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Establishment of Quick Testing Efficiency Evaluation Model and Analysis of Related Factors

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Abstract

Objective: "Rapid screening, targeted sampling, objective test" is an efficient test model. The factors affecting the efficiency include false screening rate, missing rate and rapid screening time. However, only missing rate and accuracy have been used as technical requirements to evaluate rapid screening method. In this study, efficiency was regarded as evaluation index of quick testing method. Method: The evaluation model of quick testing efficiency was established by comparing time of routine testing and quick testing. By simulation calculation, the effect factors such as rapid screening time, false screening rate, missing rate and defective rate were analyzed. Results: The calculation formula of efficiency was derived. Simulation results showed that the lower defective rate, the higher efficiency; the smaller missing rate, false screening rate, or screening time, the higher efficiency and the degree of improving efficiency is related to defective rate; sometimes, the screening time is the most important factor affecting the efficiency. In certain cases, if the false screening rate or missing rate is 50%, the efficiency can be increased by more than 10 times. Conclusions: Taken together, this study highlighted a role of efficiency which functioned as an index to evaluate rapid screening. Quick testing efficiency evaluation model can be used for the calculation efficiency, and can be used to analysis the relationship between efficiency and the influence factors, and can provide the theoretical foundation for rapid screening method development and evaluation.

Keywords

Efficiency Evaluation, Rapid Screening Time, Missing Rate, False Screening Rate

1. Introduction

Food and drug market circulation is huge but regulatory supervision is limited. WHO put forward the concept of fighting against "substandard/spurious/falsely-labelled/falsi-

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zhfied/counterfeit medical products (SSFFCs)". "Rapid screening, targeted sampling, objective test" is an efficient test pattern [1] [2]. For example, in 2010, food and drug administration carried out special supervision and sampling inspection for health food that was added illegal chemicals. Rapid screening 3000 batches of samples and 165 batches of them turned out to be positive. But through objective test, there are 98 batches of sample proved positive. It took only two weeks. Compared to test 3000 batches and many projects, efficiency is improved obviously [3].

Rapid screening methods have been widely applied to food [4] [5], medicines [6] [7], health food [8] [9] and similar health products. But the food and drug safety situation is still not optimistic [10] [11] and rapid screening methods need further research and development.

Opinions about technical requirements vary from person to person. Guide for determination of health food and cosmetics rapid screening method printed by China Food and Drug Administration in 2012 [12] set that the false positives rate should be less than 20% and the correct screening rate should be more than 80%. Theories and Practices of Rapid Screening Technologies for Adulteration in Health Products written by Xie Zhijie suggested that the false positives rate should be less than 10%; missing rate should be less than 5%; and the correct screening rate should be more than 90% [3]. However, some simple methods can be used for screening to improve efficiency evidently, but do not fit technical requirements. For example, if correct screening rate is 60% (false screening rate is 40%) [3], efficiency is improved clearly. This study considers that improving the test efficiency is the purpose of rapid screening. Correct screening rate is just one factor that affects the efficiency because positive samples need to be confirmed.

Comparing with routine testing time and quick testing time, an evaluation model to calculate quick testing efficiency is established considering the impact of rapid screening time, false screening rate, missing rate and defective rate. By simulation calculation, the relationship between efficiency and affect factors is analyzed.

2. Method

In this paper, without calculating the economic cost of testing, we only consider the time to study technical efficiency. Comparing time of routine testing and quick testing, various factors that effect efficiency are analyzed by simulation calculation

2.1. Concept Definition

- 1) Rapid screening: It means a kind of rapid test for specific project to sample. The result could be true, but sometimes it could also be false positive or false negative.
- 2) Objective test: This method is to confirm the positive result of rapid screening through a testing method in law.
- Quick testing: It means an efficient testing pattern combining rapid screening and objective test. This pattern could be used to determine the positive samples quickly and accurately.
- 4) Number of unqualified products checked out is represented by P.

- 5) False positives number is represented by F_P .
- 6) False negative number is represented by L_p .
- 7) Routine test time for all items in the lab is represented by t_1 .
- 8) Rapid screening time is represented by t_2 .
- 9) Objective testing time for positive sample is represented by t_3 .
- 10) Defective rate is represented by a%.
- 11) False screening rate: false positive expressed as a percentage of screening positive

$$\frac{F_P}{P + F_P} \times 100\%$$

12) Missing rate: false negative expressed as a percentage of true positive

$$\frac{L_P}{P+L_P} \times 100\%$$

- 13) Efficiency value is represented by *E.*
- 14) Efficiency difference with sing-factor changed is represented by D-value.

2.2. Establish Efficiency Evaluation Model of Quick Testing

When product quantity is equal, work efficiency and time is inversely ratio. Efficiency can be calculated by the ratio of routine testing time to quick testing time.

- Quick testing time is the sum of rapid screening and laboratory verification time.
 False positive and false negative affect quick testing efficiency. When the same unqualified number of products is checked out, the relationship between testing time and unqualified number is in Table 1.
- 2) Formula derivation of efficiency

$$E = \frac{\frac{P}{a\%} \times t_1}{\frac{P + L_p}{a\%} \times t_2 + (P + F_p) \times t_3}$$
 (1)

$$= \frac{P \times t_1}{P \times t_2 + L_P \times t_2 + (P + F_P) \times t_3 \times a\%}$$
 (2)

$$= \frac{t_1}{t_2 + \frac{L_P}{P} \times t_2 + t_3 \times a\% + \frac{F_P}{P} \times t_3 \times a\%}$$
(3)

Table 1. The relationship between testing time and unqualified number.

Test type	Number of sample	Test time	Total test time
Routine testing	$\frac{P}{a\%}$	t_1	$\frac{P}{a\%} \times t_{\scriptscriptstyle 1}$
Rapid screening	$\frac{P+L_{_{\mathrm{P}}}}{a\%}$	t 2	$\frac{P+L_{_{\mathrm{P}}}}{a\%}\times t_{_{2}}$
Objective testing	$P + F_p$	t ₃	$(P+F_p)\times t_3$

2.3. Simulation

First step: A quick testing method efficiency is simulated by formula (3) under the condition of missing rate is 50%, false screening rate is 50%, screening time is 0.25 h with defective rate is 1%~50%. Second step: Three quick testing methods efficiency are simulated by single-factor changed. Third step: The effect of each factor on efficiency is analyzed by calculating efficiency D-value. Fourth step: Three factors are evaluated synthetically by comparing analysis.

1) Simulation of a quick testing method

 $t_1 = 10 \text{ h}$, $t_2 = 0.25 \text{ h}$, $t_3 = 2 \text{ h}$; $L_P/p = 1$ (missing rate is 50%); $F_P/p = 1$ (false screening rate is 50%), see **Table 2**.

2) Simulation of single-factor changes

 $t_1 = 10 \text{ h}$, $t_2 = 0.25 \text{ h}$, $t_3 = 2 \text{ h}$; $L_P/p = 0.25$ (missing rate is 20%); $F_P/p = 1$ (false screening rate is 50%), see **Table 3**.

 $t_1 = 10 \text{ h}$, $t_2 = 0.25 \text{ h}$, $t_3 = 2 \text{ h}$; $L_P/p = 1$ (missing rate is 50%); $F_P/p = 0.25$ (false screening rate is 20%), see **Table 4**.

 $t_1 = 10 \text{ h}$, $t_2 = 0.1 \text{ h}$, $t_3 = 2 \text{ h}$; $L_P/p = 1$ (missing rate is 50%); $F_P/p = 1$ (false screening rate is 50%), see **Table 5**.

3) Analysis of each single-factor

D-value with different missing rate, see **Table 6**.

D-value with different false screening rate, see **Table 7**.

D-value with different screening time, see **Table 8**.

Table 2. Efficiency evaluation forms (missing rate is 50%; false screening rate is 50%; $t_1 = 10 \text{ h}$; $t_2 = 0.25 \text{ h}$; $t_3 = 2 \text{ h}$).

a%	$L_{ m P}/P$	$F_{ m P}/P$	t_2	E
1%	1	1	0.25	18.52
10%	1	1	0.25	11.11
20%	1	1	0.25	7.69
30%	1	1	0.25	5.88
40%	1	1	0.25	4.76
50%	1	1	0.25	4.00

Table 3. Efficiency evaluation forms (missing rate is reduced).

a%	$L_{ m P}/P$	$F_{ m P}/P$	t ₂	E
1%	0.25	1	0.25	28.37
10%	0.25	1	0.25	14.04
20%	0.25	1	0.25	8.99
30%	0.25	1	0.25	6.61
40%	0.25	1	0.25	5.23
50%	0.25	1	0.25	4.32

Table 4. Efficiency evaluation forms (false testing rate is reduced).

a%	$L_{ m P}/P$	$F_{ m P}/P$	<i>t</i> ₂	E
1%	1	0.25	0.25	19.05
10%	1	0.25	0.25	13.33
20%	1	0.25	0.25	10.00
30%	1	0.25	0.25	8.00
40%	1	0.25	0.25	6.67
50%	1	0.25	0.25	5.71

Table 5. Efficiency evaluation form (screening time is reduced).

a%	$L_{\mathbb{P}}/P$	$F_{ m P}/P$	t ₂	Е
1%	1	1	0.1	41.67
10%	1	1	0.1	16.67
20%	1	1	0.1	10.00
30%	1	1	0.1	7.14
40%	1	1	0.1	5.56
50%	1	1	0.1	4.55

Table 6. The efficiency difference value when missing rate is reduced.

	a%					
	1%	10%	20%	30%	40%	50%
$E(L_P/P = 0.25)$	28.37	14.04	8.99	6.61	5.23	4.32
$\mathrm{E}\left(L_{\mathrm{P}}/P=1\right)$	18.52	11.11	7.69	5.88	4.76	4.00
D-value	9.85	2.93	1.30	0.73	0.47	0.32

Table 7. The efficiency difference value when false testing rate is reduced.

	a%					
-	1%	10%	20%	30%	40%	50%
$E(F_P/P = 0.25)$	19.05	13.33	10.00	8.00	6.67	5.71
$\mathrm{E}\left(F_{\mathrm{P}}/P=1\right)$	18.52	11.11	7.69	5.88	4.76	4.00
D-value	0.53	2.22	2.31	2.12	1.91	1.71

Table 8. The efficiency difference value when screening time is reduced.

	a%					
	1%	10%	20%	30%	40%	50%
E (t ₂ = 0.1 h)	41.67	16.67	10.00	7.14	5.56	4.55
E ($t_2 = 0.25 \text{ h}$)	18.52	11.11	7.69	5.88	4.76	4.00
D-value	23.15	5.56	2.31	1.26	0.80	0.55

4) Evaluation of the three effect factors

D-value of missing rate, false screening rate and screening time are comparison analyzed. Three factors are evaluated synthetically, see **Figure 1**.

3. Results and Analysis

3.1. Simulation Efficiency Value of a Quick Testing Method at Assumption Conditions

If $t_1 = 10$ h; $t_2 = 0.25$ h; $t_3 = 2$ h, missing rate is 50% and false screening rate is 50%. Results are shown as **Table 2**, efficiency value is from 18.52 to 4.00 with defective rate changes from 1% to 50%; the smaller defective rate, the higher efficiency. The efficiency is 11.11 at defective rate is 10%. It also means false screening rate or missing rate can be more than the technical requirements (20%).

3.2. Efficiency D-Value at Different Missing Rate Conditions

Comparing Table 2 and Table 3, if false screening rate is 50%, missing rate is 50% or 20%, the lower missing rate, the higher efficiency of the quick testing. Efficiency D-value is from 9.85 to 0.32 with defective rate changes from 1% to 50%, Efficiency D-value shows that the smaller the defective rate, the more obvious effect of missing rate on the quick testing efficiency (see Table 6).

3.3. Efficiency D-Value at Different False Screening Rate Conditions

Comparing **Table 2** and **Table 4**, if missing rate is 50%, false screening rate is 50% or 20%, the lower false screening rate, the higher efficiency of the quick testing. Efficiency D-value is 0.53, 2.22, 2.31, 2.12, 1.91, 1.71 with defective rate from 1% to 50%, efficiency D-value shows that effects of false screening rate on efficiency is more obvious when the defective rate is 20% (see **Table 7**).

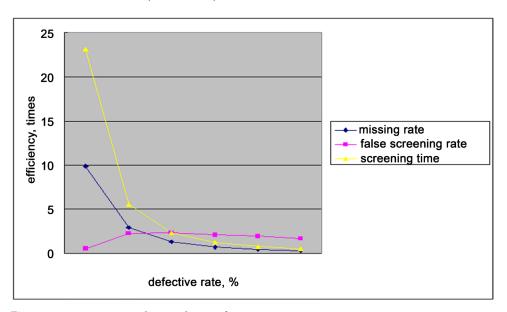


Figure 1. Comparative analysis evaluation form.

3.4. Efficiency D-value at Different Screening Time Conditions

Comparing **Table 2** and **Table 5**, if missing rate reduce is 50%; false screening rate is 50%, screening time is 0.25 h or 0.1 h, the shorter screening time, the higher efficiency. Efficiency D-value is from 23.15 to 0.55 with defective rate changes from 1% to 50%, results showed that the smaller the defective rate is, the more obvious effects of screening time on efficiency (see **Table 8**).

3.5. Comparison Analysis of the Three Effect Factors

Contrastive analysis of D-values in **Tables 6-8**, When defective rate < 20%, screening time is the most obvious impact on efficiency and the second is missing rate, false screening rate is last; when defective rate = 20%, the impact on the efficiency of screening time and false screening rate are equal and missing rate is last; when defective rate > 20%, the impact on the efficiency of the false screening rate is greater than the screening time and missing rate is last (see **Figure 1**).

4. Discussion

Although rapid screening also called quick testing in many references, in this study, quick testing means an efficient testing pattern combining rapid screening and objective test. This pattern could be used to quickly and accurately determine the positive samples.

This paper analyzes the influence of the false screening rate, missing rate, screening time under the condition of different defective rate on efficiency value. By changing the single factor respectively. The degree of improving efficiency is analyzed. The efficiency of quick testing is related to defective rate. The degree of improving efficiency is related to defective rate, too. Therefore, the defective rate is not only as a basis for the topic, but also affects the technical requirements of screening methods, so it's necessary to use big data to analyse defective rate. Although unqualified samples exist in the market unevenly sometimes. No matter how much the efficiency value is, and it can be improved certainly. The lower defective rate, the higher efficiency. The higher defective rate, the more necessary sampling inspection. For example, if the defective rate is too high, more than 50%, the efficiency of quick testing is too low and sampling inspection directly could be considered.

From an implementation standpoint, it is hard to reduce false testing rate and missing rate at the same time, reducing false testing rate usually means raising missing rate [3]. How to limit a reasonable range should take defective rate and the nature of the quick testing into consideration. For example, quick testing is applied to drug sampling inspection, sampling coverage should be increased, unqualified samples was found as far as possible; In terms of the Olympic food security, it is necessary to reduce the missing rate as far as possible; Access to agricultural products should be emphasized on the correct rate to reject unqualified samples and to ensure sales keep smoothly avoiding technical shortages.

5. Conclusions

The primary objective of the study was to establish a method for evaluating a quick testing method. The secondary aim of the study was to find the relationship between quick testing efficiency and impact factors such as missing rate, false testing rate, screening time and defective rate.

The efficiency is regarded as an evaluation index of quick testing method. The formula of calculating efficiency was deduced by comparing time of routine testing and quick testing.

Relationship between efficiency and impact factors has been analyzed by simulation calculation. Factors are changed separately. The influence of each factor on efficiency is analyzed by efficiency value. The degree of each factor's influence on the efficiency has been analyzed by efficiency D-value. At last, Comparison Analysis of the degree of each factor's influence on the efficiency has been completed.

The results provided that the lower missing rate or false screening rate, the higher efficiency of the quick testing; the shorter screening time, the higher efficiency. Difference values mean that the degree of improving efficiency is related to defective rate. Sometimes, the screening time is the most important factor affecting the efficiency. In certain cases, if the false screening rate or missing rate is 50%, the efficiency can be increased by more than 10 times.

The result of the study would enable policy and decision makers to pay more attention to efficiency; in addition, it would lead more rapid screening method to be developed. This would be one of the effective methods to ensure food and drug security.

The study can be used in food and drug; we can extend it to other fields such as disease diagnosis, environmental monitoring, etc.

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