

# COMPARATIVE STUDY OF SINGLE SURFACE AND DOUBLE SURFACE PHOTOTHERAPY IN NEONATAL HYPERBILIRUBINEMIA

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**Abstract-** Neonatal jaundice occurs in about 60% of full-term neonates and in about 80% of preterm neonates. A significant proportion of causes of hyperbilirubinemia in the term newborn are benign and reversible. However, considering the potentially irreversible toxicity of bilirubin on the central nervous system (kernicterus), newborns must be evaluated to identify which ones will need treatment. It is a result of the increased breakdown of RBC and/or decreased hepatic excretion of bilirubin. Generally NNH is a natural transition that resolves within the 1st week of life with liver maturity. Since the 1950s, phototherapy has been the therapy of choice for the newborn with indirect hyperbilirubinemia. Single phototherapy (SP) is the most commonly used method, and, when bilirubin levels are close to the threshold for exchange transfusion, intensive phototherapy is indicated<sup>6</sup>. To study and compare the rate of fall of Total Serum Bilirubin levels in neonatal hyperbilirubinemia treated with Single Surface Phototherapy and Double Surface Phototherapy. Neonates in NICU and in Postnatal wards, who are clinically icteric, will be enrolled in the study after written informed consent from the parents (Annexure 1). Data will be collected in a pre-decided proforma (Annexure 2). Details will be obtained from direct questioning of the mother and examination of the newborn and his record. Repeat bilirubin levels will be estimated as per the protocols and requirements of the NICU. Phototherapy will be stopped in accordance with the protocols followed in NICU. The rate of fall of serum bilirubin levels will be observed in both the types of phototherapy given to determine if there is any difference in the effectiveness of SSPT vs DSPT. DSPT is more effective than SSPT. Conclusions of the study will be based on the results of the study.

**Keywords:** neonatal hyperbilirubinemia, hyperbilirubinemia, single surface phototherapy, double surface phototherapy, NICU admissions, comparative study, rate of fall of serum bilirubin, hyperbilirubinemia, single surface phototherapy, double surface phototherapy, NICU admissions, comparative study.

## I. Introduction

Neonatal jaundice occurs in about 60% of full-term neonates and in about 80% of preterm neonates<sup>1</sup>. Bilirubin is derived from the breakdown of heme containing proteins in the reticuloendothelial system. The daily production

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of bilirubin in a normal newborn is 6 to 10 mg/kg/day. A significant proportion of causes of hyperbilirubinemia in the term newborn are benign and reversible<sup>2</sup>. However, considering the potentially irreversible toxicity of bilirubin on the central nervous system (kernicterus), newborns must be evaluated to identify which ones will need treatment<sup>3</sup>. It is a result of the increased breakdown of RBC and/or decreased hepatic excretion of bilirubin. Generally NNH is a natural transition that resolves within the 1st week of life with liver maturity. Since the 1950s, phototherapy has been the therapy of choice for the newborn with indirect hyperbilirubinemia<sup>4,5</sup>. Single phototherapy (SP) is the most commonly used method, and, when bilirubin levels are close to the threshold for exchange transfusion, intensive phototherapy is indicated<sup>6</sup>

The most common therapeutic modalities in the treatment of NNH are:

1. Phototherapy
2. Exchange transfusion
3. Intravenous immune globulin.
4. Bilirubin molecules in light-exposed skin undergo quick photochemical reactions:
5. configurational isomerization,
6. structural isomerization,
7. photooxidation

The size of exposed BSA + level of irradiance, determines the **spectral power** of phototherapy, which in turn influences its effectiveness. The larger the exposed BSA and stronger the light, the higher the spectral power.

**Irradiance:** Minimum level of 30microW/cm<sup>2</sup>/nm in wavelength range of 460 to 490nm must be ensured

The possible adverse effects associated with phototherapy are: skin rash, increased insensible losses, retinal damage, hyperthermia, and deposition alterations due to increased intestinal flow<sup>7,8</sup>

**Background/rationale:** Proper studies were not available to compare the effectiveness of SSPT and DSPT in NNH.

**Objectives:** To study and compare the rate of fall of Total Serum Bilirubin levels in neonatal hyperbilirubinemia treated with SSPT and DSPT.

**Methods:**

**Study design:** Prospective observational study of neonates admitted in NICU with hyperbilirubinemia.

**Setting:** The AVBRH, Sawangi is a rural Medical College located in Maharashtra. This study will be conducted in the NICU and Postnatal ward, Department of PEDIATRICS in Jawaharlal Nehru Medical College & Acharya Vinoba Bhave Rural Hospital, Wardha for a period of 2 years.

**Participants**

**Inclusion Criteri**

- All neonates admitted in NICU for treatment of neonatal hyperbilirubinemia requiring phototherapy.

**Exclusion Criteria**

- Parents denied consent.
- Neonates with hyperbilirubinemia in exchange transfusion range.

**Variables:** gestational age of the child(term/,gender, risk factors, co morbidities,

Data sources/ measurement- To **study** the rate of fall of Total Serum Bilirubin levels in NNH treated with SSPT and DSPT

- To **compare** the rate of fall of TSB levels in NNH treated with SSPT and DSPT

**Study size:** Population size (for finite population)

Sample size(n):**73**

**Quantitative variables:** Rate of fall of TSB levels in NNH treated with SSPT and DSPT.

**Statistical methods:** The data will be entered in Microsoft Excel sheet. The quantitative data analysis will be done using Student t-test and qualitative data will be analysed using Chi-square test. The p-value of <0.5 will be considered significant.

### **Expected Outcomes/Results**

DSPT is more effective than SSPT is expected according to previous similar studies. All neonates admitted in NICU for treatment of neonatal hyperbilirubinemia requiring phototherapy. Irrespective of gestational age, gender, co morbid conditions, risk factors of mother and baby, all neonates with hyperbilirubinemia will be taken into the study.

## **II. Discussion**

Many studies have demonstrated significant reduction of bilirubin level by using double phototherapy compared with single phototherapy, but the irradiance of phototherapy is lower than recommendation of AAP, and some trials studied in low birth weight infants<sup>9,10</sup>. The efficacy of phototherapy is dependent on the colour (wavelength), intensity (irradiance) of the light emitted during phototherapy, the exposed body surface area and the infant's distance from the light<sup>11,12</sup>. In phototherapy treatment of term newborns with significant hyperbilirubinemia, double phototherapy provided more rapid and effective bilirubin reduction than conventional phototherapy alone due to higher spectral irradiance and larger body surface area exposed to phototherapy<sup>13</sup>. The use of phototherapy has decreased the need for exchange transfusion in term and preterm infants with haemolytic and nonhemolytic jaundice<sup>14</sup>. Intensive phototherapy may eliminate the need for exchange transfusion<sup>15</sup>. A number of other related articles in this region were reviewed<sup>16-67</sup>.

## **III. REFERENCES**

1. Bhutani VK, Johnson LH, Keren R. Diagnosis and management of hyperbilirubinemia in the term neonate: for a safer first week. *Pediatr Clin North Am.* 2004;51:843-61, vii.
2. Mills JF, Tudehope D. Fiberoptic phototherapy for neonatal jaundice. *Cochrane Database Syst Rev.* 2001; CD002060.
3. Maisels MJ, McDonagh AF. Phototherapy for neonatal jaundice. *N Engl J Med.* 2008;358:920-8.
4. Cremer RJ, Perryman PW, Richards DH. Influence of light on the hyperbilirubinaemia of infants. *Lancet.* 1958;1:1094-7.

5. Van Kaam AH, van Beek RH, Vergunst-van Keulen JG, van der Heijden J, Lutz-Dettinger N, Hop W, et al. Fibre optic versus conventional phototherapy for hyperbilirubinaemia in preterm infants. *Eur J Pediatr*. 1998;157:132-7.
6. Ennever JF. Blue light, green light, white light, more light: treatment of neonatal jaundice. *Clin Perinatol*. 1990;17:467-81.
7. Tan KL. Phototherapy for neonatal jaundice. *Clin Perinatol*. 1991;18:423-39.
8. Ventura-Junca P, Gonzalez A. Hiperbilirubinemia neonatal. In: Tapia JL, Ventura-Junca P, editors. *Manual de neonatología*. Santiago: Mediterraneo; 2000. p. 393-413.
9. Donzelli GP, Moroni M, Pratesi S, Rapisardi G, Agati G, Fusi F. Fiberoptic phototherapy in the management of jaundice in low birthweight neonates. *Acta Paediatr* 1996; 85: 366-70.
10. Holtrop PC, Ruedisueli K, Maisels MJ. Double versus single phototherapy in low birth weight newborns. *Pediatrics* 1992; 90: 674-7.
11. Maisels MJ, McDonagh AF. Phototherapy for Neonatal Jaundice. *N Engl J Med*. 2008; 358:920-928.
12. American Academy of Pediatrics Subcommittee on Hyperbilirubinemia. Management of hyperbilirubinemia in the newborn infant 35 or more weeks of gestation. *Pediatrics*. 2004; 114:297-316.
13. Sarici SU, Alpay F, Unay B, Ozcan O, Gokcay E. Doublesurface intensive phototherapy versus singlesurface conventional phototherapy in treatment of neonatal hyperbilirubinaemia. *J Trop Pediatr*. 2000;46(1):35-39.
14. .Ambalavanan N, Carlo WA. Jaundice and Hyperbilirubinemia in the Newborn, kernicterus. In: Kliegman R M, Stanton BF, St Geme III J W, Schor N F Behrman R E, editors. *Nelson Textbook of Pediatrics*. 19th ed. Philadelphia: Saunders Elsevier; 2011:603-612.
15. Wood A J, Dennery P A, Seidman D S, Stevenson D K. Neonatal hyperbilirubinemia. *N Engl J Med*. 2001;344(8): 581-590.
16. Grabenhenrich J, Grabenhenrich L, Bühner CC, Berns M. Transcutaneous Bilirubin After Phototherapy in Term and Preterm Infants. *Pediatrics* Nov 2014, 134 (5) e1324-e1329; DOI: 10.1542/peds.2014-1677.
17. Pendse A, Jasani B, Nanavati R, Kabra NS. Comparison of Transcutaneous Bilirubin With Total Serum Bilirubin in Preterm Neonates Receiving Phototherapy. 2016; *Perinatol Vol* 17(2) 44-49.
18. Taneja S, Pande V, Kumar H, Agarkhedkar S. Correlation of various maternal factors with exaggerated hyperbilirubinemia of the newborn. *J Datta Meghe Inst Med Sci Univ* 2017;12(3):218-222.
19. Singhania S, Singhania A, Khan S, Kumar V, Singhania P. Prenatal diagnosis of cross-fused renal ectopia: Still a dilemma. *Donald Sch J Ultrasound Obstet Gynecol* 2017;11(3):225-226.
20. Jain S, Sharma SK. Challenges & options in dengue prevention & control: A perspective from the 2015 outbreak. *Indian J Med Res* 2017;145(June):718-721.
21. Gupta V, Bhake A. Molecular Diagnosis of Tubercular Lymphadenopathy from Fine-Needle Aspirates in Pediatric Patients. *Acta Cytol* 2017;61(3):173-178.
22. Varghese LA, Taksande K. A comparison between intrathecal dexmedetomidine with hyperbaric bupivacaine and intrathecal fentanyl with hyperbaric bupivacaine in lower abdominal surgeries: A prospective double-blinded study. *J Datta Meghe Inst Med Sci Univ* 2017;12(2):99-109.

23. Khan KI, Jalgaonkar PD, Agrawal S. A case of phenytoin induced multiple toxicities. *J Datta Meghe Inst Med Sci Univ* 2017;12(2):157-158.
24. Bhalerao NS, Modak A, Belekar V. Comparison between magnesium sulfate (50 mg/kg) and lignocaine (2 mg/kg) for attenuation of intubation response in hypertensive patients. *J Datta Meghe Inst Med Sci Univ* 2017;12(2):118-120.
25. Phadnis P, Kamble MA, Daigavane S, Tidke P, Gautam S. Prevalence and risk factors – Hemoglobin A1c, serum magnesium, lipids, and microalbuminuria for diabetic retinopathy: A rural hospital-based study. *J Datta Meghe Inst Med Sci Univ* 2017;12(2):121-132.
26. Kuthe S, Sonkusale M, Wanjari A, Panbude P. Large right ventricular fibroma. *J Datta Meghe Inst Med Sci Univ* 2017;12(2):154-156.
27. Phatak S, Sadavarte T, Mishra G, Yadav S, Ali Jiواني MD, Patange N. Images: Emphysematous cystitis. *J Datta Meghe Inst Med Sci Univ* 2017;12(2):159-160.
28. Taware M, Sonkusale M, Deshpande R. Ultra-fast-tracking in cardiac anesthesia “Our Experience” in a rural setup. *J Datta Meghe Inst Med Sci Univ* 2017;12(2):110-114.
29. Cladius S, Jadhav U, Ghewade B, Ali S, Dhamgaye T. Study of diabetes mellitus in association with tuberculosis. *J Datta Meghe Inst Med Sci Univ* 2017;12(2):143-147.
30. Ali S, Ghewade B, Jadhav U, Cladius S. Study of serum interferon gamma in tubercular pleural effusions. *J Datta Meghe Inst Med Sci Univ* 2017;12(2):93-98.
31. Methwani DA, Deshmukh PT. Comparative study of type I tympanoplasty with or without mastoidectomy in tubotympanic type of chronic suppurative otitis media patients. *J Datta Meghe Inst Med Sci Univ* 2017;12(2):85-88.
32. Sarkar B, Bhake A. Serum prostate-specific antigen as a tumor marker for its correlation with histopathological diagnosis of prostatomegaly. *J Datta Meghe Inst Med Sci Univ* 2017;12(4):246-252.
33. Agarwal NK, Trivedi S. The partial pressure of oxygen in arterial blood: A relation with different fraction of inspired oxygen and atmospheric pressures. *J Datta Meghe Inst Med Sci Univ* 2017;12(4):280-283.
34. Roy M, Singh BR, Gajbe UL, Thute P. Anatomical variations of ureter in central India: A cadaveric study. *J Datta Meghe Inst Med Sci Univ* 2017;12(4):277-279.
35. Wasnik RR, Akarte NR. Evaluation of serum zinc and antioxidant vitamins in adolescent homozygous sickle cell patients in Wardha, district of central India. *J Clin Diagn Res* 2017;11(8):BC01-BC03.
36. Manisha S, Bagde N, Shrivastava D. Visual inspection of cervix with acetic acid: An alternative to cytology and colposcopy in early screening of cervical cancer in low-resource setup. *J Datta Meghe Inst Med Sci Univ* 2017;12(1):32-34.
37. Sharma T, Ghewade B, Jadhav U, Chaudhari S. Clinical profile of lung cancer at acharya vinoba bhavne rural hospital. *J Datta Meghe Inst Med Sci Univ* 2017;12(1):41-44.
38. Kalucha S, Mishra KK, Gedam SR. Noncompliance in psychosis. *J Datta Meghe Inst Med Sci Univ* 2017;12(1):61-65.
39. Palan A, Agrawal NK. Control of intraoperative shivering under spinal anaesthesia- A prospective randomized comparative study of butorphanol with tramadol. *J Krishna Inst Med Sci Univ* 2017;6(1):57-65.

40. Singh P, Basak S. Extended spectrum  $\beta$ -lactamases producing klebsiella pneumoniae: A threat to patient care. *J Datta Meghe Inst Med Sci Univ* 2017;12(4):234-237.
41. Charan N, Choudhari M, Sonkusale M, Deshpande R. Anesthetic management of chronic thromboembolic pulmonary hypertension for pulmonary endarterectomy. *J Datta Meghe Inst Med Sci Univ* 2017;12(4):289-291.
42. Kashikar SV. Congenital unilateral infiltrating facial lipomatosis. *West Indian Med J* 2017;66(1):189-190.
43. Kumar G, Phatak SV, Lakhkar B, Yadav SK. Diagnostic role of magnetic resonance imaging in rotator cuff pathologies. *J Datta Meghe Inst Med Sci Univ* 2017;12(1):7-10.
44. Tidake P, Sharma S. Profile and management of primary open-angle glaucoma patients above 40 years: A rural hospital-based study. *J Datta Meghe Inst Med Sci Univ* 2017;12(1):1-6.
45. Bhriegu R, Agrawal M, Hariharan C. Assessment of maternal and perinatal outcome in postdated pregnancy. *J Datta Meghe Inst Med Sci Univ* 2017;12(1):35-40.
46. Priya N, Lamture YR, Luthra L. A comparative study of scalpel versus surgical diathermy skin incisions in clean and clean-contaminated effective abdominal surgeries in AVBRH, Wardha, Maharashtra, India. *J Datta Meghe Inst Med Sci Univ* 2017;12(1):21-25.
47. Jaipuriya P, Pate MY, Iratwar S, Mahakalkar CC, Chandankhede A. Clinical study, evaluation, and management of cases of intracranial tumors admitted at Acharya Vinoba Bhave rural hospital, Sawangi (Meghe). *J Datta Meghe Inst Med Sci Univ* 2017;12(1):26-31.
48. Gupta M, Samal S, Shrivastava D, Bagde N, Mishra N, Gupta S. The study of ovulatory pattern following use of clomiphene citrate and anastrozole in infertile women with ovulatory dysfunction: A comparative study. *J Datta Meghe Inst Med Sci Univ* 2017;12(1):17-20.
49. Gupta K, Mahakalkar C, Kaple M, Deshpande S, Ladhha P, Jain N. A comparative study of cilostazol and pentoxifylline in intermittent claudication in peripheral arterial disease. *J Datta Meghe Inst Med Sci Univ* 2017;12(1):11-16.
50. Swarnkar M, Agrawal A. Kimura's disease: A case report and review of literature. *J Krishna Inst Med Sci Univ* 2017;6(3):118-120.
51. Swarnkar M, Jain SC. Web space lipoma causing separation of toes - a rare case report with review of literature. *J Krishna Inst Med Sci Univ* 2017;6(2):107-110.
52. Tendulkar MP, Ninave SS. Prospective comparison of pressor and airway responses to IV esmolol and IV dexmedetomidine during emergence from general anaesthesia and extubation. *J Krishna Inst Med Sci Univ* 2017;6(1):49-56.
53. Taksande AB, Jagzape AT, Deshpande VK. Study of motor nerve conduction velocity in patients of thyroid dysfunction in central India. *J Datta Meghe Inst Med Sci Univ* 2017;12(4):229-233.
54. Gade SA, Chari SN, Chalak A. Use of mini-CEX as a teaching learning method in physiology for undergraduate medical students. *Natl J Physiol Pharm Pharmacol* 2017;7(5):482-485.
55. Garg S, Chakravarti A, Singh R, Masthi NRR, Goyal RC, Jammy GR, et al. Dengue serotype-specific seroprevalence among 5- to 10-year-old children in India: a community-based cross-sectional study. *Int J Infect Dis* 2017;54:25-30.

56. Regmi PR, van Teijlingen E, Mahato P, Aryal N, Jadhav N, Simkhada P, et al. The health of Nepali migrants in India: A qualitative study of lifestyles and risks. *Int J Environ Res Public Health* 2019;16(19).
57. Bagde AD, Kuthe AM, Quazi S, Gupta V, Jaiswal S, Jyothilal S, et al. State of the Art Technology for Bone Tissue Engineering and Drug Delivery. *IRBM* 2019;40(3):133-144.
58. Khanam N, Wagh V, Gaidhane AM, Quazi SZ. Knowledge, attitude and practice on uses of plastic products, their disposal and environmental pollution: A study among school-going adolescents. *J Datta Meghe Inst Med Sci Univ* 2019;14(2):57-60.
59. Khanam N, Wagh V, Gaidhane AM, Quazi SZ. Assessment of work-related musculoskeletal morbidity, perceived causes and preventive activities practiced to reduce morbidity among brick field workers. *Ind J Community Health* 2019;31(2):213-219.
60. Zodpey S, Sharma A, Zahiruddin QS, Gaidhane A, Shrikhande S. Allopathic Doctors in India: Estimates, Norms and Projections. *J Health Manage* 2018;20(2):151-163.
61. Khatib MN, Shankar AH, Kirubakaran R, Gaidhane A, Gaidhane S, Simkhada P, et al. Ghrelin for the management of cachexia associated with cancer. *Cochrane Database Syst Rev* 2018;2018(2).
62. Khatib MN, Gaidhane A, Gaidhane S, Quazi ZS. Ghrelin as a promising therapeutic option for cancer cachexia. *Cell Physiol Biochem* 2018;48(5):2172-2188.
63. Gaidhane A, Sinha A, Khatib M, Simkhada P, Behere P, Saxena D, et al. A systematic review on effect of electronic media on diet, exercise, and sexual activity among adolescents. *Indian J Community Med* 2018;43(5):S56-S65.
64. Khatib M, Sinha A, Gaidhane A, Simkhada P, Behere P, Saxena D, et al. A systematic review on effect of electronic media among children and adolescents on substance abuse. *Indian J Community Med* 2018;43(5):S66-S72.
65. Khatib MN, Kirubakaran R, Gaidhane S, Shankar AH, Quazi Syed Z. Yoga for improving functional capacity, quality of life and cardiovascular outcomes in people with heart failure. *Cochrane Database Syst Rev* 2017;2017(7).
66. Uddin S, Mahmood H, Senarath U, Zahiruddin Q, Karn S, Rasheed S, et al. Analysis of stakeholders networks of infant and young child nutrition programmes in Sri Lanka, India, Nepal, Bangladesh and Pakistan. *BMC Public Health* 2017;17.
67. Puri S, Fernandez S, Puranik A, Anand D, Gaidhane A, Quazi Syed Z, et al. Policy content and stakeholder network analysis for infant and young child feeding in India. *BMC Public Health* 2017;17.