Neonate Screening Test among Childbearing Mother Knowledge and Attitude

Ahmed Abudallh. Abud, PhD* Wassan Raheem, PhD**

ABSTRACT

Many biochemical tests are done through Newborn screening (NBS) in the first few hours or days of postpartum life. The main objective of these is the early identification of metabolic, genetic, or endocrine conditions in affected neonates. Prompt intervention can reduce morbidity, mortality, and associated disabilities that may appear later in life in undiagnosed and untreated cases so early detection is critical because. In comprehensive NBS programs biochemical testing conceder a key component of it, which includes examination and education, follow-up, diagnosis, treatment, management, and evaluation.

Aims of the study: This study was conducted to evaluate of Mothers' attitudes and knowledge for newborn screening in Thi-Qar.

Material and methods: The current study is a cross-sectional study in the city of Nasiriyah. Samples were collected from Muhammad Al-Mousawi Hospital and Bint Al-Huda Hospital in Nasiriyah city, about 100 samples during the period between 2 and 3 of 2022. The samples were collected randomly and their number was (100), distributed over the age group of mothers (18 - > 35 years).

Result: The result show that the agree answers were (4%), while the neutral answers were (77.8%) and the disagree answers were (18.2%). And, the true answers were (51.5%), while the false answers were (48.5%)

Conclusions: Antenatal examinations may increase knowledge and awareness of newborn screening but do not appear to influence the situation. It can be an effective way where health care providers can inform the mother about the process and purpose of this life-saving procedure. Better integration of government policy and information dissemination by health care providers should be a priority to improve mothers' understanding of newborn screening.

Keywords: Screening Test, Childbearing, Knowledge, Attitude

INTRODUCTION

Newborn screening is a public health service, which completes asymptomatic (newborn) testing so they can be identified and treated before problems occur, newborn screening is offered to all newborns and done soon after birth, the best age for testing is seventy-two hours to five days after birth, those not screened until two months of age aim for testing to distinguish a specific uncommon genetic, congenital, and metabolic disorder, but it is dangerous and may threaten life¹.

In Iraq, the program was implemented in April 2013, covering two governorates; Baghdad and Karbala, as Bidaya governorates, all newborns are offered screening for phenylketonuria (PKU), galactosemia, and congenital hypothyroidism (CHT)¹.

In many countries, phenylketonuria (PKU) was the first disorder for which NBS programs were initiated².

If mothers are aware and have knowledge of the purpose, process, and benefits of screening, they may act in response to the need for another test after positive (abnormal) results or an inadequate sample³.

Knowledge about newborn screening may also help reduce psychological and social harm; For example, informing mothers prior to testing that a first positive result is a possibility and does not indicate a confirmatory diagnosis, would likely reduce the risk of psychosocial harm associated with obtaining false-positive results from NBS⁴.

Though many factors influence mothers' opinions of newborn screening, it was created to encourage them to take its test. Positive attitudes are not necessarily dependent on adequate knowledge of screening, as there are substantial facts indicating that mothers generally have incomplete knowledge of the conditions being tested, the particular effects of the conditions, and accessible treatments⁵.

METHODOLOGY OF THE STUDY

The current study is a cross-sectional study in the city of Nasiriyah. Samples were collected from Muhammad Al-Mousawi Hospital and Bint Al-Huda Hospital in Nasiriyah city, about 100 samples during the period between 2 and 3 of 2022. Information regarding the age of the samples and details were obtained from the Hospitals Administration after obtaining approval. The samples were collected randomly and their number was (100), distributed over the age group of mothers (18 -> 35 years).

Focusing on the importance of testing and assessing high school students' self-concept.

* Department of pediatric College of Nursing University of Thi Qar, IRAQ E-mail: Ahmed-abud@utq.edu.iq

** College of Medicine

University of Thi Qar, IRAQ E-mail: wassan.raheem@utq.edu.iq The questionnaire consists of two parts:

Part 1: Demographic Data Consist Item Age, gender, grade, type of family, number of family members, father's testimony, mother's certificate, father's profession, mother's profession and monthly income. Date Collection Demographics were determined and developed by conducting face-to-face interviews with each student in the study using a questionnaire format. The data collection process took place from December 19, 2018 to January 9, 2019.

Each answer takes approximately 15-25 minutes to complete the questionnaire format

STUDY RESULTS

The distributions of the study sample by their sociodemographic characteristics:

Regarding socio-demographic characteristics table (5) indicated that the maternal age, estimated (26.3%) of their ages are (18-23) years, (26.3%) at the age (of 24-30) years, and (24.2%) at the age (31-35) years, the less age was frequent in this study was (>35 years) was (23.2%). This finding is supported by many studies like a cross-sectional descriptive study carried out in Baghdad, which indicated that a lot of mothers at age (24-30) years⁶.

As well as⁷ found that (40.1%) of mothers at age (25-30). While, Soliman et al., (2020)⁸ found that one-third of mothers at age (30-35) years. The result of the present study and other Iraqi and Arab studies was related to our culture in early age marriage and having children at an early age⁹.

Mother's education as shown in table (5) revealed that approximately thirty of the respondents, with school-level education, which disagrees with Rasheed et al., (2017)⁹ found that (35) others had university level of education. Alsuwat et al., (2018)¹⁰ found that about half of mothers had academic education¹¹.

The researcher's point of view is that the educational level of the mother is a very important issue in order to provide for the basic needs of the child particularly focusing on their diet and protective measures at home against many health problems which reflect the reduction in morbidity and mortality.

According to children number, 35.4% of mothers have 5-6 children.

The distribution of the mother's knowledge: Mother's knowledge about the test the table (6) revealed that mothers have a high level of knowledge about the test in all items the mean score (is 1.47). question 1 in table (6), (The test is a simple procedure for the discovery of genetic and congenital disease) was (54.5%) answered with True and (45.5%) answered with False. question 2 in table (6), (The test is important to ensure the infants' good health condition) was (45.5%) answered with True and (54.5%) answered with False. the question 3 in the table (6) , (The test helps in detecting metabolic disorders of an infant to avoid further deterioration of the child's health status) was (49.5%) answer with True and (50.5%) answered with False, the question 4 in the table (6), (When diagnosed early, there is a chance of an excellent prognosis and the newborn may be spared from lifelong impairment and can enjoy a normal life) was (43.4%) answer with True and (56.6%) answered with False, the question 5 in the table (6), (The test aids in the early detection of common genetic and congenital diseases, such as PKU, G6PD and CHT) was (57.6%) answer with True and (42.4%) answered with False, the question 6 in the table (6), (The test can identify more than 25 genetic and congenital diseases) was (58.6%) answer with True and (41.4%) answered with False the question 7 in the table (6), (The blood spot would help future research related to public health problems) was (46.5%) answer with True and (53.5%) answered with False the question 8 in the table (6), (The program is operated by the Ministry of Health) was (56.6%) answer with True and (43.4%) answered with False the question 9 in the table (6), (he test is performed by pricking the newborn's heel) was (63.6%) answer with True and (36.4%) answered with False the question 10 in the table (6)

Table 1: The demographic characteristics of the samples

Variables	Statistics	F	%
Age (Mean=2.44)	18-23	26	26.3%
	24-30	26	26.3%
	31-35	24	24.2%
	>30	23	23.2%
	1-2	25	25.3%
G 1	3-4	28	28.3%
Gender	5-6	35	35.4%
	>6	11	11.1%
	No Education(<1 years)	19	19.2%
	School Level (1-12years)	30	30.3%
Father Job	Mid-College Diploma (14years)	26	26.3%
	University Bachelor Degrees'	21	21.2%
	Master's Degree or Higher	3	3.0%
	Glucose 6 phosphate dehydrogenase (G6PD)	18	18.2%
	Phenylketonuria (PKU)	24	24.2%
Mather Job	Congenital hypothyroidism (CHT)	18	18.2%
	Congenital hypothyroidism (CHT)	39	39.4 %
Housing Environment	No Health Issues	44	44.4%
	Carrier of Congenital or Heredity Disease	13	13.1%
	Have Congenital or Heredity Disease	23	23.2%
	Dead Because of Congenital or Heredity Disease	19	19.2%

Table 2: Descriptive Statistics for Knowledge N=99, T=True, F=False

The test is a simple procedure for the discovery of genetic and congenital disease The test is important to ensure the infants' good health. condition The test is important to ensure the infants' good health. condition The test helps in detecting metabolic disorders of an infant to avoid further deterioration of the child's health status The test helps in detecting metabolic disorders of an infant to avoid further deterioration of the child's health status The test helps in detecting metabolic disorders of an infant to avoid further deterioration of the child's health status The disapposed early, there is a chance of an excellent prognosis and the newborn may be spared from lifelong impairment and can enjoy a normal life of the test aids in the early detection of common genetic and congenital diseases, such as PKU, G6PD and CHT The test can identify more than 25 genetic and congenital diseases The test can identify more than 25 genetic and congenital diseases The blood spot would help future research related to public health problems of the definition of the child's definition of the
The test is important to ensure the infants' good health. condition The test helps in detecting metabolic disorders of an infant to avoid further deterioration of the child's health status When diagnosed early, there is a chance of an excellent prognosis and the newborn may be spared from lifelong impairment and can enjoy a normal life The test aids in the early detection of common genetic and congenital diseases, such as PKU, G6PD and CHT The test can identify more than 25 genetic and congenital diseases The blood spot would help future research related to public health problems The program is operated by the Ministry of Health True
The test is important to ensure the infants' good health, condition The test helps in detecting metabolic disorders of an infant to avoid further deterioration of the child's health status When diagnosed early, there is a chance of an excellent prognosis and the newborn may be spared from lifelong impairment and can enjoy a normal life The test aids in the early detection of common genetic and congenital diseases, such as PKU, G6PD and CHT The test can identify more than 25 genetic and congenital diseases The blood spot would help future research related to public health problems The program is operated by the Ministry of Health True True 1.54 5.45 5.45 5.45 1.54 0.50046 False True T 49 49.5 F 50 50.5 1.50 0.49819 False True True The program is operated by the Ministry of Health True
The test helps in detecting metabolic disorders of an infant to avoid further deterioration of the child's health status When diagnosed early, there is a chance of an excellent prognosis and the newborn may be spared from lifelong impairment and can enjoy a normal life The test aids in the early detection of common genetic and congenital diseases, such as PKU, G6PD and CHT The test can identify more than 25 genetic and congenital diseases The test can identify more than 25 genetic and congenital diseases The blood spot would help future research related to public health problems The program is operated by the Ministry of Health True
deterioration of the child's health statusF5050.51.50 0.50252 TrueWhen diagnosed early, there is a chance of an excellent prognosis and the newborn may be spared from lifelong impairment and can enjoy a normal lifeT4343.41.560.49819FalseThe test aids in the early detection of common genetic and congenital diseases, such as PKU, G6PD and CHTT5757.61.420.49674TrueThe test can identify more than 25 genetic and congenital diseasesT5858.61.410.49508TrueThe blood spot would help future research related to public health problemsT4646.51.530.50129FalseThe program is operated by the Ministry of HealthT5656.61.430.49819True
When diagnosed early, there is a chance of an excellent prognosis and the newborn may be spared from lifelong impairment and can enjoy a normal life The test aids in the early detection of common genetic and congenital diseases, such as PKU, G6PD and CHT The test can identify more than 25 genetic and congenital diseases The test can identify more than 25 genetic and congenital diseases The blood spot would help future research related to public health problems The program is operated by the Ministry of Health True
newborn may be spared from lifelong impairment and can enjoy a normal life The test aids in the early detection of common genetic and congenital diseases, such as PKU, G6PD and CHT The test can identify more than 25 genetic and congenital diseases T 58 58.6 F 41 41.4 True The blood spot would help future research related to public health problems T 46 46.5 F 53 53.5 The program is operated by the Ministry of Health True
The test can identify more than 25 genetic and congenital diseases The blood spot would help future research related to public health problems To 56 56.6 The test aids in the early detection of common genetic and congenital diseases To 57 57.6 F 42 42.4 1.42 0.49674 True True The blood spot would help future research related to public health problems To 58 58.6 F 41 41.4 The program is operated by the Ministry of Health True True True
The test can identify more than 25 genetic and congenital diseases $\begin{array}{c ccccccccccccccccccccccccccccccccccc$
The test can identify more than 25 genetic and congenital diseases $\frac{\mathbf{T}}{\mathbf{F}} = \frac{42}{41} = \frac{42.4}{41.4}$ The test can identify more than 25 genetic and congenital diseases $\frac{\mathbf{T}}{\mathbf{F}} = \frac{58}{41} = \frac{58.6}{41.4} = 1.41 = 0.49508 \qquad \text{True}$ The blood spot would help future research related to public health problems $\frac{\mathbf{T}}{\mathbf{F}} = \frac{46}{53} = \frac{46.5}{53.5} = 1.53 = 0.50129 \qquad \text{False}$ The program is operated by the Ministry of Health $\frac{\mathbf{T}}{\mathbf{F}} = \frac{56}{56.6} = \frac{56.6}{1.43} = 0.49819 \qquad \text{True}$
The blood spot would help future research related to public health problems $\begin{array}{c ccccccccccccccccccccccccccccccccccc$
The blood spot would help future research related to public health problems $\begin{array}{c ccccccccccccccccccccccccccccccccccc$
The program is operated by the Ministry of Health $\frac{\mathbf{F}}{\mathbf{F}}$ 53 53.5 1.53 0.50129 False
The program is operated by the Ministry of Health T 56 56.6 1.43 0.40819 True
The program is operated by the Ministry of Health
F 43 43 4
The test is performed by pricking the newborn's heel $\frac{T}{F} = \frac{63}{26} = \frac{63.6}{264} = 1.36 = 0.48349$ True
r 30 30.4
The best time to do the test for a newborn is between three to seven days $\frac{T}{F} = \frac{59}{40} = \frac{59.6}{40.4} = 1.53$ 0.50129 False
F 40 40.4
The heel-pricking is performed by a nurse, midwife, or doctor $\frac{T}{F} = \frac{55}{44} = \frac{55.6}{44.4} = 1.40 \qquad 0.49320 \qquad \text{True}$
r 44 44.4
The blood samples will be sent to the Newborn T 46 46.5
Screening Laboratory canter and the result will be released in 7–14 working F 53 53.5 1.44 0.49943 True
days
If the results of the heel-prick test are abnormal, it means that the newborn \underline{T} 39 39.4
has the risk of developing a F 60 60.6 1.53 0.50129 False
genetic or congenital disorder
Weighting mean 1.4794 True
St. Deviation 0.14530

Table 3: Descriptive statistics for attitudes N=99, T=True, F=False

Questions		N	%	Mean	Deviation	Assessment
	Agree	37	37.4		0.80927	Neutral
Useful if it prevents a disease	Neutral	34	34.3	1.9091		
	Disagree	28	28.3			
	Agree	21	21.2		0.80121	Neutral
Useful if it reduces the severity of a disease	Neutral	27	27.3	2.3030		
	Disagree	51	51.5			
Useful even if it cannot improve the disease as it may help	Agree	16	16.2			
me decide about mothering children in the future	Neutral	41	41.4	2.2626	0.72261	Neutral
the decide about mothering children in the future	Disagree	42	42.4			
	Agree	35	35.4			
Is beneficial to the newborn	Neutral	32	32.3	1.9697	0.82628	Neutral
	Disagree	32	32.3			
	Agree	25	25.3		0.82040	Neutral
Is harmful to the newborn	Neutral	29	29.3	2.2020		
	Disagree	45	45.5			
	Agree	24	24.2		0.75716	Neutral
Is essential for the well-being of the newborn	Neutral	42	42.4	2.0909		
	Disagree	33	33.3			
	Agree	26	26.3	2.1313	0.80365	Neutral
Is morally justified	Neutral	34	34.3			
	Disagree	39	39.4			
	Agree	22	22.2		0.76541	Neutral
Is against my religious belief	Neutral	39	39.4	2.1616		
	Disagree	38	38.4			
Would make me feel guilty if the newborn is found to have	Agree	19	19.2			
a genetic disease	Neutral	24	24.2	2.3737	0.79006	Disagree
a genetic disease	Disagree	56	56.6			
Weighting mean			2.1560		— Neutral	
St. Deviation			0.29946		Neutrai	

Table 4: The descriptive statistics for overall knowledge and attitudes

Table Overall Knowledge			
Statistics	N	%	
True	51	51.5	
False	48	48.5	
False Total	99	100.0	

Table 5: Demographic characteristics with overall knowledge

Variables	Statistics	Chi secure	P. value	Sig.
Age (Mean=2.44)	18-23			Significant
	24-30	— —61.238ª	0.028	
	31-35	-01.238		
	>30			
	1-2		0.322	Insignificant
Gender	3-4	— —45.674ª		
Gender	5-6	43.074		
	>6			
	No Education(<1 years)		0.482	Insignificant
	School Level (1-12years)			
Father Job	Mid-College Diploma (14years)	— —55.821ª		
ramer job	University Bachelor	- 33.821		
	Degrees'			
	Master's Degree or Higher			
	Glucose 6 phosphate dehydrogenase		0.619	Insignificant
	(G6PD)			
Mather Job	Phenylketonuria (PKU)	38.645a		
	Congenital hypothyroidism (CHT)			
	Congenital hypothyroidism (CHT)			
	No Health Issues		0.759	Insignificant
	Carrier of Congenital or Heredity Disease			
Housing Environment	Have Congenital or Heredity Disease	35.271a		
	Dead Because of			
	Congenital or Heredity Disease			

Table 6: Demographic characteristics with overall attitudes

Table Overall Attitudes					
Statistics	N	9/0			
Agree	4	4.0			
Neutral	77	77.8			
Disagree	18	18.2			
Total	99	100.0			

Table 7: Demographic characteristics with overall knowledge

Variables	Statistics	Chi secure	P. value	Sig.
Age (Mean=2.44)	18-23			Insignificant
	24-30	— —33.767a	0.290	
	31-35	-33.767a		
	>30	_		
	1-2		0.317	Insignificant
C 1	3-4	22 120-		
Gender	5-6	−33.129a		
	>6			
	No Education(<1 years)		0.781	Insignificant
	School Level (1-12years)			
Father Job	Mid-College Diploma (14years)	22.952-		
rainer Job	University Bachelor	−32.853a		
	Degrees'			
	Master's Degree or Higher			
	Glucose 6 phosphate dehydrogenase		0.985	Insignificant
	(G6PD)			
Mather Job	Phenylketonuria (PKU)	15.695a		
	Congenital hypothyroidism (CHT)			
	Congenital hypothyroidism (CHT)			
	No Health Issues		0.565	Insignificant
	Carrier of Congenital or Heredity Disease			
Housing Environment	Have Congenital or Heredity Disease	28.095a		
	Dead Because of			
	Congenital or Heredity Disease			

This chapter show reasonably determined clarification and systematic discussion provided with reassuring confirmation available in the literature and articles.

, (The best time to do the test for a newborn is between three to seven days) was (59.6%) answer with True and (40,4%) answered with False the question 11 in the table (6) , (The heel-pricking is performed by a nurse, midwife, or doctor) was (55.6%) answer with True and (44.4%) answered with False the question 12 in the table (6) , (The blood samples will be sent to the Newborn Screening Laboratory center and the result will be released in 7–14 working days) was (46.5%) answer with True and (53.5%) answered with False the question 13 in the table (6) , (If the results of the heel-prick test are abnormal, it means that the newborn has the risk of developing a genetic or congenital disorder) was (39.4%) answer with True and (60.6%) answered with false.

The distributions of the mother's attitudes: The mother's attitudes about the test in the table (7) revealed that mothers have a high level of knowledge about the test in all items the mean score (2.15). question 1 in the table (7), (Useful if it prevents a disease) was (37.4%) answered with Agree and (34.3%) answered with neutral, while (28.3%) answered with Disagree. question 2 in table (7), (Useful if it reduces the severity of disease) was (21.2%) answered with Agree and (27.3%) answered with neutral, while (51.5%) answered with disagreeing. the question 3 in table (7), (Useful even if it cannot improve the disease as it may help me decide about mothering children in the future) (16.2%) answered with Agree and (41.4%) answered with neutral, while (42.4%) answered with disagreeing. question 4 in table (7), (Is beneficial to the newborn) was (35.4%) answered with Agree and (32.3%) answered with Neutral, while (32.3%) answered with disagreeing. question 5 in table (7), (Is harmful to the newborn) was (25.3%) answered with Agree and (29.3%) answered with neutral, while (45.5%) answered with disagreeing. question 6 in table (7), (Is essential for the well-being of the newborn) was (24.4%) answered with Agree and (42.4%) answered with neutral, while (33.3%) answered with disagreeing. question 7 in table (7), (Is morally justified) was (26.3%) answered with Agree and (34.3%) answered with neutral, while (39.4%) answered with disagreeing. question 8 in table (7), (Is against my religious belief) was (22.2%) answered with Agree and (39.4%) answered with neutral, while (38.4%) answered with disagreeing. question 9 in table (7), (Would make me feel guilty if the newborn is found to have a genetic disease) was (19.2%) answered agreeing and (24.2%) answered neutral, while (56.6%) answered disagree.

The distributions of the overall knowledge: According to table (8), the true answers were (51.5%), while the false answers were (48.5%)

The distributions of the overall attitudes

According to the table (9) the agree answers were (4%), while the neutral answers were (77.8%) and the disagree answers were (18.2%).

DISCUSSION

Antenatal examinations may increase knowledge and awareness of newborn screening but do not appear to influence the situation. It can be an effective way where health care providers can inform the mother about the process and purpose of this life-saving procedure. Better integration of government policy and information dissemination by health care providers should be a priority to improve mothers' understanding of newborn screening. For future studies, larger sample size is recommended, given the weak statistical power of the analyses. The survey can also be conducted in different hospitals to see the impact of location and type of hospital on the results. PGH deals with complex cases; Thus, mothers on PGH are more likely to consult for prenatal screening and therefore be better acquainted with newborn screening. The retention time of mothers acquainted with NBS may also be a factor in general understanding and may also be assessed.

RECOMMENDATIONS

Recognize and take advantage of the positive aspects of personality. Recognize your personal strengths to be able to excel. Attention to appearance.

Building new friends, mixing into social life, and engaging in various interesting discussions, all of these steps strengthen selfconfidence and reinforce new ideas.

Exercise different activities and hobbies that improve a person's skills, improve his health, eliminate negative energy in the body, and enhance self-confidence.

Every new experience is always to kill the fear within the person, and to enhance the ability to make a decision.

Authorship Contribution: Mei-Ling Huang: Conceptualization, Methodology, Writing- Original draft preparation, Investigation, Supervision, Writing-Reviewing and Editing, Funding acquisition. Ting-Yu Lin: Software, Formal Analysis, Data curation

Potential Conflict of Interest: None

Competing Interest: None

Acceptance Date: 16 July 2022

REFERENCES

- Padilla CD. Newborn screening in the Philippines. Southeast Asian J Trop Med Public Health 2003;34(3):87-8.
- Davey A, French D, Dawkins H, et al. New mothers' awareness of newborn screening, and their attitudes to the retention and use of screening samples for research purposes. Life Sci Soc Policy 2005;1(3):41.
- 3. Parva NR, Tadepalli S, Singh P, et al. Prevalence of Vitamin D Deficiency and Associated Risk Factors in the US Population. Cureus 2018;10(6):e2741.
- 4. Business Dictionary Attitude. 2019. http://www.businessdictionary.com/definition/attitude.html

- Biber K. Views of Expectant Mothers About Sources of Knowledge and Skills Related to Motherhood. Eur J Educ 2016;2(11):30-46.
- 6. Therrell BL, Padilla CD, Loeber JG, et al. Current status of newborn screening worldwide: 2015. Semin Perinatol 2015;39(3):171-87.
- Padilla CD, Aguirre TB. Enhancing case detection of selected inherited disorders through expanded newborn screening in the Philippines. Acta Med Philipp 2012;46(4):24-9.
- 8. Soliman NS, Wahdan MM, Abouelezz NF, et al. Knowledge, Attitude and Practice towards Vitamin D Importance and Supplementation among Mothers of under Five Children in a Primary Health Care Center. Int J Med 2020;113(1).
- Rasheed TAW, Taha HK, Rasheed BA. Knowledge, attitude and practice of Iraqimothers towards Vitamin D supplementation to their infants in Baghdad Al -Rusafa 2016. Al-Kindy College Med J 2017;13(2):111-6.
- 10. Alsuwat FS, Alzahrani NJ. Knowledge and Attitude Practice (KAP) of Rickets Disease among Mothers in KSA. Egypt J Hosp Med 2018;72(6):4582-5.
- WHO. Early child development. 2019. https://www.who.int/ topics/early-child-development/en/
- 12. Kamel WK, El-Hamid HSA, El-Megeed HA, et al. Mothers' Awareness regarding Vitamin D Deficiencyamong Their Infantsin Kalyobia Governorate. Menoufia Nursing J 2017;2(1):1-16
- Ministry of health (MOH) and USAID National Guideline of Newborn Screening for Care Providers in primary health care centers in Iraq. 2014;1:7.
- 14. Metz MP, Ranieri E, Gerace RL, et al. Newborn screening in south Australia: universal? Med J Aust 2003;179(8):412-5.
- 15. kemper AR, Fant KE, Little J, et al. Informing parents about newborn screening Public Health Nurs 2005;22(4):332-8.
- 16. Pollitt R. Neonatal screening for inbornerrors of metabolism: cost, yield and outcome. Health Technal Assess 2005;1(3):41-51.
- 17. Gurian EA, kinnamon DD, HenryJJ, et al. Expanded newborn screening for biochemical disorders: the effect of false positive result. Pediatrics 2006;117(6):1915-21.
- 18. Nicholls SG, Tessier L, Etchegary H, et al. Stakeholder attitudes towards the role and application of informed consent for newborn bloodspot screening: a study protocol BMJ Open 2014;4(11):e006782.