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Original Research Article

# Fasting Insulin and Glucose Concentrations among Adult Males of the Native Mising Population of Assam

Haren Baruah<sup>1</sup>, Debapriya Bandyopadhyay<sup>2</sup>, Thokchom Opendro Singh<sup>3</sup>, Anuradha Bharosay<sup>4</sup>, Anju Barhoi<sup>5</sup>

<sup>1</sup>Assistant Professor, Department of Biochemistry, Subharti Medical College, Meerut.

<sup>2</sup>Assistant Professor, Department of Biochemistry, All India Institute of Medical Sciences, Bhubaneswar.

<sup>3</sup>Post Graduate Student, Department of Biochemistry, Subharti Medical College, Meerut.

<sup>4</sup>Associate Professor, Department of Biochemistry, Chirayu Medical College, Bhopal.

<sup>5</sup>Associate Professor, Department of Biochemistry, Assam Medical College, Dibrugarh.

Corresponding Author: Debapriya Bandyopadhyay

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#### ABSTRACT

**Introduction:** The fasting Insulin and Glucose levels are very good indicators of an individuals' glucose metabolic status. Glucose metabolism among different ethnic groups may vary due to difference in genetic makeup as well as their lifestyles. A study was carried out on a specific ethnic group called "*Mising tribe*" of Assam, to evaluate metabolic status in this unique population.

**Methods:** In this work, the fasting serum Insulin and plasma Glucose were recorded for a group of 50 randomly selected non-diabetic males of the indigenous *Mising* tribe of upper Assam, of the age group 40-50 years. Their biophysical profiles and their physical activity status were recorded and compared with their fasting Insulin and Glucose levels.

**Result:** The mean fasting serum insulin was found to be  $13.6 \pm 4.62 \mu U/ml$ , mean fasting glucose was  $98.39 \pm 6.74$  mg/dl, the mean fasting insulin: glucose ratio in our study was  $0.121 \pm 0.07$  which is again within the normal range (i.e. < 0.4), comparable to the findings of other workers. The fasting insulin levels were found to be inversely proportional to the Physical activity status.

**Conclusion:** Sedentary, so called urban lifestyle maybe associated with higher fasting Insulin levels, which might have a strong role in the etiopathogenesis of pre-dominantly urban diseases like diabetes, obesity and cancers.

Key Words: Insulin, Insulin: Glucose ratio, Mising tribe, Modified Activity Questionnaire (MAQ)

#### **INTRODUCTION**

Insulin, the protein proved to have hormonal action, secreted by the islet of Langerhans of pancreas is directly related to several diseases. On one hand, ageing, poor diet, stress among other factors can deprive cells of their Insulin sensitivity producing Diabetes Mellitus; on the other hand, as a result of insulin resistance, the pancreas produces more insulin than normal so there are higher levels of Insulin circulating in the bloodstream or hyperinsulinemia. Hyperinsulinemia and insulin resistance may lead to several problems, including elevated triglycerides, <sup>(1-4)</sup> low HDL, <sup>(5,6)</sup> type II diabetes, <sup>(7-9)</sup> and obesity. <sup>(10-13)</sup>

Regular exercise and active lifestyle is known to improve glycemic control, delay the onset or even prevent Diabetes mellitus.

The *Mising* tribe is an isolated, hardworking, ethnic, native population of upper Assam. No published data is available on their Biophysical profiles and Physical activity status. So we wanted to study the levels of fasting Insulin and Glucose levels in this indigenous population, and correlate these levels with their Biophysical profiles and physical activity status, since this could throw some light on the role of regular physical activity on the pathophysiology of lifestyle based metabolic diseases.

## **MATERIALS AND METHODS**

The 'Misings' are one of the colorful Mongoloid Tribes of the northeast India. They have their own history socio-cultural origin, life and of language. This ethnic group from N-E region were previously known as 'Miris'; they are mentioned as such in the list of schedule tribes of India under constitution order 1950. The present study was conducted among the male members of this group in Mising dominated villages of upper Assam, namely

- Natun Chiring Gaon, P.O. & District Dibrugarh
- Pani Miri Gaon, P.O. Kolakhowa Chariali, District Dibrugarh

The study was conducted during the period June to July 2013, in Assam Medical College and Hospital, Dibrugarh, Assam and Subharti Medical College, Meerut, Uttar Pradesh.

The sample size was calculated after knowing the total population of male *Mising* persons in the age group of 40-50 years in *Mising* dominated villages, where the study was carried out. As the proposed study was a prevalence study, a 10% sample from the targeted population was adequate to achieve the objective of the study.

The number of missing populated villages of the region was identified and from these villages cases were selected by the random process and thereafter consecutive house in clockwise direction chosen for the purpose of selecting the subjects.

Only healthy male *Mising* subjects between 40-50 years were included in this study. Subjects with diabetes mellitus, any form of malignancy and first degree family history of diabetes mellitus were excluded.

A home visit was made prior to sample collection and the purpose of visit and procedure of testing was explained. An informed consent for participating in the test was recorded. The name, age, height, weight and other biophysical parameters, family history suggestive of Diabetes. Thyroid disorders, Pancreatitis, Jaundice and other relevant details were recorded in a proforma developed for the purpose. Subjects known to be having diabetes and or on treatment were excluded from the study.

Blood pressure was measured using a mercury sphygmomanometer in comfortable sitting position in the right arm.

Sample Collection and Biochemical testing:

The subjects were instructed to stay in fasting state for the early

morning venous blood sample to be collected on the decided day. With aseptic precautions (A) 2ml venous blood in polystyrene tube of 25mm x 75mm size for insulin estimation and (B) 2ml venous blood in NaF vial for glucose estimation were collected from antecubital vein after overnight fasting (at least 8 hours).

Sample after collection:

- A. For insulin estimation, the serum was kept at room temperature and allowed to clot. Centrifugation done. serum was was the collected in a separate vial and preserved at  $-20^{\circ}$ C at Dept of Biochemistry, Assam Medical College, Dibrugarh. The sample collection of the entire study group was done over a span of 4 days, then the collected serum were transferred to portable carrier avoid vaccine to denaturation of protein. They were brought to Subharti Medical College, Meerut by air and Insulin was estimated in a ROBONIK ELISA washer and using solid reader. phase sandwich ELISA diagnostic kits (EIA 2935) from DRG Germany, on the same day.
- B. For the estimation of blood sugar 2ml blood collected in NaF vials were centrifuged at 5000 RPM for 10 mins and the plasma was used for estimation of Glucose by GOD-POD method, using Glucose estimation kit from SPAN, in a Semi-automatic analvzer within 6 hours of collection, at Dept of Biochemistry, Assam Medical College, Dibrugarh.

Assessment of Physical activity: The Modified Activity Questionnaire (MAQ), <sup>(14)</sup> adapted from

the Pima Indian Ouestionnaire, for suitability and appropriateness of the various indigenous physical activities undertaken by the population under study, was filled to assess the physical status of the selected activity individuals. The MAQ has been shown to be both reliable and valid for adults. (15) The occupational and leisure estimates calculated from this comprehensive interviewer-administered questionnaire can be combined to arrive at an overall estimate of physical activity levels. The MAQ can estimate the individual's physical activity level over the past one year, expressed as hours per week, by a crude estimate of the metabolic cost of each activity undertaken and expressed as MET hours per week. The MET is the ratio of working metabolic rate of an activity divided by the resting metabolic rate.

The MAQ used in this study, as well as detailed instructions for administrating the questionnaire and calculating the results can be found in "A Collection of Physical Activity Questionnaires for Health-Related Research". <sup>(15)</sup>

## RESULTS

This present community based cross-sectional study includes 50 healthy male subjects, within the age group of 40-50 years, of the isolated tribal *Mising* population of upper Assam. In this study a survey of biochemical profiles on glucose and insulin in basal state and their variation with physical activity status among members of this isolated tribal population has been conducted.

The following tables illustrate the results of this study:

Table 1: Mean Fasting Serum Insulin level ( $\mu$ U/ml), Mean Fasting Blood Glucose Level (mg/dl), and Mean Fasting Insulin: Glucose ratio of 50 subjects, based on their occupation.

Groups	Total no of	Fasting Insulin	Fasting Blood	Fasting Insulin:
	subjects	(µU/ml)	Glucose (mg/dl)	Glucose Ratio
Service	12	12.78 (±8.36)	$107.92 \pm 19.83$	0.201 (±0.083)
Cultivators	38	10.18 (±7.92)	96 ±9.23	0.158 (±0.069)

Table 2: Comparison between subjects with family history (F/H) and without family history of Diabetes Mellitus (DM) (not 1<sup>st</sup> degree relative).

	Groups	Total no of	Fasting Insulin	Fasting Blood Glucose	Fasting Insulin: Glucose		
		subjects	(µU/ml)	(mg/dl)	Ratio		
	Subjects having family history of Diabetes Mellitus	5	$13.95 \pm 12.54$	$115\pm6.96$	$0.214 \pm 0.14$		
ŀ	Subjects without family history	45	$10.45 \pm 6.08$	97.28 ± 6.37	0.117 ± 0.07		
	of Diabetes Mellitus						
			P < 0.01	P is 0.001	P < 0.05		

Table 2 shows the comparison between the two groups of people having family history of diabetes mellitus and no family history of diabetes mellitus. It is to be noted that subjects having family history were not first degree relatives. It is observed that persons having family history of diabetes mellitus have got higher mean fasting glucose, mean fasting insulin and a mean fasting Insulin:Glucose ratio. The mean difference is almost 10mg/dl in fasting glucose and 9.5  $\mu$ U/ml in Fasting Insulin. Fasting glucose level shows a very high significance (p < 0.001) when compared between subjects with positive family history and negative family history of diabetes mellitus, while fasting insulin level is found to be highly significant (p < 0.01).

Table 3: Shows the mean values of Fasting insulin, Fasting glucose and Fasting Insulin:Glucose ratio, compared between mild hypertensive and normotensive subjects, there are little difference in these values carrying no significance (p>0.05).

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Groups	Total no of	Fasting Insulin	Fasting Blood	Fasting Insulin:			
	subjects	(µU/ml)	Glucose (mg/dl)	Glucose Ratio			
Hypertensive	5	$11.43 \pm 7.25$	$102.2 \pm 5.16$	$0.140\pm0.080$			
Normotensive	45	$10.73 \pm 6.78$	$98.49 \pm 6.81$	$0.121\pm0.076$			
		p>0.05	p>0.05	p>0.05			

Among the 50 subjects in the present study, 5 subjects are found to be hypertensive, taking cut-off value as 140/90mmHg, the rest 45 were normotensive.

 Table 4: Shows comparison of mean Fasting Insulin, Fasting Glucose and Fasting Insulin: Glucose ratio with corresponding Body mass Index.

	Groups	Total no of	Fasting Insulin	Fasting Blood	Fasting Insulin:	
	BMI $(kg/m^2)$	subjects	(µU/ml)	Glucose (mg/dl)	Glucose Ratio	
	(A) 22.1-25	30	11.03±6.67	98.88±6.38	0.123±0.074	
	(B) < 22	17	9.93±7.27	97.61±7.24	0.119±0.083	
(C) > 25 3		13.70±5.55 105.16±7.22		0.128±0.065		

p>0.05 for Fasting Glucose (A Vs. B)p>0.05 for Fasting Glucose (A Vs. C)p>0.05 for Fasting Insulin (A Vs. B)p>0.05 for Fasting Insulin (A Vs. C)p>0.01 for Fasting I:G ratio (A Vs. B)p>0.05 for Fasting I:G ratio (A Vs. C)

It is observed that the values of mean fasting glucose gradually increasing from Group-B to A and A to C. Again mean fasting insulin values between A and C are almost same but it is little bit lower in Group-B. Similarly there is almost same value of insulin: glucose ratio between A & C, but slight lower value in Group-B. Statistically, it has been observed that there are no significant (p > 0.05)relationship between mean fasting glucose and mean fasting insulin levels when group A is compared with group B or C. Similarly, no significance (p > 0.05) is observed in their glucose: insulin ratio also.

Table 5: Shows the comparison among subjects performing various amount of physical activity, which has been recorded as per the modified PIMA Indian questionnaire. The Group D subjects have activity >20 MET hours, Group E: 15-20 MET hours, Group F: < 15 MET hours per week. The commonly undertaken work related activity was agriculture related, while leisure activities included bicycling, u8walking, chopping wood, carrying water, rearing animals (cows and pigs), gardening and sporting activities.

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	Activity	No	of	Physical A	Activity	Mean	Mean Fasting	Mean I:G	
	Status	Subject		(hours per	wk) <sup>#</sup>	Fasting	Insulin	Ratio	
				Work	Leisure	Glucose			
	Heavy (D)	36		13.1	8.2	$96.99 \pm 9.23$	9.82±4.93	$0.087 \pm 0.069$	
	Moderate (E)	10		9.2	6.5	$99.65 \pm 6.89$	12.19±6.33	0.112±0.071	
	Mild (F)	4		6.9	5.6	113.66±7.34	$16.2 \pm 7.43$	$0.164 \pm 0.081$	

# median hours/week activity average over past one year. p>0.05 for glucose (F Vs. E) p>0.05 for glucose (F Vs. D) p<0.01 for insulin (F Vs. E) p<0.05 for insulin (F Vs. D) p<0.01 for I:G ratio (F Vs. E) p<0.05 for I:G ratio (F Vs. D)

Statistically gives it no significance for fasting glucose (p > p)0.050) when subjects of group (D) were compared with group (E) or group (F). But there is high significance (p < 0.01) for fasting insulin level when Group-D was compared with F and significant correlation (p < 0.05) when compared with group E. Again for insulin: glucose ratios show high significant variation (p < 0.01) when Group-D is compared with F and significant (p < 0.05) correlation when compared with Group-E.

Also, on comparing the recorded total physical activity values (MET) with fasting glucose levels we find a clear trend of worsening fasting glucose with decreasing physical activity levels among the study group.

## **DISCUSSION**

This study is a unique work, since it is an endeavor to investigate the metabolic status of the indigenous ethnic native population of upper Assam- the *Mising* tribe, by simultaneously estimating Insulin and Glucose levels in the fasting state and correlate the same with their physical activity status. There are some studies done on their food habits

and social setup, <sup>(16)</sup> but no information is available on the Insulin levels, Fasting blood glucose levels and its correlation with Body Mass Index and Physical Activity status of this population. This study also throws light upon the ill effects of so-called urban sedentary lifestyle, where food, water and other essentials are easily available, on (17-19) metabolism of Glucose. The population included in this study, is a typical example of people who are undergoing a lifestyle change, from having rapid cultivation and hunting as their mainstay occupation, some are switching to office jobs, shifting to urban areas and adopting lifestyle which might involve more mental and psychological stress but much less physical activity. Probably as a consequence of this, we found a higher level of circulating insulin in this sub-group, compared to their farming counterparts, similar to other studies. <sup>(20-22)</sup> Moreover, the socioeconomic status of cultivator group is also lower than the service group which may be another factor. Our results are similar to what the previous workers had shown, that there exists and inverse relationship between socioeconomic status and the prevalence of diabetes.<sup>(23)</sup>

Since we have excluded subjects with diabetes or with diabetic first degree relative, the fasting Glucose levels in all subjects studied were within the normal reference range.

The incidence of positive family history of diabetes mellitus (other than 1st degree relatives) was 5%. This incidence seems to be low compared to in other studied communities like Parsees and Sikhs. This is probably due to non-prevalence of inter-caste marriages among the *Misings*.

Ingestion and absorption of dietary Glucose triggers the release of a burst of the peptide Insulin which enables Glucose uptake by liver and muscle cells. Once glucose has been shifted inside cells for storage or energy release, insulin levels should drop to very low or even to undetectable levels. <sup>(24)</sup> Only a tiny amount of basal insulin is required for glucose homeostasis. In our study, we found that among the people with weekly physical activity more than 20 MET hours, the mean fasting Insulin was 9.82±4.93 µIU/mL. But for people with less than 15 MET hours activity per week, the mean fasting Insulin was 16.2±7.43 µIU/mL. Insulin being a strongly anabolic hormone, high levels can promote cell growth, predisposing to certain cancers. <sup>(25-29)</sup> Higher Insulin levels also trigger the synthesis of beta-amyloid proteins in brain and may contribute to the development of Alzheimer's disease. (30) Overproduction of Insulin is even a contributory factor to prostate enlargement because of its effects in promoting the overgrowth of prostate cells.<sup>(31)</sup> Therefore several of these so called urban diseases, may have a very intricate lifestyle based etiopathogenesis.

The study of 'Fasting Insulin and Glucose concentrations among adult males of the native Mising population of Assam' is a preliminary one, carried out among a small section of missing community. So, it is thought that there are reasonable scope for further studies in this field which would uphold new vistas for a better understanding of the etio-pathogenesis and prevention of highly prevalent so called urban noncommunicable diseases like Diabetes Mellitus and Obesity.

#### CONCLUSION

A physically active lifestyle is an absolute must to prevent metabolic diseases like diabetes mellitus and obesity; or to delay the onset or slow the progression of degenerative diseases and cancer.

#### REFERENCES

- Godsland IF, Crook D, Walton C, Wynn V, Oliver MF. Influence of insulin resistance, secretion, and clearance on serum cholesterol, triglycerides, lipoprotein cholesterol, and blood pressure in healthy men. Arterioscler Thromb. 1992 Sep;12 (9):1030-5.
- Salonen JT, Lakka TA, Lakka HM, Valkonen VP, Everson SA, Kaplan GA. Hyperinsulinemia is associated with the incidence of hypertension and dyslipidemia in middle-aged men. Diabetes. 1998 Feb;47(2):270-5.
- Stannard SR, Johnson NA. Insulin resistance and elevated triglyceride in muscle: more important for survival than "thrifty" genes? J Physiol. 2004 Feb 1;554 (Pt 3):595-607.
- Mykkänen L, Kuusisto J, Haffner SM, Pyörälä K, Laakso M. Hyperinsulinemia predicts multiple atherogenic changes in lipoproteins in elderly subjects. Arterioscler Thromb. 1994 Apr; 14(4):518-26.
- Karhapää P, Malkki M, Laakso M. Isolated low HDL cholesterol. An insulin-resistant state. Diabetes. 1994 Mar; 43(3):411-7.

- Ko GT, Cockram CS, Woo J, Chan JC. Obesity, insulin resistance and isolated low high-density-lipoprotein cholesterol in Chinese subjects. Diabet Med. 2001 Aug; 18(8):663-6.
- Goldstein BJ. Insulin resistance as the core defect in type 2 diabetes mellitus. Am J Cardiol. 2002 Sep 5; 90(5A):3G-10G.
- 8. Haffner SM, Stern MP, Mitchell BD, Hazuda HP, Patterson JK. Incidence of type II diabetes in Mexican Americans predicted by fasting insulin and glucose levels, obesity, and body-fat distribution. Diabetes. 1990 Mar; 39(3):283-9.
- 9. Hansen J, Rinnov A, Krogh-Madsen R, C P Fischer, A S Andreasen, R M G Berg, et al. Plasma follistatin is elevated in patients with type 2 diabetes: relationship to hyperglycemia, hyperinsulinemia, and systemic low-grade inflammation. Diabetes Metab Res Rev. 2013 Sept; 29(6):463–472.
- 10. Chu N, Spiegelman D, Hotamisligil GS, Rifai N, Stampler M, Rimm EB. Plasma insulin, leptin, and soluble TNF receptors levels in relation to obesity-related atherogenic and thrombogenic cardiovascular disease risk factors among men. Atherosclerosis. 2001 Aug; 157(2):495-503.
- 11. Despres JP, Pascot A, Lemieux I. Risk factors associated with obesity: a metabolic perspective. Ann Endocrinol. 2000 Dec; 61 Suppl 6:31-8.
- 12. Modan M, Halkin H, Almog S, Lusky A, Eshkol A, Shefi M, et al. Hyperinsulinemia. A link between hypertension obesity and glucose intolerance. J Clin Invest. 1985 Mar; 75(3):809-17.
- 13. Feuers RJ, Desai VG, Chen FX, Hunter JD, Duffy PH, Oriaku ET. Effects of dietary restriction on insulin resistance in obese mice. J

Am Aging Assoc. 2000 Apr; 23(2):95-101.

- 14. Kriska AM, Knowler WC, LaPorte RE, Drash AL, Wing RR, Blair SN, Kuller Bennett PH. LH. Development of questionnaire to examine relationship of physical and diabetes in activity Pima Indians. Diabetes Care. 1990: 13:401-411.
- 15. Kriska AM, Caspersen CJ. Introduction to the collection of physical activity questionnaires: In A Collection of Physical Activity Questionnaires for Health-Related Research. Kriska AM, Caspersen CJ, Eds. Med Sci Sports Exerc 29.
- 16. Pratisha Kumari. Kitchen and Dining Space: As A Way of Eating Manner in Mising Community. IOSR Journal of Humanities and Social Science. 2012; 4(5):23-28.
- 17. Kristine Færch, Allan Vaag, Jens J Torben Hansen, Holst, Torben Jørgensen, Knut Borch-Johnsen. Natural History of Insulin Sensitivity and Insulin Secretion in Progression from Normal the Glucose Tolerance to Impaired Glycemia and Impaired Fasting Glucose Tolerance: The Inter 99 Study. Diabetes Care. 2009; 32(3): 439-444.
- 18. James B M, Denis C M, David M N, Deirdre R B, Reubin A. The Natural history of Progression from Normal Glucose Tolerance to Type 2 Diabetes in the **Baltimore** Study Longitudinal of Aging. Diabetes. 2003; 52: 1475-1484.
- Ele Ferrannini, Monica Nannipieri, Ken Williams, Clicerio Gonzales, Steve M. Haffner, Michael P Stern. Mode of onset of type 2 diabetes from normal or impaired glucose tolerance. Diabetes. 2004; 53:160-165.
- 20. Kim SM, Lee JS, Lee J, Na JK, Han JH, Yoon DK, et al. Prevalence of diabetes and impaired fasting

glucose in Korea: Korean National Health and Nutrition Survey 2001. Diabetes Care 2006; 29(2): 226–231.

- 21. Tran Quang Binh, Pham Tran Phuong, Bui Thi Nhung, Dang Dinh Thoang, Pham Van Thang, Tran Khanh Long, et al. Prevalence and correlates of hyperglycemia in a rural population, Vietnam: implications from a cross–sectional study. BMC Public Health. 2012 Nov; 12: 939.
- 22. Andrea M Kriska, Stewart B Harris, Anthony JG Hanley, Bernard Zinman. Physical Activity, Physical Fitness, and Insulin and Glucose Concentrations in an Isolated Native Canadian Population Experiencing Rapid Lifestyle Change. Diabetes Care. 2001; 24:1787–1792.
- 23. Steven M. Haffner. Epidemiology of Type 2 Diabetes: Risk Factors. Diabetes Care. 1998; 21(3): C3-C6.
- 24. Available at: <u>http://www.a4m.com</u> /assets/pdf/bookstore/thera6\_ch4.pdf ? SESSION\_MAIN= dap3mjgnb96luhbje8tcklgp75. Accessed April 29, 2013.
- 25. Nilsen TI, Vatten LJ. Prospective study of colorectal cancer risk and physical activity, diabetes, blood glucose and BMI: exploring the hyperinsulinaemia hypothesis. Br J Cancer. 2001 Feb 2; 84(3):417-22.

- 26. Balkau B, Kahn HS, Courbon D, Eschwege E, Ducimetier P. Hyperinsulinemia predicts fatal liver cancer but is inversely associated with fatal cancer at some other sites. Diabetes Care. 2001 May; 24(5):843-9.
- 27. Kaaks R. Plasma insulin, IGF-I and breast cancer. Gynecol Obstet Fertil. 2001 Mar; 29(3):185-91.
- 28. Czyzyk A, Szczepanik Z. Diabetes mellitus and cancer. Eur J Intern Med. 2000 Oct; 11(5):245-52.
- 29. Bruce WR, Wolever TM, Giacca A. Mechanisms linking diet and colorectal cancer: the possible role of insulin resistance. Nutr Cancer. 2000; 37(1):19-26.
- 30. David G Cook, James B Leverenz, Pamela J McMillan, J Jacob Kulstad, Sasha Ericksen, Richard A Roth, et al. Reduced hippocampal insulindegrading enzyme in late-onset Alzheimer's disease is associated with the apolipoprotein E-epsilon4 allele. American Journal of Pathology. 2003 Jan; 162(1):313-9.
- 31. Hammarsten J, Hogstedt B. Hyperinsulinaemia as a risk factor for developing benign prostatic hyperplasia. Eur Urol. 2001 Feb; 39(2):151-8.

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