décision de pooler plusieurs groupes (???).



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# Equivalence Testing and the Second Generation P-Value.

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## Abstract

To move beyond the limitations of null-hypothesis tests, statistical approaches have been developed where the observed data are compared against a range of values that are equivalent to the absence of a meaningful effect. Specifying a range of values around zero allows researchers to statistically reject the presence of effects large enough to matter, and prevents practically insignificant effects from being interpreted as a statistically significant difference. We compare the behavior of the recently proposed second generation *p*-value (Blume, D'Agostino McGowan, Dupont, & Greevy, 2018) with the more established Two One-Sided Tests (TOST) equivalence testing procedure (Schuirmann, 1987). We show that the two approaches yield almost identical results under optimal conditions. Under suboptimal conditions (e.g., when the confidence interval is wider than the equivalence range, or when confidence intervals are asymmetric) the second generation *p*-value becomes difficult to interpret. The second generation *p*-value is interpretable in a dichotomous manner (i.e., when the SGPV equals 0 or 1 because the confidence intervals lies completely within or outside of the equivalence range), but this dichotomous interpretation does not require calculations. We conclude that equivalence tests yield more consistent *p*-values, distinguish between datasets that yield the same second generation *p*-value, and allow for easier control of Type I and Type II error rates.

**Keywords:** equivalence testing, second generation *p*-values, hypothesis testing, TOST, statistical inference

To test predictions researchers predominantly rely on null-hypothesis tests. This statistical approach can be used to examine whether observed data are sufficiently surprising under the null hypothesis to reject an effect that equals exactly zero. Null-hypothesis tests have an important limitation, in that this procedure can only reject the hypothesis that there is no effect, while scientists should also be able to provide statistical support for *equivalence*. When testing for equivalence researchers aim to examine whether an observed effect is too small to be considered meaningful, and therefore is practi-

cally equivalent to zero. By specifying a range around the null hypothesis of values that are deemed practically equivalent to the absence of an effect (i.e.,  $0 \pm 0.3$ ) the observed data can be compared against an *equivalence range* and researchers can test if a meaningful effect is absent (Hauck & Anderson, 1984; Kruschke, 2018; Rogers, Howard, & Vessey, 1993; Serlin & Lapsley, 1985; Spiegelhalter, Freedman, & Parmar, 1994; Wellek, 2010; Westlake, 1972).

Second generation *p*-values (SGPV) were recently proposed as a statistic that represents “the proportion of

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data-supported hypotheses that are also null hypotheses” (Blume et al., 2018). The researcher specifies an equivalence range around a null hypothesis of values that are considered practically equivalent to the null hypothesis. The SGPV measures the degree to which a set of data-supported parameter values falls within the interval null hypothesis. If the estimation interval falls completely within the equivalence range, the SGPV is 1. If the confidence interval falls completely outside of the equivalence range, the SGPV is 0. Otherwise the SGPV is a value between 0 and 1 that expresses the overlap of data-supported hypotheses and the equivalence range. When calculating the SGPV the set of data-supported parameter values can be represented by a confidence interval (CI), although one could also choose to use credible intervals or Likelihood support intervals (SI). When a confidence interval is used, the SGPV and equivalence tests such as the Two One-Sided Tests (TOST) procedure (Lakens, 2017; Meyners, 2012; Quertemont, 2011; Schuirmann, 1987) appear to have close ties, because both tests compare a confidence interval against an equivalence range. Here, we aim to examine the similarities and differences between the TOST procedure and the SGPV. We limit our analysis to continuous data sampled from a bivariate normal distribution. The TOST procedure also relies on the confidence interval around the effect. In the TOST procedure the data are tested against the lower equivalence bound in the first one-sided test, and against the upper equivalence bound in the second one-sided test (Lakens, Scheel, & Isager, 2018). For an excellent discussion of the strengths and weaknesses of different frequentist equivalence tests, including alternatives to the TOST procedure, see Meyners (2012). If both tests statistically reject an effect as extreme or more extreme than the equivalence bound, you can conclude the observed effect is practically equivalent to zero from a Neyman-Pearson approach to statistical inferences. Because one-sided tests are performed, one can also conclude equivalence by checking whether the  $1-2\alpha$  confidence interval (e.g., when the alpha level is 0.05, a 90% CI) falls completely within the equivalence bounds. Because both equivalence tests as the SGPV are based on whether and how much a confidence interval overlaps with equivalence bounds, it seems worthwhile to compare the behavior of the newly proposed SGPV to equivalence tests to examine the unique contribution of the SGPV to the statistical toolbox.

### The relationship between $p$ -values from TOST and SGPV when confidence intervals are symmetrical

The second generation  $p$ -value (SGPV) is calculated as:

$$p_{\delta} = \frac{|I \cap H_0|}{|I|} \times \max \left\{ \frac{|I|}{2|H_0|}, 1 \right\}$$

where  $I$  is the interval based on the data (e.g., a 95% confidence interval) and  $H_0$  is the equivalence range. The first term of this formula implies that the second generation  $p$ -value is the width of the confidence interval that overlaps with the equivalence range, divided by the total width of the confidence interval. The second term is a “small sample correction” (which will be discussed later) that comes into play whenever the confidence interval is more than twice as wide as the equivalence range. To examine the relation between the TOST  $p$ -value and the SGPV we can calculate both statistics across a range of observed effect sizes. Building on the example by Blume et al. (2018), in Figure 1  $p$ -values are plotted for the TOST procedure and the SGPV. The statistics are calculated for hypothetical one-sample  $t$ -tests for observed means ranging from 140 to 150 (on the x-axis). The equivalence range is set to  $145 \pm 2$  (i.e., an equivalence range from 143 to 147), the observed standard deviation is assumed to be 2, and the sample size is 30. For example, for the left-most point in Figure 1 the SGPV and the TOST  $p$ -value is calculated for a hypothetical study with a sample size of 30, an observed standard deviation of 2, and an observed mean of 140, where the  $p$ -value for the equivalence test is 1, and the SGPV is 0.

Our conclusions about the relationship between TOST  $p$ -values and SGPV hold for second generation  $p$ -values calculated from confidence intervals, and assuming data is sampled from a bivariate normal distribution. Readers can explore the relationship between TOST  $p$ -values and SGPV for themselves in an online Shiny app: [http://shiny.ijs.tue.nl/TOST\\_vs\\_SGPV/](http://shiny.ijs.tue.nl/TOST_vs_SGPV/).

The SGPV treats the equivalence range as the null-hypothesis, while the TOST procedure treats the values outside of the equivalence range as the null-hypothesis. For ease of comparison we can plot 1-SGPV (see Figure 2) to make the values more easily comparable. We see that the  $p$ -value from the TOST procedure and the SGPV follow each other closely. When we discuss the relationship between the  $p$ -values from TOST and the SGPV, we focus on their correspondence at three values, namely where the TOST  $p = 0.025$  and SGPV is 1, where the TOST  $p = 0.5$  and SGPV = 0.5, and where the TOST  $p = 0.975$  and SGPV = 1. These three values are important for the SGPV because they indicate the values at which the SGPV indicates the data should be interpreted as compatible with the null hypothesis (SGPV =

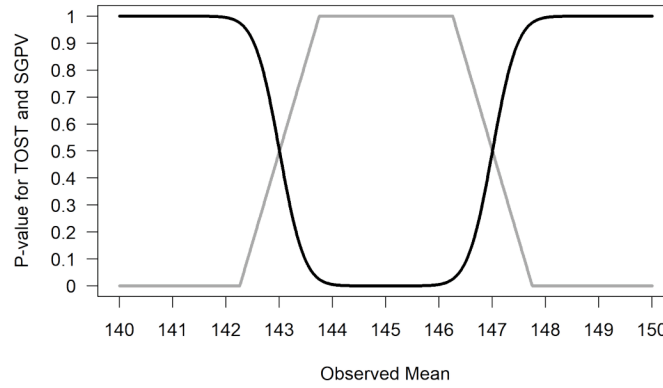


Figure 1. Comparison of  $p$ -values from TOST (black line) and SGPV (grey line) across a range of observed sample means (x-axis) tested against a mean of 145 in a one-sample  $t$ -test with a sample size of 30 and a standard deviation of 2, illustrating that when the TOST  $p$ -value = 0.5, the SGPV = 0.5, when the TOST  $p$ -value is 0.975, 1-SGPV = 1, and when the TOST  $p$ -value = 0.025, 1-SGPV = 0.

1), or with the alternative hypothesis (SGPV = 0), or when the data are strictly inconclusive (SGPV = 0.5).

These three points of overlap are indicated by the horizontal dotted lines in Figure 2 at TOST  $p$ -values of 0.975, 0.5, and 0.025.

When the observed sample mean is 145, the sample size is 30, and the standard deviation is 2, and we are testing against equivalence bounds of 143 and 147 using the TOST procedure for a one-sample  $t$ -test, the equivalence test is significant,  $t(29) = 5.48$ ,  $p < .001$ . Because the 95% CI falls completely within the equivalence bounds, the SGPV is 1 (see Figure 1). On the other hand, when the observed mean is 140, the equivalence test is not significant (the observed mean is far outside the equivalence range of 143 to 147),  $t(29) = -8.22$ ,  $p = 1$  (or more accurately,  $p > .999$  as  $p$ -values are bounded between 0 and 1). Because the 95% CI falls completely outside the equivalence bounds, the SGPV is 0 (see Figure 1).

#### SGPV as a uniform measure of overlap

It is clear the SGPV and the  $p$ -value from TOST are closely related. When confidence intervals are symmetric we can think of the SGPV as a straight line that is

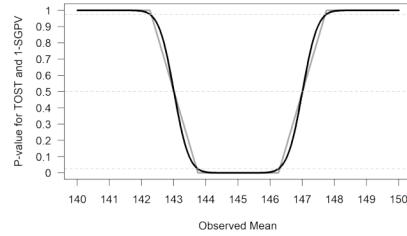


Figure 2. Comparison of  $p$ -values from TOST (black line) and 1-SGPV (grey line) across a range of observed sample means (x-axis) tested against a mean of 145 in a one-sample  $t$ -test with a sample size of 30 and a standard deviation of 2.

directly related to the  $p$ -value from an equivalence test for three values. When the TOST  $p$ -value is 0.5, the SGPV is also 0.5 (note that the reverse is not true). The

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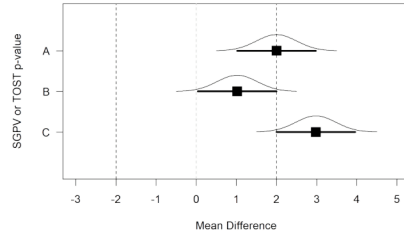


Figure 3. Means, normal distribution, and 95% CI for three example datasets that illustrate the relationship between  $p$ -values from TOST and SGPV.

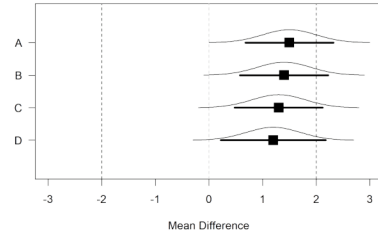


Figure 4. Means, normal distribution, and 95% CI for samples where the observed population mean is 1.5, 1.4, 1.3, and 1.2.

SGPV is 50% when the observed mean falls exactly on the lower or upper equivalence bound, because 50% of the symmetrical confidence interval overlaps with the equivalence range. When the observed mean equals the equivalence bound, the difference between the mean in the data and the equivalence bound is 0, the  $t$ -value for the equivalence test is also 0, and thus the  $p$ -value is 0.5 (situation A, Figure 3).

Two other points always have to overlap. When the 95% CI falls completely inside the equivalence region, and one endpoint of the confidence interval is exactly equal to one of the equivalence bounds (see situation B in Figure 3) the TOST  $p$ -value (which relies on a one-sided test) is always 0.025, and the SGPV is 1. Note that when sample sizes are small or equivalence bounds are narrow, small  $p$ -values for the TOST or a SGPV = 1 might not be observed in practice if too few observations are collected. The third point where the SGPV and the  $p$ -value from the TOST procedure should overlap is where the 95% CI falls completely outside of the equivalence range, but one endpoint of the confidence interval is equal to the equivalence bound (see situation C in Figure 3), when the  $p$ -value will always be 0.975, and the SGPV is 0. Note that this situation is in essence a minimum-effect test (Murphy, Myers, & Wolach, 2014). The goal of a minimum-effect is not just to reject a difference of zero, but to reject the smallest effect size of interest (i.e., the equivalence bounds). An equivalence test and minimum effect test against the same equivalence bound are complementary, and when a TOST  $p$ -value is larger than 0.975, the  $p$ -value for the minimum effect test is smaller than 0.05 (and therefore the minimum effect test provides no additional information that can not be derived from the  $p$ -value from the equivalence test).

The SGPV summarizes the information from an equivalence test (and the complementary minimum-effect test). These can be two relevant questions to ask, although it often makes sense to combine an equivalence test and a null-hypothesis test instead (Lakens et al., 2018).

For example, in Figure 4 we have plotted four SGPV's. From A to D the SGPV is 0.76, 0.81, 0.86, and 0.91. The difference in the percentage of overlap between A and B (-0.05) is identical to the difference in the percentage of overlap between C and D as the mean gets 0.1 closer to the test value (-0.05). As the observed mean in a one-sample  $t$ -test lies closer to the test value, from situation A to D, the difference in the overlap changes uniformly. As we move the observed mean closer to the test value in steps of 0.1 across A to D the  $p$ -value calculated for normally distributed data are not uniformly distributed. The probability of observing data more extreme than the upper bound of 2 is (from A to D) 0.16, 0.12, 0.08, and 0.05. As we can see, the difference between A and B (0.04) is not the same as the difference between C and D (0.03). Indeed, the difference in  $p$ -values is the largest as you start at  $p = 0.5$  (when the observed mean falls on the test value), which is why the line in Figure 1 is the steepest at  $p = 0.5$ . Note that where the SGPV reaches 1 or 0,  $p$ -values closely approximate 0 and 1, but never reach these values.

#### When different $p$ -values for equivalence tests yield the same SGPV

There are three situations where  $p$ -values for TOST differentiate between observed results, while the SGPV does not differentiate. The first two situations were discussed before and can be seen in Figure 1. When

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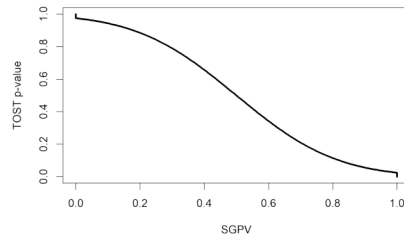


Figure 5. The relationship between  $p$ -values from the TOST procedure and the SGPV for the same scenario as in Figure 1.

the SGPV is either 0 or 1,  $p$ -values from the equivalence test fall between 0.975 and 1 or between 0 and 0.025. Where the SGPV is 1 as long as the confidence interval falls completely within the equivalence bounds, the  $p$ -value for the TOST continues to differentiate between results as a function of how far the confidence interval lies within the equivalence bounds (the further the confidence interval is from both bounds, the lower the  $p$ -value). The easiest way to see this is by plotting the SGPV against the  $p$ -value from the TOST procedure. The situations where the  $p$ -values from the TOST procedure continue to differentiate based on how extreme the results are, but the SGPV is a fixed value are indicated by the parts of the curve where there are vertical straight lines at second generation  $p$ -values of 0 and 1.

A third situation in which the SGPV remains stable across a range of observed effects, while the TOST  $p$ -value continues to differentiate, is whenever the CI is wider than the equivalence range, and the CI overlaps with the upper and lower equivalence bound. When the confidence interval is more than twice as wide as the equivalence range the SGPV is set to 0.5. Blume et al. (2018) call this the “small sample correction factor”. However, it is not a correction in the typical sense of the word, since the SGPV is not adjusted to any “correct” value. When the normal calculation would be “misleading” (i.e., the SGPV would be small, which normally would suggest support for the alternative hypothesis, but at the same time all values in the equivalence range are supported), the SGPV is set to 0.5 which according to Blume and colleagues signals that the SGPV is “uninformative”. Note that the CI can be twice as wide as the equivalence range whenever the sample size is small (and the confidence interval width is large) or when

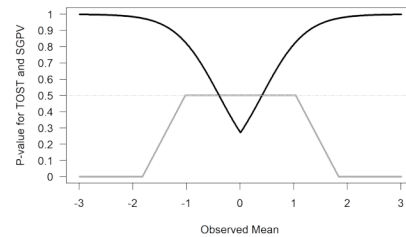


Figure 6. Comparison of  $p$ -values from TOST (black line) and SGPV (grey line) across a range of observed sample means (x-axis). Because the sample size is small ( $n = 10$ ) and with a standard deviation of 2 the CI is more than twice as wide as the equivalence range (set to -0.4 to 0.4), the SGPV is set to 0.5 (horizontal light-grey line) across a range of observed means.

then equivalence range is narrow. It is therefore not so much a “small sample correction” as it is an exception to the typical calculation of the SGPV whenever the ratio of the confidence interval width to the equivalence range exceeds 2:1 and the CI overlaps with the upper and lower bounds.

We can examine this situation by calculating the SGPV and performing the TOST for a situation where sample sizes are small and the equivalence range is narrow, such that the CI is more than twice as large as the equivalence range (see Figure 6). When the two statistics are plotted against each other we can see where the SGPV is the same while the TOST  $p$ -value still differentiates different observed means (indicated by straight lines in the curve, see Figure 7). We see the SGPV is 0.5 for a range of observed means where the  $p$ -value from the equivalence test still varies. It should be noted that in these calculations the  $p$ -values for the TOST procedure are *never* smaller than 0.05 (i.e., they do not get below 0.05 on the y-axis). In other words, we cannot conclude equivalence based on any of the observed means. This happens because we are examining a scenario where the 90% CI is so wide that it never falls completely within the two equivalence bounds.

As Lakens (2017) notes: “in small samples (where CIs are wide), a study might have no statistical power (i.e., the CI will always be so wide that it is necessarily wider than the equivalence bounds).” None of the  $p$ -values based on the TOST procedure are below 0.05, and thus, in the long run we have 0% power.

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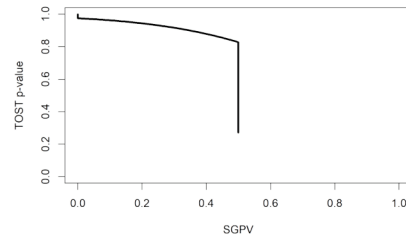


Figure 7. The relationship between  $p$ -values from the TOST procedure and the SGPV for the same scenario as in Figure 6.

The  $p$ -value from the TOST procedure still differentiates observed means, while the SGPV does not, when the CI is wider than the equivalence range (so the precision is low) and overlaps with the upper and lower equivalence bound, but the CI is *not* twice as wide as the equivalence range. In the example below, we see that the CI is only 1.79 times as wide as the equivalence bounds, but the CI overlaps with the lower and upper equivalence bounds (Figure 8). This means the SGPV is not set to 0.5, but it is constant across a range of observed means, while the TOST  $p$ -value is not constant across this range.

If the observed mean would be somewhat closer to 0, or further away from 0, the SGPV remains constant (the CI width does not change, and it completely over-

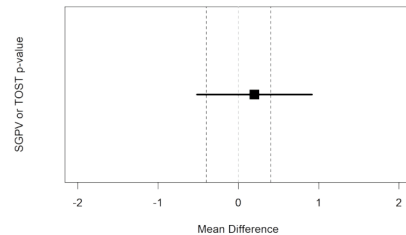


Figure 8. Example of a 95% CI that overlaps with the lower and upper equivalence bound (indicated by the vertical dotted lines).

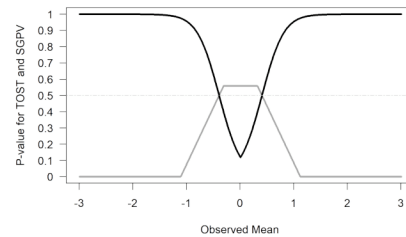


Figure 9. Comparison of  $p$ -values from TOST (black line) and SGPV (grey line) across a range of observed sample means (x-axis). The sample size is small ( $n = 10$ ), but because the sd is half as big as in Figure 7 (1 instead of 2) the CI is less than twice as wide as the equivalence range (set to -0.4 to 0.4). The SGPV is not set to 0.5 (horizontal light grey line) but reaches a maximum slightly above 0.5 across a range of observed means.

laps with the equivalence range) while the  $p$ -value for the TOST procedure does vary. We can see this in Figure 9 below. The SGPV is not set to 0.5 across a range of means. How high the SGPV will be for a CI that is not twice as wide as the equivalence range, but overlaps with the lower and upper equivalence bounds, depends on the width of the CI and the equivalence range.

If we once more plot the two statistics against each other we see the SGPV is 0.56 for a range of observed means where the  $p$ -value from the equivalence test still varies, as indicated by the straight section of the line (Figure 10).

To conclude this section, there are situations where the  $p$ -value from the TOST procedure continues to differentiate, while the SGPV does not. Therefore, interpreted as a continuous statistic, the SGPV is more limited than the  $p$ -value from the TOST procedure.

#### The relation between equivalence tests and SGPV for asymmetrical confidence intervals around correlations

So far we have only looked at the relation between equivalence tests and the SGPV when confidence intervals are symmetric (e.g., for confidence intervals around mean differences). For correlations, which are bound between -1 and 1, confidence intervals are only symmetric for a correlation of exactly 0. The confidence interval for a correlation becomes increasingly asymmetric



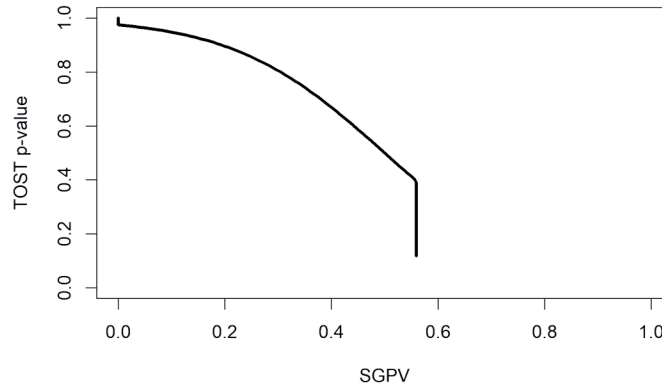


Figure 10. The relationship between  $p$ -values from the TOST procedure and the SGPV for the same scenario as in Figure 9.

as the observed correlation nears -1 or 1. For example, with ten observations, an observed correlation of 0 has a symmetric 95% confidence interval ranging from -0.63 to 0.63, while an observed correlation of 0.7 has an asymmetric 95% confidence interval ranging from 0.13 to 0.92. Note that calculating confidence intervals for a correlation involves a Fisher's  $z$ -transformation, which transforms values such that they are approximately normally  $z$ -distributed, which allows one to compute symmetric confidence intervals. These confidence intervals are then retransformed into a correlation, where the confidence intervals are asymmetric if the correlation is not exactly zero.

The effect of asymmetric confidence intervals around correlations is most noticeable at smaller sample sizes. In Figure 11 we plot the  $p$ -values from equivalence tests and the SGPV (again plotted as 1-SGPV for ease of comparison) for correlations. The sample size is 30 pairs of observations, and the lower and upper equivalence bounds are set to -0.45 and 0.45, with an alpha of 0.05. As the observed correlation in the sample moves from -0.99 to 0 the  $p$ -value from the equivalence test becomes smaller, as does 1-SGPV. The pattern is quite similar to that in Figure 2. The  $p$ -value for the TOST procedure and 1-SGPV are still related as discussed above, with TOST  $p$ -values of 0.975 and 0.025 corresponding to a 1-

SGPV of 1 and 0, respectively. There are two important differences, however. First of all, the SGPV is no longer a straight line, but a curve, due to the asymmetry in the 95% CI. Second, and most importantly, the  $p$ -value for the equivalence test and the SGPV do no longer overlap at  $p = 0.5$ .

The reason that the equivalence test and SGPV no longer overlap is due to asymmetric confidence intervals. If the observed correlation falls exactly on the equivalence bound the  $p$ -value for the equivalence test is 0.5. In the equivalence test for correlations the  $p$ -value is computed based on a  $z$ -transformation which better controls error rates (Goertzen & Cribbie, 2010). This transformation is computed as follows, where  $r$  is the observed correlation and  $\rho$  is the theoretical correlation under the null:

$$z = \frac{\frac{\log(\frac{1+r}{1-r})}{2} - \frac{\log(\frac{1+\rho}{1-\rho})}{2}}{\sqrt{\frac{1}{n-3}}}$$

Because the  $z$ -distribution is symmetric, the probability of observing the observed or more extreme  $z$ -score, assuming the equivalence bound is the true effect size, is 50%. However, because the  $r$  distribution is not symmetric, this does not mean that there is always a 50% probability of observing a correlation smaller or larger

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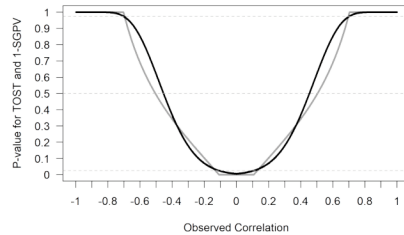


Figure 11. Comparison of  $p$ -values from TOST (black line) and 1-SGPV (grey curve) across a range of observed sample correlations (x-axis) tested against equivalence bounds of  $r = -0.45$  and  $r = 0.45$  with  $n = 30$  and an alpha of 0.05.

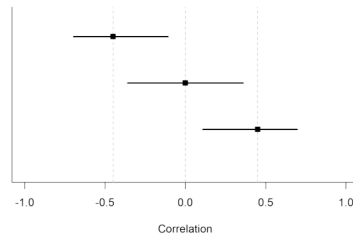


Figure 12. Three 95% confidence intervals for observed effect sizes of  $r = -0.45$ ,  $r = 0$ , and  $r = 0.45$  for  $n = 30$ . Only the confidence interval for  $r = 0$  is symmetric.

than the true correlation. As can be seen in Figure 12, the proportion of the confidence interval that overlaps with the equivalence range is larger than 50% when the observed correlations are  $r = -0.45$  and  $r = 0.45$ , meaning that the two second generation  $p$ -values associated with these correlations are larger than 50%. Because the confidence intervals are asymmetric around the observed effect size of 0.45 (ranging from 0.11 to 0.70) according to Blume et al. (2018) 58.11% of the data-supported hypotheses are null hypotheses, and therefore 58.11% of the data-supported hypotheses are compatible with the null premise.

The further away from 0, the larger the SGPV when the observed mean falls on the equivalence bound. The

SGPV is the proportion of values in a 95% confidence interval that overlap with the equivalence range, but not the probability that these values will be observed. In the most extreme case (i.e., a sample size of 4, and equivalence bounds set to  $r = -0.99$  and  $0.99$ , with a true correlation of 0.99) 97.60% of the confidence interval overlaps with the equivalence range, even though in the long run only 36% of the correlations observed in the future will fall in this range.

It should be noted that in larger sample sizes the SGPV is closer to 0.5 whenever the observed correlation falls on the equivalence bound, but this extreme example nevertheless clearly illustrates the difference between question the SGPV answers, and the question a  $p$ -value answers. The conclusion of this section on asymmetric confidence intervals is that a SGPV of 1 or 0 can still be interpreted as a  $p < 0.025$  or  $p > 0.975$  in an equivalence test, since the SGPV and  $p$ -value for the TOST procedure are always directly related at the values  $p = 0.025$  and  $p = 0.975$ . Although Blume et al. (2018) state that “the degree of overlap conveys how compatible the data are with the null premise” this definition of what the SGPV provides does not hold for asymmetric confidence intervals. Although a SGPV of 1 or 0 can be directly interpreted, a SGPV between 0 and 1 is not interpretable as “compatibility with the null hypothesis” under the assumption of a bivariate normal distribution, and the generalizability of this statement needs to be examined beyond normal bivariate distributions. Indeed, Blume and colleagues write in the supplemental material that “The magnitude of an inconclusive second-generation  $p$ -value can vary slightly when the effect size scale is transformed. However definitive findings, i.e. a  $p$ -value of 0 or 1 are not affected by the scale changes.”

#### What are the Relative Strengths and Weaknesses of Equivalence Testing and the SGPV?

When introducing a new statistical method, it is important to compare it to existing approaches and specify its relative strengths and weaknesses. Here, we aimed to compare the SGPV against equivalence tests based on the TOST procedure. First of all, even though a SGPV of 1 or 0 has a clear interpretation (we can reject effects outside or inside the equivalence range), intermediate values are not as easy to interpret (especially for effects that have asymmetric confidence intervals). In one sense, they are what they are (the proportion of overlap), but it can be unclear what this number tells us about the data we have collected. This is not too problematic, since the main use of the SGPV (e.g., in all examples provided by Blume and colleagues) seems to be to examine whether the SGPV is 0, 1, or inconclusive.



As already mentioned, this interpretation of a SGPV is very similar to the Neyman-Pearson interpretation of an equivalence test and a minimum effect tests (which are complementary). The difference is that where a SGPV of 1 can be interpreted as  $p < .025$ , equivalence tests provide exact  $p$ -values, and they continue to differentiate between for example  $p = 0.024$  and  $p = 0.002$ . Whether this is desirable depends on the perspective that is used. From a Neyman-Pearson perspective on statistical inferences the main conclusion is based on whether or not  $p < \alpha$ , and thus an equivalence test and SGPV can be performed by simply checking whether the confidence interval falls within the equivalence range, just as a null-hypothesis test can be performed by checking whether the confidence interval contains zero or not. At the same time, it is recommended to report exact  $p$ -values (American Psychological Association, 2010), and exact  $p$ -values might provide information of interest to readers about how precisely how surprising the data, or more extreme data, is under the null model. Some researchers might be interested in combining an equivalence test with a null-hypothesis significance test. This allows a researcher to ask whether there is an effect that is statistically different from zero, and whether effect sizes that are considered meaningful can be rejected. Equivalence tests combined with null-hypothesis tests classify results into four possible categories, and for example allow researchers to conclude an effect is significant *and* equivalent (i.e., statistically different from zero, but also too small to be considered meaningful; see Lakens et al., 2018).

An important issue when calculating the SGPV is its reliance on the “small sample correction”, where the SGPV is set to 0.5 whenever the ratio of the confidence interval width to the equivalence range exceeds 2:1 and the CI overlaps with the upper and lower bounds. This exception to the normal calculation of the SGPV is introduced to prevent misleading values. Without this correction it is possible that a confidence interval is extremely wide, and an equivalence range is extremely narrow, which without the correction would lead to a very low value for the SGPV. Blume et al. (2018) suggest that under such a scenario “the data favor alternative hypotheses”, even when a better interpretation would be that there is not enough data to accurately estimate the true effect compared to the width of the equivalence range. Although it is necessary to set the SGPV to 0.5 whenever the ratio of the confidence interval width to the equivalence range exceeds 2:1, it leads to a range of situations where the SGPV is set to 0.5, while the  $p$ -value from the TOST procedure continues to differentiate (see for example Figure 6). An important benefit of equivalence tests is that it does not need

such a correction to prevent misleading results.

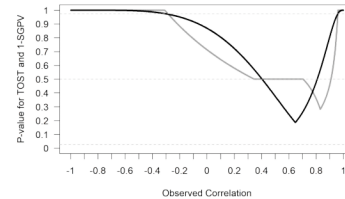


Figure 13. Comparison of  $p$ -values from TOST (black line) and 1-SGPV (grey curve) across a range of observed sample correlations (x-axis) tested against equivalence bounds of  $r = 0.4$  and  $r = 0.8$  with  $n = 10$  and an alpha of 0.05.

As a more extreme example of the peculiar behavior of the “small sample correction” as currently implemented in the calculation of the SGPV, see Figure 13. In this figure observed correlations (from a sample size of 10) from  $-0.99$  to  $0.99$  are tested against an equivalence range from  $r = 0.4$  to  $r = 0.8$ . We can see the SGPV has a peculiar shape because it is set to 0.5 for certain observed correlations, even though there is no risk of a “misleading” SGPV in this range. This example suggests that the current implementation of the “small sample correction” could be improved. If, on the other hand, the SGPV is mainly meant to be interpreted when it is 0 or 1, it might be preferable to simply never apply the “small sample correction”.

Blume et al. (2018) claim that when using the SGPV “Adjustments for multiple comparisons are obviated” (p. 15). However, this is not correct. Given the direct relationship between TOST and SGPV highlighted in this manuscript (where a TOST  $p = 0.025$  equals SGPV = 1, as long as the SGPV is calculated based on confidence intervals, and assuming data are sampled from a continuous bivariate normal distribution), not correcting for multiple comparisons will inflate the probability of concluding the absence of a meaningful effect based on the SGPV in exactly the same way as it will for equivalence tests. Whenever statistical tests are interpreted as support for a hypothesis (e.g., SGPV = 0 or SGPV = 1), it is possible to do so erroneously, and if researchers want to control error rates, they need to correct for multiple comparisons.

### Conclusion

We believe that our explanation of the similarities between the TOST procedure and the SGPV provides context to interpret the contribution of second generation  $p$ -values to the statistical toolbox. The novelty of the SGPV can be limited when confidence intervals are asymmetrical or wider than the equivalence range. There are strong similarities with  $p$ -values from the TOST procedure, and in all situations where the statistics yield different results, the behavior of the  $p$ -value from the TOST procedure is more consistent. We hope this overview of the relationship between the SGPV and equivalence tests will help researchers to make an informed decision about which statistical approach provides the best answer to their question. Our comparisons show that when proposing alternatives to null-hypothesis tests, it is important to compare new proposals to already existing procedures. We believe equivalence tests achieve the goals of the second generation  $p$ -value while allowing users to easily control error rates, and while yielding more consistent statistical outcomes.

### Authors Note

All code associated with this article, including the reproducible manuscript, is available from [https://github.com/Lakens/TOST\\_vs\\_SGPV](https://github.com/Lakens/TOST_vs_SGPV) and <https://osf.io/8crkg/>. The preprint can be found at <https://psyarxiv.com/7k6ay/>.

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### Author Contributions

DL conceptualized the idea, both authors wrote and revised this manuscript.

### References

- American Psychological Association (Ed.). (2010). *Publication manual of the American Psychological Association* (6th ed.). Washington, DC: American Psychological Association.
- Blume, J. D., D'Agostino McGowan, L., Dupont, W. D., & Greevy, R. A. (2018). Second-generation  $p$ -values: Improved rigor, reproducibility, & transparency in statistical analyses. *PLOS ONE*, 13(3), e0188299. doi:[10.1371/journal.pone.0188299](https://doi.org/10.1371/journal.pone.0188299)
- Goertzen, J. R., & Cribbie, R. A. (2010). Detecting a lack of association: An equivalence testing approach. *British Journal of Mathematical and Statistical Psychology*, 63(3), 527–537. doi:[10.1348/000711009X475853](https://doi.org/10.1348/000711009X475853)
- Hauck, D. W. W., & Anderson, S. (1984). A new statistical procedure for testing equivalence in two-group comparative bioavailability trials. *Journal of Pharmacokinetics and Biopharmaceutics*, 12(1), 83–91. doi:[10.1007/BF01063612](https://doi.org/10.1007/BF01063612)
- Kruschke, J. K. (2018). Rejecting or Accepting Parameter Values in Bayesian Estimation. *Advances in Methods and Practices in Psychological Science*, 1(2), 270–280. doi:[10.1177/2515245918771304](https://doi.org/10.1177/2515245918771304)
- Lakens, D. (2017). Equivalence Tests: A Practical Primer for  $t$  Tests, Correlations, and Meta-Analyses. *Social Psychological and Personality Science*, 8(4), 355–362. doi:[10.1177/1948550617697177](https://doi.org/10.1177/1948550617697177)
- Lakens, D., Scheel, A. M., & Isager, P. M. (2018). Equivalence Testing for Psychological Research: A Tutorial. *Advances in Methods and Practices in Psychological Science*, 1(2), 259–269. doi:[10.1177/2515245918770963](https://doi.org/10.1177/2515245918770963)
- Meyners, M. (2012). Equivalence tests A review. *Food Quality and Preference*, 26(2), 231–245. doi:[10.1016/j.foodqual.2012.05.003](https://doi.org/10.1016/j.foodqual.2012.05.003)
- Murphy, K. R., Myors, B., & Wolach, A. H. (2014). *Statistical power analysis: A simple and general model for traditional and modern hypothesis tests* (Fourth edition.). New York: Routledge, Taylor & Francis Group.
- Quertemont, E. (2011). How to Statistically Show the Absence of an Effect. *Psychologica Belgica*, 51(2), 109–127. doi:[10.5334/pb-51-2-109](https://doi.org/10.5334/pb-51-2-109)
- Rogers, J. L., Howard, K. I., & Vessey, J. T. (1993). Using significance tests to evaluate equivalence between two experimental groups. *Psychological Bulletin*, 113(3), 553–565. doi:[http://dx.doi.org/10.1037/0033-2909.113.3.553](https://doi.org/http://dx.doi.org/10.1037/0033-2909.113.3.553)

- Schuurmann, D. J. (1987). A comparison of the two one-sided tests procedure and the power approach for assessing the equivalence of average bioavailability. *Journal of Pharmacokinetics and Biopharmaceutics*, 15(6), 657–680. doi:<https://doi.org/10.1007/BF01068419>
- Serlin, R. C., & Lapsley, D. K. (1985). Rationality in psychological research: The good-enough principle. *American Psychologist*, 40(1), 73–83. doi:<http://dx.doi.org/10.1037/0003-066X.40.1.73>
- Spiegelhalter, D. J., Freedman, L. S., & Parmar, M. K. (1994). Bayesian approaches to randomized trials. *Journal of the Royal Statistical Society. Series A (Statistics in Society)*, 357–416. doi:[10.2307/2983527](https://doi.org/10.2307/2983527)
- Wellek, S. (2010). *Testing statistical hypotheses of equivalence and noninferiority* (2nd ed.). Boca Raton: CRC Press.
- Westlake, W. J. (1972). Use of Confidence Intervals in Analysis of Comparative Bioavailability Trials. *Journal of Pharmaceutical Sciences*, 61(8), 1340–1341. doi:[10.1002/JPS.2600610845](https://doi.org/10.1002/JPS.2600610845)

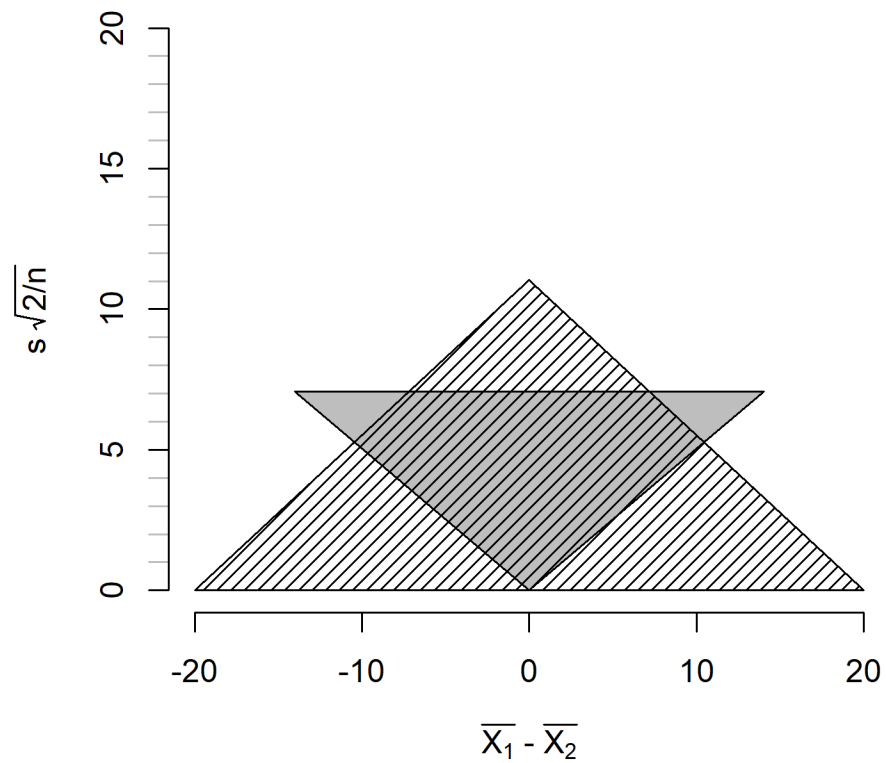


Figure 1. Région d'équivalence pour l'approche de la puissance (zone grisée) et pour le TOST (zone hachurée), pour l'exemple où  $|\theta|=20$ ,  $n = 50$  et  $\alpha = .05$

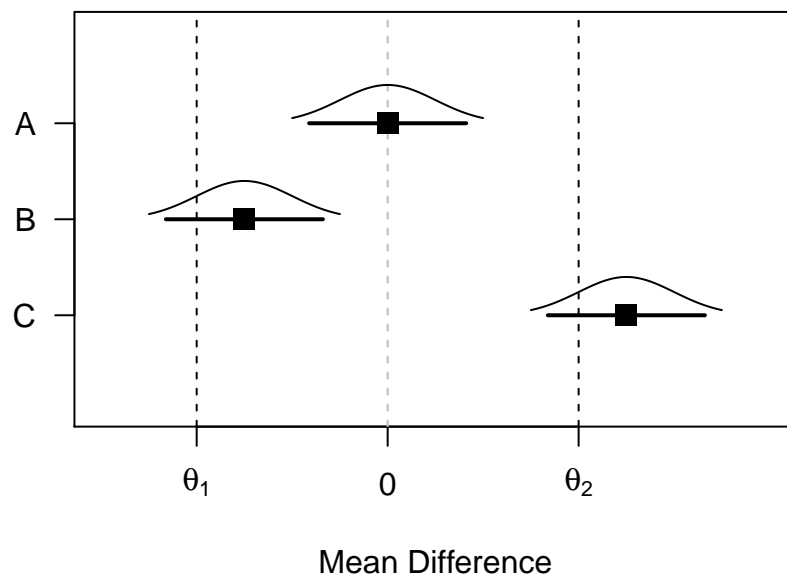


Figure 2. Différence de moyennes ( $\bar{X}_1 - \bar{X}_2$ ), distribution d'échantillonnage et IC à  $1 - 2\alpha\%$  autour de la différence de moyennes ( $\bar{X}_1 - \bar{X}_2$ ) pour 3 exemples de TOST.