

areas of necrosis, multiple epitheloid and giant cell granulomas and diffuse lymphocytic infiltration consistent with tuberculosis. Step sections of the testis were normal. Three weeks following the surgery, repeat beta-hCG was normal (0.3 mIU/ml). Urine examination for acid-fast bacilli (AFB) and polymerase chain reaction for AFB were negative. The patient was started on four-drug anti-tubercular therapy.

Epididymal involvement in tuberculosis is through primary hematogenic spread to the globus minor which has a rich vascular supply. Diagnosis is based on the presentation and isolation of bacteria in the morning urine specimen or culture of material from discharging sinuses.¹ However, a pre-operative diagnosis can often not be made necessitating an inguinal orchidectomy with the suspicion of a testicular tumor.²

Beta-hCG is usually undetectable in normal adult men and elevated serum levels in any form of tuberculosis have not been reported. Affronti and DeBlaker demonstrated an association between mycobacterium tuberculosis and hCG in 1986.³ They noticed the production of hCG like substances by two non-tumor associated, virulent mycobacteria apart from other species of tumor associated aerobic bacteria. They postulated this production to be a variable character among bacterial species and a sign of conservation of similar features seen in prokaryotes and unicellular eukaryotes which also produce hormone like substances. The histopathological diagnosis of tuberculosis makes this the first case of epididymal tuberculosis with elevated beta-hCG in the

literature.

Jaya Kumar, Rajeev Kumar

Department of Urology, All India Institute of Medical Sciences, Ansari Nagar, New Delhi-110029, India.

E-mail: rajeevk@bol.net.in

REFERENCES

1. Gow JE. Genitourinary tuberculosis. In: Walsh PC, Retik AB, Vaughan Jr. ED, Wein AJ, (Ed). Campbell's Urology. 7th (Ed). Philadelphia: WB Saunders Company 1998:816-7.
2. Senzaki H, Watanabe H and Ishiguro Y. A case of very rare tuberculosis of the testis. Nippon Hinyokika Gakkai Zasshi 2001;92:534-7.
3. Affronti LF and DeBlaker DF. Immunological detection of hCG like substances in aerobic bacteria both tumor and non-tumor origin. Microbios 1986;48:173-82.

SEROPREVALENCE OF HIV, HBV, HCV AND SYPHILIS IN VOLUNTARY BLOOD DONORS

Sir,

Screening of blood is now mandatory for many diseases and is undertaken routinely in blood banks. Many studies have been done on human immunodeficiency virus (HIV), syphilis, Australia antigen (HBsAg), hepatitis C virus (HCV) separately, but the knowledge about the interrelationship between these transfusion transmitted diseases (TTD's) is limited, the present study was undertaken to find out the prevalence and correlation between various infectious markers in healthy blood donors.

A total of 44064 blood units collected in department of transfusion medicine, Dayanand

Table no 1 showing the incidence a percentage of various infectious disease markers in healthy blood donors

	Year 2001 (Jan-Dec)	Year 2002 (Jan-Dec)	Year 2003 (Jan-Oct)	Total/ %age
Total units bled	15042	13830	15192	44064
HBsAg positive	159	85	46	290/0.66%
AntiHBc positive	40	08	01	49/0.11%
Anti HCV positive	224	161	98	483/1.09%
HIV ½ positive	13	16	08	37/0.084%
VDRL positive	120	154	99	373/0.85%

Medical College & Hospital, Ludhiana, during the period of January 2001 to October 2003 were studied. No professional or honorary donor was bled. Screening of all the blood units for anti HIV-1/2, HBsAg, anti HBc, anti HCV & syphilis was done by a fully Automated Microplate Elisa Processor (ARIO model) from SEAC RADIM group using commercially available kits. Any serum found reactive by the first assay was retested using a second assay based on a different antigen preparation and/or different test principle. HIV seropositivity was seen to be 37/44064 (0.084%) and few of these were also confirmed by western blot test. HBsAg sropositivity was 290/44064 (0.66%), anti HBc positivity was 49/44064 (0.11%), anti HCV positivity was found to be 483/44064 (1.09%) and syphilis positivity was found to be 373/44064 (0.85%) as shown in Table 1. Also a definite correlation between positivity of HIV & syphilis was observed, but no correlation was seen between HIV and HBsAg/anti HBc/anti HCV positivity. The positivity of anti HBc was found to be more than positivity of HBsAg suggesting the ability to detect Hepatitis B virus (HBV) infection in window period.

Otuonye et al¹ observed that amongst the 150 (21.5%) patients positive with sexually transmitted diseases, 82 (54.65%) were found to be positive for HIV antibodies. Patil et al² also observed a positive correlation between

HIV & VDRL positivity. Therefore, serological screening for syphilis serves as a surrogate test for HIV infected donors. Jain et al³ and Gosavi et al⁴ found prevalence of anti HCV 1.57% in New Delhi and 15.9% in Mumbai, respectively. Risbud⁵ found that there was lack of evidence for sexual transmission of hepatitis C virus in patients attending STD clinics in Pune, India. Kothari⁶ observed that out of a total of 200 blood donors, 3% were positive for HBsAg, 1% for HIV, 2% for HCV and 4.5% for syphilis. So, taking into consideration rising prevalence of these infectious markers, a routine screening of all the donated blood units for anti HIV-1/2, HBsAg, anti HBc, anti HCV and syphilis should be done, which will assist blood transfusion services in improving blood product safety and donor recalls.

Nalini Gupta, Vijay Kumar, Amarjit Kaur

Dr. Nalini Gupta, House no- 2935, Sector 37-C, Chandigarh-160023, India.
E-mail: nalini203@rediffmail.com

REFERENCES:

1. Otuonye NM, Olukoya DK, Odunukwe WN, Idigbe EO, Udeaja MN, Bamidele M, Onyewuchie JI, Oparaugu CT, Ayelari OS, Oyekunle B. HIV association with conventional STDs (sexual transmitted diseases) in Lagos state, Nigeria. West Afr J Med 2002;21:153-6.
2. Patil AV, Pawar S D, Pratinidhi A K: Study of

- prevalence, trend and correlation between infectious disease markers of blood donors. *Ind. J. Hemat. Blood Transf.* 1996;14:95-102.
3. Jain A, Rana SS, Chakravarty P, Gupta RK, Murthy NS, Nath MC, Gururaja S, Chaturvedi N, Verma U, Kar P. The prevalence of hepatitis C virus antibodies among the voluntary blood donors of New Dehli, India. *Eur J Epidemiology* 2003;18:695-7.
 4. Gosavi MS, Shah SK, Shah SR et al; Prevalence of Hepatitis C virus infection in Mumbai. *Ind J Med Sci* 1997;51:378-85.
 5. Risbund A, Pereira M, Meherdale S, Gangakhedkar R, Ghate M, Joshi S, et al. Lack of evidence for sexual transmission of hepatitis C virus in patients attending STD clinics in Pune, India. *Sex Trans Infect* 2003;79:425.
 6. Kothari A, Ramachandran UG, Gupta P, Singh B, Talwar V. Seroprevalence of cytomegalovirus among voluntary blood donors in Dehli, India. *J Health Popul Nutr* 2002;20:348-51.

Indian Journal of Medical Sciences

Publication of The Indian Journal of Medical Sciences Trust

ISSN: 0019-5359

Published by:
Medknow Publications
www.medknow.com

SUBSCRIPTION RATES FOR 2004

	Annual	Individual issues*
In India (Rs.)	1000.00	350.00
Abroad (US \$)	100.00	35.00

*Please include Rs.50/\$5 for postage and handling

- Journal is indexed with Index Medicus/MEDLINE and Biological Abstracts
- Published monthly, subscriptions are for calendar year only
- Please return the subscription form to:
MEDKNOW PUBLICATIONS
12, Manisha Plaza, MN Road, Kurla (W), Mumbai 400070, India.
- Cheque should favour "Medknow Publications, Mumbai"
- Please allow at least six to eight weeks for commencement of new subscription.
- Back issues are available from 2001 onwards.
- Claims for missing issues can be made only within one month of publication

Subscription Form

Name of the subscriber
Current institutional attachment
Designation
Delivery address

City Pin code
State Country

Phone no. (with STD/ISD code)
E-mail address

Subscription details
Subscription period One year
Subscription type India / Foreign
Subscription starts from January (year)

Payment details
Cheque No Dated
Drawn on Amount

Signature Date:

PRACTITIONERS SECTION

MEGALOBlastic ANEMIA - PART I

ASHA SHAH

Megaloblastic anemia is most commonly due to deficiency of Folate or Cobalamin (vitamin B12). It is characterized by peculiar type of morphologic changes i.e. presence of megaloblasts or larger than normal sized precursors of red cells and granulocyte series in the bone marrow.

PHYSIOLOGY OF VITAMIN B12 AND FOLIC ACID

Vitamin B12

It is synthesized by micro-organisms. Higher plants and animals cannot synthesize it and must depend on external sources. Milk, vegetables, cereals, pulses, etc. are poor sources of B12. Curd, cheese, and such milk products contain B12. Contamination of legumes by B12 synthesizing bacteria is another source of B12 for a vegetarian. Non vegetarian articles of food like liver, kidney meat, oysters, crabs, egg yolk etc are rich sources of B12. B12 is synthesized by colonic bacteria but this is not available for absorption and is excreted. Fecal contamination of food and water which is responsible for infections like amebiasis etc. is ironically a source of B12 for majority of population of our country!

The daily requirement of B12 is only 1 microgram.

B12 absorption requires the presence of

intrinsic factor in the gastric juice. Hydrochloric acid and enzymes are required to split B12 from its combination with proteins in food. B12 is absorbed from lower part of ileum.'

Majority of B12 is stored in the liver (about 4-5 mg). Megaloblastosis occurs when the body B12 stores fall below 0.1 mg.

Folic acid

Folic acid is present in nearly all foods. Articles rich in folate are liver, yeast and green leafy vegetables. Prolonged cooking of food in large quantities of boiling water destroys folate and leads to folate-deficient diet. This is an important cause of dietary folate deficiency.

The daily requirement of folate is about 50 micrograms. Folate requirements increase in conditions with increased cell turnover for e.g. hemolytic anemias, malignancies, infections, etc.

Folic acid is absorbed from the entire length of the small intestine. Folate synthesized by the bacteria present in the colon is not absorbed.

The total body folate stores are about 5-10 mg and about one third of this is in the liver.

CAUSES OF B12 AND FOLATE DEFICIENCY