

The Use of Signal Detection Theory to Evaluate Aphasia Diagnostic Accuracy and Clinician Bias

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ERRATUM

In Volume 21 (1993) the article "The Use of Signal Detection Theory to Evaluate Aphasia Diagnostic Accuracy and Clinician Bias," by Donald A. Robin and Malcolm R. McNeil, was printed incompletely. The correct version follows.

Diagnosing aphasia involves differentiating it from other speech-language pathologies that resemble it. This differentiation is difficult because the symptom complex required for most definitions of aphasia include a variety of behaviors shared by other language and communication impairments such as those associated with confusion, dementia, right hemisphere disorders, traumatic brain injury, schizophrenia, malingering, and some forms of motor speech disorders. The validity and reliability of the differential diagnoses among the neurogenic and psychiatric pathologies have been approached from a variety of perspectives. For example, Halpern, Darley, and Brown (1973) used the pattern of performance across a variety of speech, language, and other cognitive tasks to construct a differential profile of performance for confused, demented, and aphasic patients. Wertz and Rosenbek (1971) also profiled differentiation of apraxia of speech from aphasia.

Recently, discriminant function analyses have been used to construct a statistical *profile* for the differentiation of aphasia from conditions resembling it. The *Porch Index of Communicative Ability (PICA)* (Porch, 1971) has been used to differentiate aphasia from malingering (Porch, Frieden, & Porec, 1977), normal (Brauer, McNeil, Duffy, Keith, & Collins, 1990), and

from right hemisphere damage (Brauer et al., 1988). The *Revised Token Test (RTT)* (McNeil & Prescott, 1978) has been used to differentiate left hemisphere damaged-aphasic, right hemisphere damaged-nonaphasic and normal individuals using discriminant analyses (McNeil, Brauer, & Prescott, 1988).

While the statistical profile analyses provide useful clinical insight into general areas of impaired and unimpaired functions, they have limited information for generating differential diagnosis standards. The discriminant function studies offer information on the accuracy and bias of a particular test; however, they provide no opportunity to evaluate individual clinicians. It would be useful to have a method of assessing the accuracy and bias (the tendency to favor one outcome over another) for individual clinicians. The theory of signal detection (TSD) quantifies accuracy and response bias for test or individual clinicians. This paper (1) outlines the major tenets of TSD, (2) discusses TSD application to diagnosis in aphasia, and (3) presents the results of an initial application of TSD in assessing accuracy and bias in aphasia diagnosis using *PICA* test data.

OVERVIEW OF TSD AND ITS RELATION TO DIAGNOSTICS

The assumption was made that no diagnostic system, test, or clinician is perfectly accurate no matter how **objective** each may be. Moreover, the valid evaluation of the accuracy and bias of diagnostic systems is critical in understanding their potential *clinical value* and diagnostic accuracy must precede the evaluation of treatment efficacy and cost-benefits (Swets, 1988; Swets & Pickett, 1982).

Statistical decision theory, translated into TSD, can be traced to Blackwell in the early fifties (Blackwell, 1953; Blackwell, Prichard, & Ohmart, 1954). For three decades Swets and colleagues (e.g., Green & Swets, 1966; Swets, 1959, 1988; Swets & Pickett, 1982) have pioneered the theory's application to diagnostic systems. Originally developed in World War II to assist in submarine detection, TSD has been used widely in psychophysical experiments (e.g., Gescheider, 1985; Green & Swets, 1966) in order to assess the *true* accuracy of detecting a stimulus.

In any discrimination task there are two possible states: Noise (N) and Signal-plus-Noise (SN). The observer's job is to determine if SN is present or just N. Noise arises from external sources (e.g., the wind, rain, traffic or human noise such as speech or babble) as well as internal sources (e.g., from the observer's physiology). Given these two states (N and SN), there are four possible judgment outcomes at any given time. As

Table 1. Four Possible Outcomes When Presented With Signal Plus Noise (SN) or Noise (N)

		<i>Subject Response</i>	
		SN	N
Actual State	SN	HIT	MISS
	N	FP	TN

shown in Table 1, an observer may say SN when SN is present, a *HIT* (correct detection or true positive). Or the observer may say N when SN is present, a *MISS* (false negative). Likewise, the observer may say SN when N is present, (a *FALSE POSITIVE* [FP] or false alarm) or the observer may say N when N is present, (a *TRUE NEGATIVE* [TN] or correct rejection).

Knowledge of the raw numbers in each of the cells compared to the total number of trials allows one to calculate the probability of each event. As shown in the hypothetical case in Table 2, there are a total of 200 trials: 100 are SN and 100 are N. The observer said SN 60 times and N 40 times in 100 SN trials. Thus, in this example, the probability of a hit, $P(\text{hit})$, is .60 and the probability of a miss, $P(\text{miss})$, is .40. Of the 100 trials in which N was present, the observer said SN 25 times and N 75 times; thus, $P(\text{FP})$ is .25 and $P(\text{TN})$ is .75. $P(\text{miss}) = 1 - P(\text{hit})$ and $P(\text{TN}) = 1 - P(\text{FP})$. Therefore, the percent of hits and FPs are needed to determine true sensitivity (accuracy) and bias. A point that cannot be overly stressed is that accuracy cannot be assessed based on percent correct (hits) alone; FPs must be accounted for as well. This point will be elaborated below.

Knowledge of hits, misses, FPs, and TNs (hits and FPs) allows for TDS application to evaluate a diagnostic system. Consider SN to be the abnormal case (e.g., an individual with aphasia) and N to be any other state (e.g., normal). How N is defined can be varied systematically to provide different degrees of precision when evaluating a diagnostic system. Given any diagnostic decision, the clinician is faced with the problem of choosing SN or N, and any of the four outcomes (hits, misses, FP, TN) are possible.

A hypothetical example of diagnostic accuracy will serve to demonstrate the need for a better accuracy measure than percent hits alone. Clinician Y is 100% correct in the identification of aphasia when aphasia

Table 2. Hypothetical Data and Response Frequencies When Presented With Signal Plus Noise (SN) or Noise (N)

		<i>Subject Response</i>		TOTAL #
		SN	N	
Actual State	SN	60	40	100
	N	25	75	100

$P(\text{HIT}) = .60$ $P(\text{MISS}) = .40$ $P(\text{FP}) = .25$ $P(\text{TN}) = .75$
 $P(\text{MISS}) = 1 - P(\text{HIT})$ $P(\text{TN}) = 1 - P(\text{FP})$

is present (i.e., $P[\text{hit}] = 1.0$ and $P[\text{miss}] = 0.0$). Clinician Z is only 70% correct in saying aphasia is present when the person actually has aphasia ($P[\text{hit}] = .70$ and $P[\text{miss}] = .30$). Y, under most traditional criteria, would be considered superior. However, if you were also told that Y identified 90% of the normal persons who came into his clinic as aphasic (i.e., $P[\text{FP}] = .9$ and $P[\text{TN}] = .1$) and Z had a FP rate of only 5% ($P[\text{FP}] = .05$ and $P[\text{TN}] = .95$), you might reconsider your evaluation of the adequacy of these individuals and be forced to conclude that Z was a better diagnostician than Y. Thus, the percent correct alone is an inadequate measure of accuracy. One must account for FPs as well as hits. As will be shown, TSD allows one to derive a better measure of accuracy based on hits plus FPs than percent hits alone.

TDS also considers a second importance area, the bias with which one approaches a task. In TSD, *bias* conventionally refers to the tendency to favor a positive outcome (hits). Both tests and individual clinicians have biases. Numerous factors affect these biases, including: (1) a clinician's background and training, (2) the assumptions a clinician or test (or both) brings to a given diagnostic situation, (3) a clinician's environment (e.g., work setting), (4) a clinician's mood, and/or (5) a clinician's cost-benefit assessment of a given situation.

Consider the following hypothetical diagnostic situation. General X is a military radar operator with the job of detecting incoming missiles. General X has family living in the area of his surveillance and wants to make sure that no missiles come close to populated areas, especially those where his family live. Because of his personal and professional situation he adopts a lenient response bias. That is, he wants to make sure that

when a missile is detected, he will not miss it. Thus, even though a defensive missile will be launched every time he detects an incoming missile, he launches his defensive missile any time he detects a signal on his radar screen. Because aircraft other than missiles are also detected on the screen, the signal detected may or may not be a missile. As a result, General X has a high FP rate, but never misses an incoming missile. For the General, the cost of missing an incoming missile is greater than the benefit of not taking down an allied or neutral plane. When General X goes off duty, Private A assumes the radar watch. Although the defense of the country from attack is paramount in both General X's and Private A's minds, Private A's family is scheduled to arrive by air some time during his work shift. He knows that every time he launches a defensive missile he is likely to hit his target because they are very accurate. He also reasons that the attack missiles are fairly inaccurate and the chance of hitting a populated area is low. Private A adopts a bias that dictates that he will launch a defensive missile only when he is absolutely positive that the signal detected on the radar screen is an offensive missile. His hit rate is lower than General X's, but so is his rate of FPs. Private A's bias dictates that the cost of a FP far outweighs the benefit derived from identifying every single incoming missile. The distributions of decisions based on N and SN form the data on which TSD is computed.

Figure 1 shows two distributions that represent N on the left and SN on the right. The distributions are similar in shape and their variance is equal. (Note that TSD can handle nonparametric data as well.) Notice that the two curves overlap to some degree. That is, in all situations there are instances where one may report SN when only N is present or only N when SN is present. One main index of accuracy, d' , is represented by the difference between the peaks of the N and SN distributions. The greater the overlap of N and SN distributions, the less the distance between the peaks, and the lower the d' . The distributions at the top of Figure 1 have the highest d' and represent the highest accuracy. The bottom distributions with the most overlap have the lowest d' and represent the poorest accuracy. The middle distributions represent some intermediary accuracy level. Accuracy as measured by d' can range from complete overlap of the distributions ($d' = 0.00$) to as high as about 4.67.

Figure 2 represents theoretical N (top) and SN (bottom) distributions. The perpendicular line represents the response bias which in TSD terms is called β . Responses to the right of β are positive for SN (clinician says yes, SN present) whereas those to the left favor N (clinician says no). At the intersection of the two distributions β is 1, which refers to equal bias for SN and N. As β decreases, a more lenient criterion is adopted (hits and FPs increase) and as β increases a more stringent criterion is adopted (hits and FPs decrease). Also note that the areas of the distribution that represent different cells in contingency Table 1 are shown in this figure.

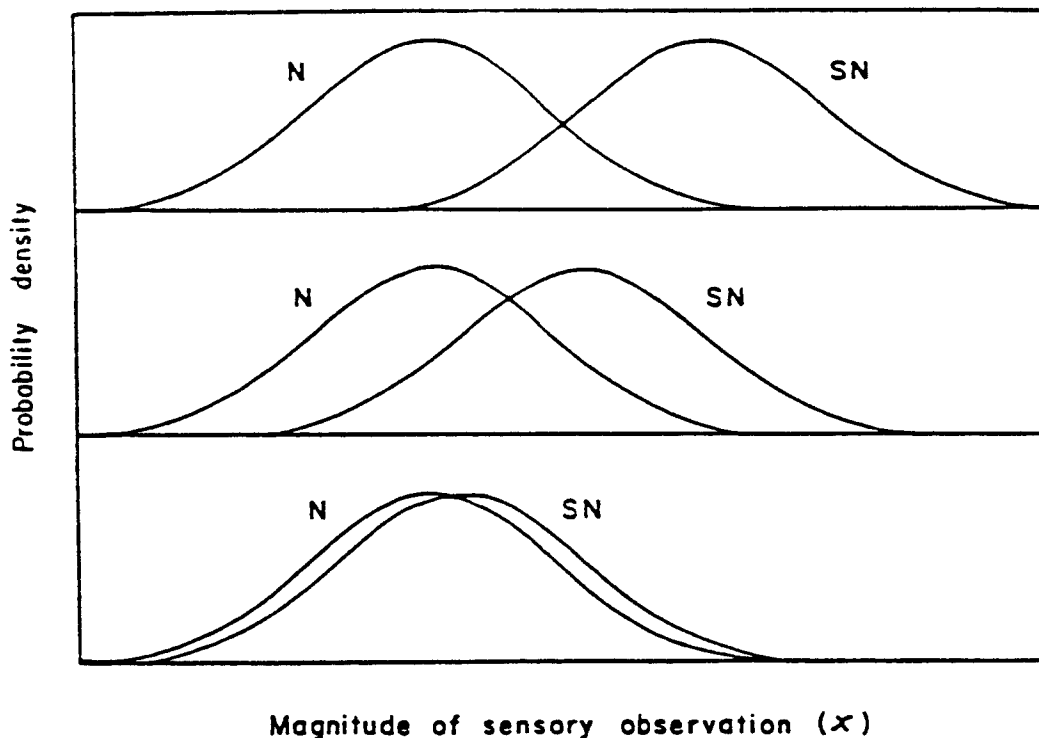


Figure 1. Three different theoretical distributions of noise (N) and signal-plus-noise (SN). Accuracy (d') is defined as the distance between the two peaks of the N and SN distributions. The further the distance between the two peaks, the better the accuracy of and the higher the d' . Thus, the top N and SN distributions represent the highest accuracy and d' is relatively large. The lowest N and SN distributions are almost overlapping and accuracy is poor, d' is close to 0. Reprinted from G. A. Gescheider, *Psychophysics: Method, theory and application* (2nd ed.), p. 87, with permission of Lawrence Erlbaum Associates, Hillsdale, New Jersey.

These probabilities depend on one's bias. Moreover, accuracy does not change as a function of shifts in bias, although the relative percent of hits and FPs do.

One other aspect of TSD needs to be discussed before the data are presented. In order to apply TSD to diagnostic situations, receiver operating characteristic (ROC) curves—a plot of the $P(\text{hits})$ by the $P(\text{FP})$ —need to be developed. The measure d' , reflects the distance between the N and SN distributions, which is reflected in the position of the ROC curve on the graph. Figure 3 is a graph of an ROC curve with a d' of 1.0. The diagonal straight line is a d' of zero, where the two distributions have complete overlap. The N and SN distributions that give rise to the position of the ROC curve are shown along with three different biases (β s)

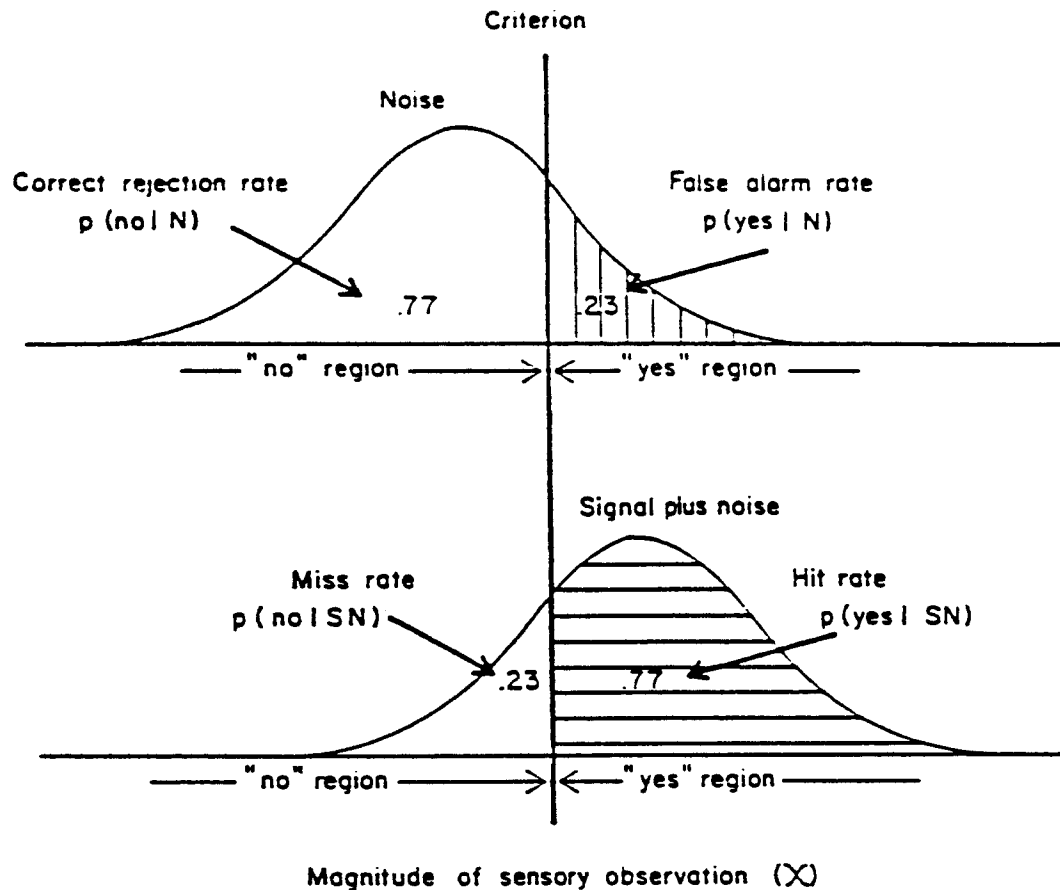


Figure 2. Theoretical distributions of noise (N) on top and signal-plus-noise (SN) bottom. Note that the distance between the peaks represents accuracy or d' . The perpendicular line represents bias (labeled as criterion in the figure) or β . For responses to the right of the line the clinician says yes, SN present and to the left of the line the clinician says no, SN not present (i.e., N present). Note the areas of overlap between the two distributions. Thus, all four conditions (Hit, FP, miss, and TN) are represented. Hits, (hit rate in figure [$P = .77$]) are shown by the lined portion of the SN distribution; FPs, false alarm in figure ($P = .23$), by the lined portion of the N distribution; misses, miss rate in figure ($P = .23$), by the blank portion of the SN distribution; and TN, correct rejection in figure ($P = .77$), by the blank portion of the N distribution. As the perpendicular line moves to the left a more lenient bias is adopted, β goes up. In this case both the hits and FPs increase. As the perpendicular line moves to the right, β goes down and a more stringent bias is adopted. In this case both hits and FPs decrease. Reprinted from G. A. Gescheider, *Psychophysics: Method, theory and application* (2nd ed.), p. 90, with permission of Lawrence Erlbaum Associates, Hillsdale, New Jersey.

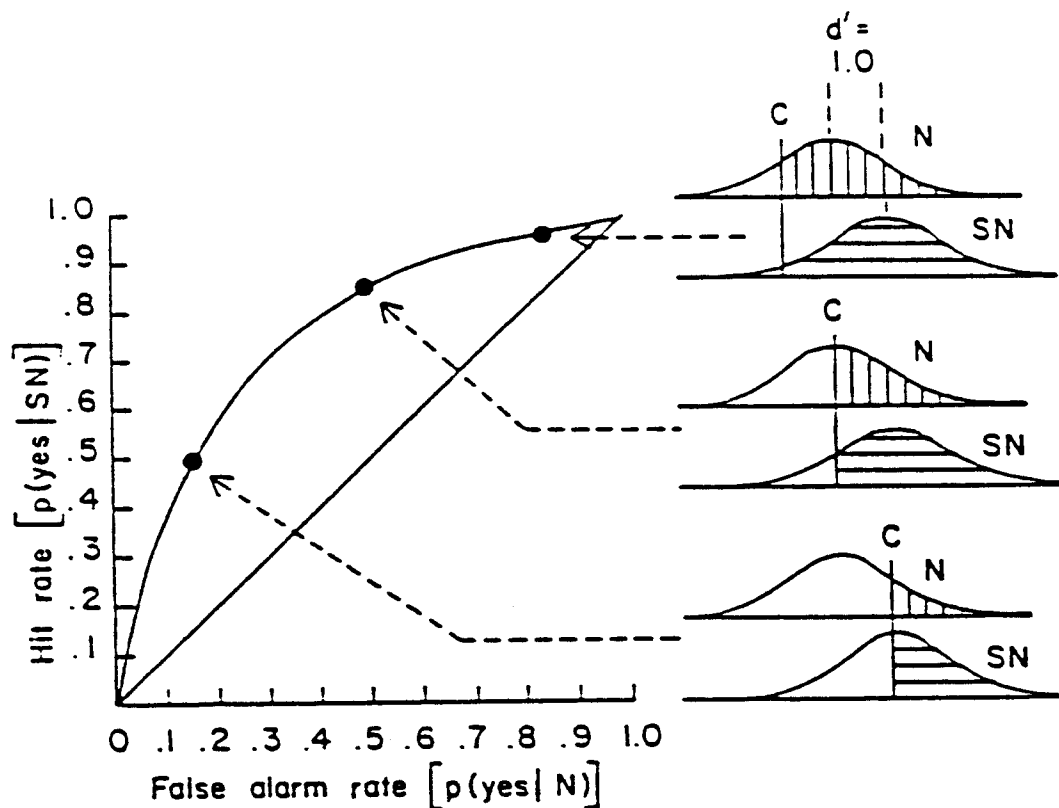


Figure 3. Relation between the ROC curve and the theoretical N and SN distributions that represent a d' of 1.0. The straight diagonal represents a d' of 0 as hits and FPs are equal. Each of the three data points represent a different bias at the same accuracy level (the distance between the peaks of the N and SN distributions is constant), only the perpendicular line moves. The most lenient bias, highest β is represented by the upper distributions for which both hit and FP rates are high. The most stringent bias, lowest β is represented by the bottom distributions in which both hit and FP rates are low. Reprinted from G. A. Gescheider, *Psychophysics: Method, theory and application* (2nd ed.), p. 95, with permission of Lawrence Erlbaum Associates, Hillsdale, New Jersey.

that lie along the curve. Different points along the curve represent differences in bias or β , with the same d' . Thus, d' is an accuracy measure independent of bias. The construction of ROC curves allows an assessment of accuracy *and* bias. The uppermost N and SN distributions have the bias line far to the left, thus indicating a low β and a lenient bias. Moving down the figure to the lower distributions, β increases, indicating more stringent biases. Figures 4 and 5 show ROC curves for d' 's of 2.0 and 0 respectively, each with three bias points.

If diagnostician's data are shown to fit the theoretical curves, then d' and β may be utilized without developing the entire curve. Convenient

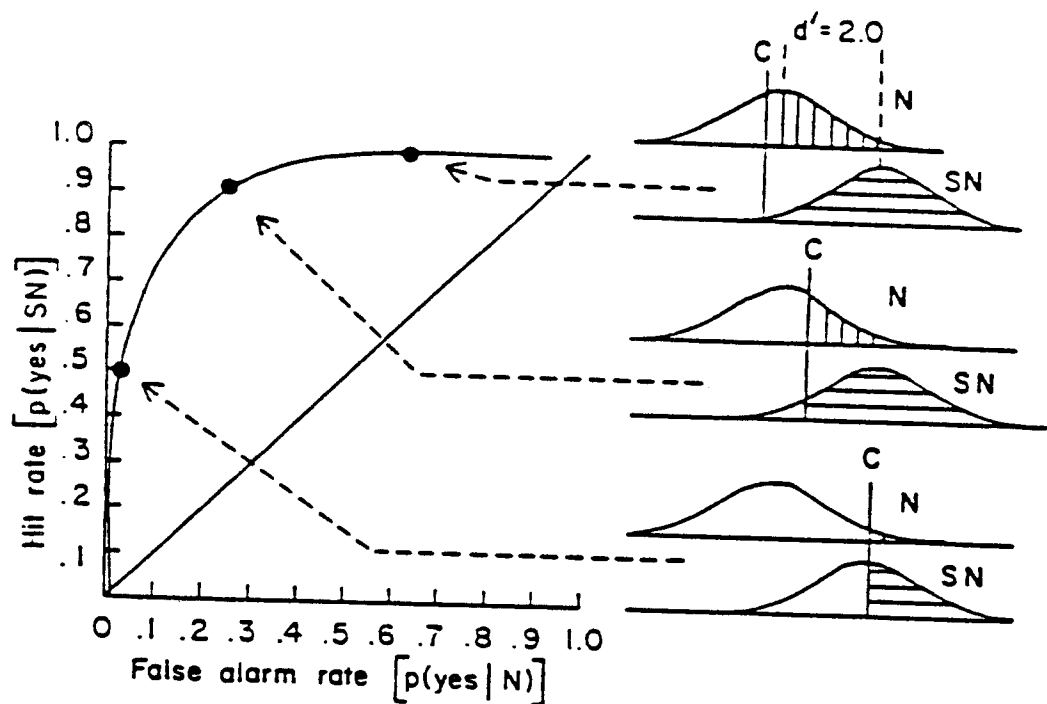


Figure 4. Same as Figure 3, but $d' = 2.0$. Reprinted from G. A. Gescheider, *Psychophysics: Method, theory and application* (2nd ed.), p. 94, with permission of Lawrence Erlbaum Associates, Hillsdale, New Jersey.

tables for d' and β exist for any known hit and FP rate. To illustrate TSD application to aphasia diagnostics and to explore its value, ROC curves, d' and β data for individual clinicians using the PICA to diagnose aphasia are presented.

METHOD

Judges

Four individual judges formally trained and experienced in the administration, scoring, and interpretation of the PICA, evaluated item, subtest, and overall scores from 246 different PICA tests. A fifth judge (subject 3) was a PICA-trained, but generally inexperienced graduate student seeking a master's degree in speech-language pathology.

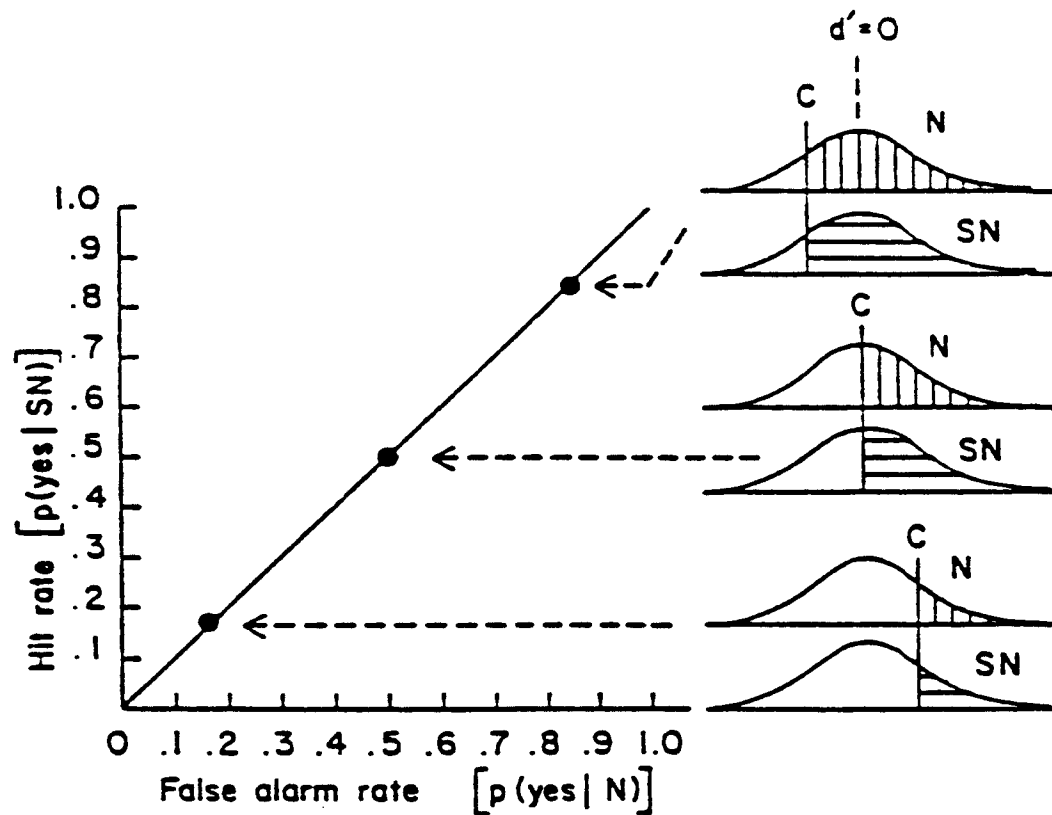


Figure 5. Same as Figure 3, but $d' = 0.0$. Reprinted from G. A. Gescheider, *Psychophysics: Method, theory and application* (2nd ed.), p. 96, with permission of Lawrence Erlbaum Associates, Hillsdale, New Jersey.

PICA Data

To determine the accuracy of responses, one needs to know, apriori, which PICA tests were derived from normal and which were from aphasic individuals. In other words, a *gold standard* for diagnosis is required. Although much theory and philosophy underlies the final categorization, only the general criteria and methods used for making the decisions are discussed. First, the normal subject's PICA performance was provided by Duffy and Keith from their 1980 standardization for normal individuals. Subjects met the criteria for normal as defined in that investigation. The aphasic PICA data were derived from the speech-language pathology patient records at the VA hospital in Madison, Wisconsin. The aphasia diagnosis was made or certified by either Wertz, Rosenbek, or Collins. Although the PICA was used in the patient's diagnosis, in no case was the categorization based solely on PICA data. A variety of other stan-

standardized, unstandardized and informal assessment tools, interviews, case histories, and medical records were also utilized to arrive at the final diagnosis. Aphasia was defined in a manner similar to Darley (1982). That is, all aphasic subjects had multimodality language deficits that were disproportionate to other intellectual deficits. Onset was sudden and secondary to a focal dominant hemisphere lesion. Of the 246 total *PICA* tests to be evaluated, 103 were from aphasic adults (SN distribution) and 143 were from normal non-brain-damaged individuals (N distribution).

Judges were first required to indicate whether the test presented was from an aphasic or normal individual (subjects said yes for aphasia and no for normal). The clinicians then rated the confidence of their decision on a five-point equal-appearing interval scale from 1 (*not very sure*) to 5 (*very sure*). The rating scale method is a cost effective means of applying TSD to clinical situations in order to generate the ROC curve and to test the validity of the approach (Swets, 1988; Swets & Pickett, 1982). Each point on the rating scale represents a different bias. An individual who is *very sure* of a decision will have a stringent bias (β will be high) while the bias associated with a rating of *not very sure* will be very lenient (β will be low). In this manner up to four points along an ROC curve, each with differing bias, were obtained. The maximum number of points is four as the most lenient criterion associated with the rating of 1 will always result in a $P(\text{hit})$ and $P(\text{FP})$ equaling 1.0.

From the rating scale data, the $P(\text{hit})$ and $P(\text{FP})$ for each person at the different biases were determined. ROC curves for each person were plotted and the data were compared to the theoretical curves. From these, accuracy and bias measures were derived. The d' and β for each person representing their typical levels were determined.¹

RESULTS AND DISCUSSION

Table 3 shows each subject's d' and β for the diagnostic decision (yes/no portion of the study). These data generally are thought to reflect the typical accuracy and bias of each judge. High accuracy is indicated by a d' better than 3.5. None of these subjects were close to achieving a high accuracy. The highest d' was 2.78. This judge, subject 5, had the least experience with aphasia and the *PICA* other than the student. Subject 4 with approximately 12 years of experience with the *PICA*, who uses it as a

1. The computer program to calculate the best fit for the data and determine accuracy and bias were kindly provided by Drs. Donald D. Dorfman and Kevin Burnbaum at the University of Iowa in the Departments of Psychology and Radiology, respectively. The original program can be found in Dorfman and Alf (1969).

Table 3. Accuracy (d') and Response Bias (β) for Each Subject at Typical Response Level

<i>SUBJ #</i>	d'	β
1	1.96	4.10
2	2.66	2.56
3	2.42	3.60
4	1.00	0.46
5	2.79	1.72

Note: Subject 3 was a student.

primary diagnostic tool, had a very low d' of 1.0. At this low level of accuracy, at typical bias, this judge had a P(hit) of .86 and a P(FP) of .62. Thus, this judge identified 86% of the aphasic tests as aphasic but also identified 63% of the normal tests as aphasic. By contrast, subject 5, who had the highest accuracy, had a P(hit) of .89 (89% correct aphasic identification) but a P(FP) of only .06 (6% incorrect identification of normal subjects) at their typical bias. The student clinician (subject 3) had a similar accuracy level to most of the other judges.

Table 3 also shows the bias with which the subjects typically approached the task. While β s were generally indicative of a relatively stringent bias (>1.0), there was a range of biases among the subjects. Subject 4 had a very lenient bias ($\beta = .46$) while Subject 1 was fairly stringent ($\beta = 4.10$). Thus, some individuals were more willing than others to identify an individual as having aphasia given the *PICA* information they reviewed.

Figure 6 shows the ROC curves for Subjects 4, triangles, and 5, circles. The dotted and solid lines represent the theoretical curves for a d' of 1.00 and 2.79, respectively. The diagonal line is a d' of zero. These data produce a good fit with the theoretical curves; in fact, each of the 5 subjects' data produced a good fit with the theoretical curves. It is interesting to note that Subject 4 reported using a relatively lenient response bias (typical $\beta = .46$) and that this bias caused the high degree of inaccuracy. Recall, however, that d' is independent of bias. At subject 4's most lenient bias ($\beta = .32$), d' was the same as at his more stringent biases ($\beta = .43, .46, \text{ or } .70$). Subject 5's data illustrate that one may adopt a stringent bias ($\beta = 6.78$) and have only 75% correct, or one may use a lenient bias ($\beta = .48$) and achieve a 96% hit rate but produce the same accuracy. Comparison of Subjects 4 and 5 at the similar bias ($\beta = .46$ and $.53$, respectively) was independent of their very different accuracy levels ($d' = 1.0$ and 2.78 , respectively). Thus, bias can be assessed independently of accuracy using TSD.

Figure 7 represents the theoretical ROC curves for each of the five subjects. Notice how the position of the ROC curve varies as a function of

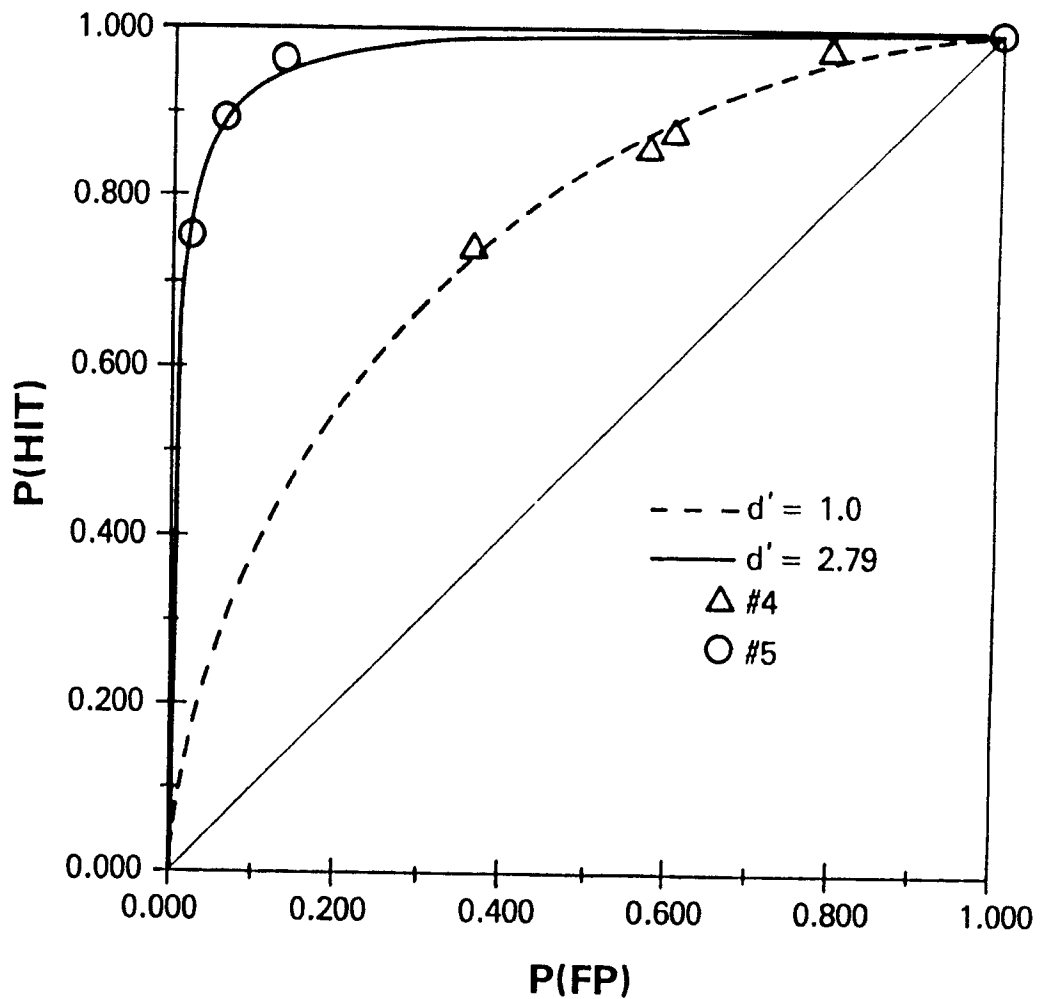


Figure 6. ROC curves for subjects #4 and #5 generated from *actual data* and *theoretical data*.

accuracy. As d' is lowered, the curve approaches the zero line. Different points along each curve represent differing biases. All subjects achieved a hit rate of 80%–90% using their typical bias. Their FPs varied, however, and thus accuracy differed among them.

In summary, TSD provided a method of investigating accuracy and bias in aphasia diagnosis. *PICA* tests alone did not result in acceptable accuracy levels given the extant literature suggesting that a d' greater than 3.5 is achieved in most good diagnostic systems. However, individual clinicians can use TSD to check their own bias and accuracy. Future studies could systematically evaluate accuracy and bias when differing amounts of information are available to the diagnostician. For instance, accuracy and bias could be determined when other standardized aphasia test data are added to those of the *PICA*. Accuracy and bias could then be

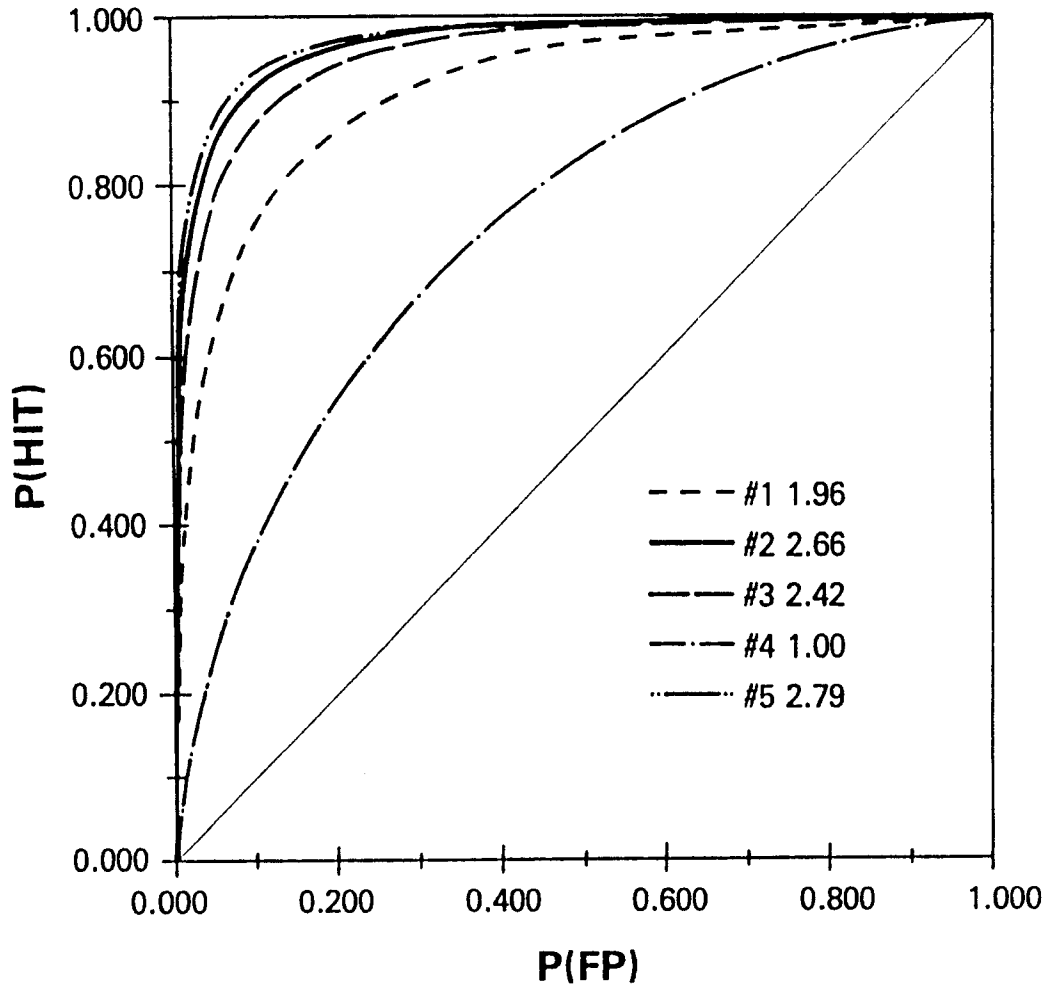


Figure 7. Best fit *theoretical* ROC curves for all subjects based on *actual* data.

determined when a case history or a brain scan is part of the information provided. One could vary *noise* by asking judges to determine if aphasia is present or if the patient is normal, has dementia, or has right hemisphere involvement. Accuracy and bias will vary as a function of the amount of information provided and the amount of noise in the system.

TSD is also useful in cost-benefit assessment. One can objectively determine which measures give the highest accuracy. The minimum amount of information that provides the highest accuracy measures can be determined for any given diagnostic decision.

This paper examined the basic principles of TSD and its application to aphasia diagnosis. Future studies will examine if the assumptions of d' (equal variance of N and SN) hold true. If not, other measures of accuracy may be more appropriate. Moreover, instead of using β as the bias measure, $\ln\beta$ (natural log of β) may prove to be a better metric as

lenient biases are indicated by β s between 0 and 1.0 and stringent biases by β greater than 1.0. The investigation of these and several related methodological issues in the application of TSD to aphasia assessment should assist the clinical aphasiologist in their selection of the most sensitive and specific tools and procedures for their clinical practice.

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