ARMS-PCR Based SNP Analysis of MTHFR C677T Allele Using Syber Green in Pancreatic Tumour

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Authors’ contributions

This work was carried out in collaboration between all authors. All authors read and approved the final manuscript.

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ABSTRACT

In human, carcinoma of pancreas, a rare disease and mortality rate is quite high in Indian population. Epidemiological studies support the hypothesis that folate metabolism regulate DNA stability and prevent cancer. Because folate have been linked to dietary supplement and defect in folate metabolism have been increase risk of developing cancer shows adverse effect on health conditions. In the present study we have assess methylene tetrahydrofolate reductase gene polymorphism (MTHFR) 677 C➔T using ARMS PCR based SNP analysis using Syber green in the cases of pancreatic tumour to determine the “risk factor”. Interestingly, our findings reveals that 33.0% frequency (one case) showing mutation of MTHFR 677 TT genotype (rare type) in homozygous condition with Tm value 82.50°C for mutant 677T allele shifted to 677C allele (83.50°C). Two cases (66%) showing CC (wild type) allele and Tm value 82.83°C for 677C allele.

In MTHFR 677TT is a rare mutation and individuals show very low enzymatic activity due to the substitution of alanine to valine. The study further continue to confirm the mutation by

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visualization on agarose gel electrophoresis of the same PCR product, again showing (Lane- 2) the band of 50 kb of mutant 677TT allele in the same case, suggesting an relevant role of folate metabolism and subsequent impairment of aberrant DNA methylation during carcinogenesis with increasing “high risk” for the development of carcinoma due to allele TT mutation of MTHFR. However, large samples size is required to further confirm the association.

Keywords: Pancreatic tumour; ARMS PCR; MTHFR polymorphism.

1. INTRODUCTION

Pancreatic tumour, a most common form of cancer mortality and their incidence varies in different countries due to different life style including food habit and other environmental factors [1,2]. To reduce the global burden of pancreatic cancer in the coming decades it is become necessary to identify the “risk factor” for the prevention of such disease in this region. Since methylenetetrahydrofolate reductase (MTHFR) play an important role to regulate folate metabolism by irreversible catalyzing the conversion of 5,10- methylenetetrahydrofolate to 5- methylenetetrahydrofolate and is also involved against oxidative stress to maintain pancreatic homeostasis during injury [3,4]. Recent studies of MTHFR 677 C → T gene polymorphism on cancer risk are highly controversial and conflicting findings have been documented in the current literature [5-8]. However, earlier studies of same authors on MTHFR gene in other carcinoma of cervix and ovary rather than pancreas has been associated with risk factor [9,10]. Therefore, the interest has been generated to assess a short case study to find out the mutation of polymorphic genes variation in methylenetetrahydrofolate reductase (MTHFR) using sensitive allele refractory mutation system – polymerase chain reaction (ARMS PCR).

2. MATERIALS AND METHODS

2.1 Samples

Blood samples were obtained in sterile EDTA vials from pancreatic cancer patients (n=3) and controls of the same age group after clinical diagnosis from the OPD of All India Institute of Medical Sciences who refers these cases to Genetic Laboratory of the Department of Pathology & Lab Medicine for the study of MTHFR C677T polymorphism by RT PCR (Bio Red Mini Opticon,USA) Syber green method. Genomic DNA were isolated using Bioneer kits (Korea) and quantified using Nanodrop spectrophotometer (Thermoscientific, USA) and samples were stored at -20°C till the SNP analysis was performed.

2.2 Primer Design

The primers for tetra plex real-time PCR assay were designed for genotyping of MTHFR 677C T (http://cedar.genetics.soton.ac.uk/public_html/primer1.html) and further confirmed by BLAST program at http:// www.ncbi.nlm.nih.gov/blast to determine the specificity of the primers. To increase the specificity of the reaction a mismatch at the 2 position of the 3’ end both the allele-specific primers were selected and further confirmed by software [11]. Earlier, the effect of the mismatch at the 2 position of the 3’ end of the allele-specific primers on the specificity of the reaction already documented in the literature [12]. To obtain amplicons with distinct melting points, the hypothetical ‘Tm’ values were calculated using known software or Web sites (for example, http://eu.idtdna.com/analyzer/Applications/OligoAnalyzer/). The selection of the primers were based on the amplicons ‘Tm’ values and following primers used in present study: MTHFR-T, 5’ – GCACCTGAAAGGAAGGTGTCTGCGGGCGT-3’ ; MT MTHFR-C-polyG, 5’ – GGCCTGGCAGGGGAAAGCTGGTAGATGA TGAAATAGG-3’ ; MTHFR-cf, 5’ - TGTCATCCTATTGGCAGTTACCACCAA-3’ ; MTHFR-cr, 5’ - CCATGTCCGTGCATGCTTCACAAA-3’ [13].These group of tetra-primer ARMS PCR using SYBR Green based on melting-point (Tm) analysis was part of strategy of our interest to detect SNP of mutant of MTHFR allele (wild-type & mutant alleles and also the same PCR product further confirmed by agarose gel electrophoresis.

2.3 Real-time PCR Method

A total volume of 20 µl containing 10 µl of SYBR Green PCR Master Mix (Bio red USA), 1 µl of each primer per reaction, 40 ng of genomic DNA,
and distilled water was taken to performed Real-time PCR. The PCR protocol on the Light Cycler (BioRed USA) was as an initial denaturation step (95 °C for 7 min) was followed by amplification and quantification steps repeated for 30–40 cycles (95°C for 10 s, 60°C for 10 s, 72°C for 20 s, with a single fluorescence measurement at the end of the elongation step at 72°C, a melting-curve analyzed the data and reaction was terminated by cooling to 40°C. Melting curves were constructed by lowering the temperature to 65°C and later increasing the temperature by 0.2°C/s to 98°C to measuring the change in fluorescence consistently. Tm values were assigned to develop plot generated by the RTPCR of the negative derivation of fluorescence versus temperature (dF/dT) of the melting curve for amplification products measured at 530 nm.

3. RESULTS

In the present study of MTHFR gene polymorphism carry out in few cases of carcinoma of pancreas using highly sensitive and reliable method using tetra-primer PCR which amplifies both wild-type and mutant alleles 677C>T of MTHFR gene with a controls of the same group in a single PCR tube. The method employs four set of primers to amplify a larger non-allele- specific fragment containing the mutation site and allele-specific amplicons representing each of the two allelic forms. The study reveals that of Tetra-primers PCR generated amplicons with Tm value 82.83°C for the 677C allele (controls) and Tm value 82.50°C for the 677T allele, and 83.50°C for 677C allele in cases of carcinoma of pancreas. Because Tm values are close to one another, a short GC tail is added to the inner 677C allele-specific primer and genotype support on the Tm specific amplicons with unique shape of the melting peaks as documented in Figs. 1a & b.

The above findings based on ARMS PCR further confirmed by agarose gel electrophoresis in the same PCR product (both cases & controls). In one case (33%) showing rare 677TT mutant allele as shown in Lane-2 and having and of 50bp band, while in two cases (wild type) 677CC (66%) allele were observed in homozygous condition (Lane-1 & 3). In case of controls samples only 677CC (wild type) bands are visualized in lane 4, 5 & 6 (Fig. 2).

4. DISCUSSION

Etiopathology of tumour biology is highly complex and pancreatic carcinoma is one of the important for surgical point of view neoplasia associated high risk of mortality rate. Epidemiological studies shows that an association between low folate intake and an increased risk for the development of cancer. In this manuscript, we have undertaken a case-control study to investigate the role of MTHFR 677C → T gene polymorphisms and their susceptibility in pancreatic tumour. MTHFR, thermolabile enzyme that play a crucial role in the folate metabolic pathway involving DNA synthesis and methylation.

![Fig. 1a & b. ARMS T-Plex real - time PCR assay for MTHFR C677T genotyping using wild-type (CC) and mutant (TT) allele (1a). The effect of addition of a short GC tail on the Tm value of the 677C allele with melting-curve analysis for the allele specific and non allele specific amplicons between cases and controls samples (Fig. 1b)
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Interestingly, in the recent year we have observed the genetic susceptibility in various types of tumor lead to a growing attention to the MTHFR C677T gene polymorphisms associated risk in tumour biology. Interestingly, polymorphisms in MTHFR may interact with folate status to influence genomic methylation. A recent study by Frisco et al reports that subjects with low folate status and carrying the TT variant of the MTHFR 677 C→T (ala→val) polymorphism had relative hypomethylation of DNA extracted from whole blood compared with homozygous wild types, an effect that was mitigated by high folate status [14]. Methyleneetetrahydrofolate reductase (MTHFR) mutation are commonly associated with folate metabolism to increase “risk factor” for the development of neural tube defects, recurrent pregnancy loss and development of pancreatic carcinoma with genetic interaction between MTHFR alleles has been poorly defined in Eastern region of India.

Folic acid metabolism plays an important role in the maintenance of genomic stability. Dietary factors including folate play an important protective role and there are also plausible explanatory mechanisms. Factor influencing development of such cancer risk is still not known but certainly associated with genetic and epigenetic variation regulating folate metabolism [15]. The MTHFR 677 TT genotype led to elevated homocysteine levels and DNA hypomethylation in folate-depleted subjects lead to provide the genomic instability, leading to carcinogenesis. Because one case showing 677TT mutation (33%) of MTHFR polymorphisms determining variant “T” genotype as a risk factor for carcinoma of pancreas and changes in the mutation frequency may be either due racial differences or environmental factor including dietary supplement between population. Metaanalysis study revealed that there is a controversial finding between different groups of the authors regarding involvement of MTHFR 677C→T gene polymorphism in development of adenocarcinoma of pancreas [16].

ARMS PCR is one of the most sensitive and reliable techniques being used to determine SNP analysis using Syber green in the present study. Two allele-specific (inner) primers are designed in opposite orientations and in combination with the outer primers, simultaneously amplify both the wild-type & mutant amplicons in single reaction to determine genotypes (homozygote/heterozygote) in disease condition such as carcinoma of pancreas. Using the intercalating fluorescence dye SYBR Green I and the Light Cycler system, the accumulation of amplicons in
the reaction can be monitored over time. After real-time PCR amplification, the Light Cycler continuously monitors the decrease in fluorescence resulting from the release of SYBR Green during DNA melting point analysis by slowly increasing the temperature. Because the allele-specific (inner) primers are designed in opposite orientations, they generate amplicons targeting opposite sides of the mutation point and the amplicons generally have different Tm (melting temperatures) depending on their GC content, length, and sequence. To verify our results we also run the PCR product on 3% agarose gel.

5. CONCLUSION

Multiple proceedings cause critical loss of gene activity and thereby predispose to cancer. How the normal cells become progressively transformed to malignant cell as a magnitude of damage to the genome in term of a gain, loss, or mutation of the genetic information? The finding of such small study is quite interesting because of using ARMS PCR to detect rare mutation of MTHFR 677TT allele and has important implications to assess “risk factor” for the development of carcinoma of pancreas in Indian population. However, further study is continued and required to collect more sample size to make the study significant.

CONSENT

It is not applicable.

ETHICAL APPROVAL

It is not applicable.

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COMPETING INTERESTS

Authors have declared that no competing interests exist between the authors.

REFERENCES


