

RESEARCH ARTICLE

Open Access

Novel missense mutation in the *RSPO4* gene in congenital hyponychia and evidence for a polymorphic initiation codon (p.M1I)

Tahir Naeem Khan¹, Joakim Klar², Sadia Nawaz¹, Muhammad Jameel¹, Muhammad Tariq¹, Naveed Altaf Malik¹, Shahid M Baig¹ and Niklas Dahl^{2*}

Abstract

Background: Anonychia/hyponychia congenita is a rare autosomal recessive developmental disorder characterized by the absence (anonychia) or hypoplasia (hyponuchia) of finger- and/or toenails frequently caused by mutations in the *R-spondin 4* (*RSPO4*) gene.

Methods: Three hypo/anonychia consanguineous Pakistani families were ascertained and genotyped using microsatellite markers spanning the *RSPO4* locus on chromosome 20p13. Mutation screening of the *RSPO4* gene was carried out by direct sequencing of the entire coding region and all intron-exon boundaries.

Results: Mutations in the *RSPO4* gene were identified in all families including a novel missense mutation c.178C>T (p.R60W) and two recurrent variants c.353G>A (p.C118Y) and c.3G>A (p.M1I). The c.3G>A variant was identified in unaffected family members and a control sample in a homozygous state.

Conclusions: This study raises to 17 the number of known *RSPO4* mutations and further expands the molecular repertoire causing hypo/anonychia. The c.353G>A emerges as a recurrent change with a possible founder effect in the Pakistani population. Our findings suggest that c.3G>A is not sufficient to cause the disorder and could be considered a polymorphism.

Keywords: Anonychia, Hyponychia, Mutation, *RSPO4* gene, Polymorphism

Background

The formation of nails is initiated in the upper limb followed by the development at the hindlimb and the morphogenesis is similar in primates and rodents [1]. This process is dependent on *RSPO4* as shown by mutations in the *RSPO4* gene in autosomal recessive anonychia/hyponychia (MIM 206800) [2-7]. *RSPO4* is a mediator of the highly conserved WNT signalling pathway and binds to the FZD family of receptors and the low-density lipo-protein related receptors (LRPs) 5 and 6 [8-10]. WNT signalling is critical for nail development and mutations in different WNT associated genes have been identified in disorders involving the nails. In addition to

RSPO4 mutations in anonychia/hyponychia, it was recently shown that mutations in *FZD6* cause isolated autosomal recessive nail dysplasia (MIM 614157) [11]. Furthermore, *WNT10A* (MIM 6062689) mutations are associated with odontoonychodermal dysplasia (OODD; MIM 257980) [12-14] and mutations in the WNT-associated transcription factors *LMX1B* (MIM 602575) and *MSX1* (MIM 142983) cause Nail-Patella (MIM 161200) and Witkop syndrome (MIM 189500), respectively [15,16].

We investigated three consanguineous families of Pakistani origin segregating autosomal recessive hyponychia of finger- and toenails. We report herein on three *RSPO4* missense mutations in a homozygous state of which one is novel. A second mutation is likely to be a founder mutation and a third variant, previously described as associated with anonychia, was identified in family members with normal nails as well as in a control individual.

* Correspondence: niklas.dahl@igp.uu.se

²Department of Immunology, Genetics and Pathology, The Rudbeck Laboratory and Science for Life Laboratory, Uppsala University, Uppsala 751 85, Sweden

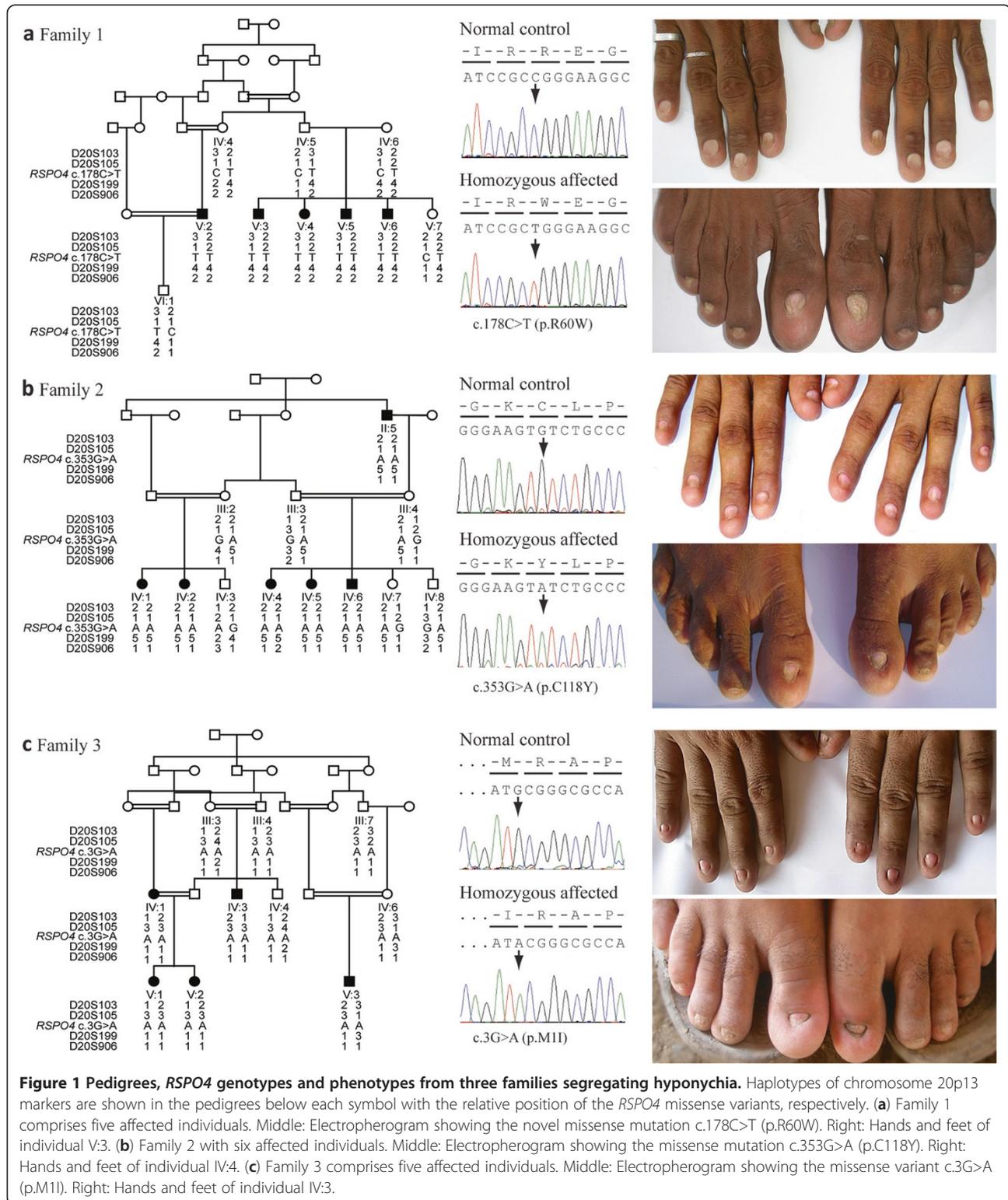
Full list of author information is available at the end of the article

Methods

Study subjects

Three consanguineous pedigrees of Pakistani origin with several members affected with hyponychia were investigated (Figure 1a-c). The nail phenotype ranged from

almost complete absence of some nail plates to reduced size of nail plates. Affected individuals had no other ectodermal abnormality including the teeth, skin and hair and they reported normal sweating. Informed consent was obtained from all individuals who participated in



this study under a protocol approved by the local ethical committee at National Institute for Biotechnology and Genetic Engineering (NIBGE), Faisalabad, Pakistan.

Genotyping

Blood samples from a total of 32 available members from the three families were obtained and DNA was extracted using standard protocols. All family members were initially genotyped using microsatellite markers D20S103, D20S105, D20S199 and D20S906 spanning the *RSPO4* gene on chromosome 20p13. Mutation screening of the *RSPO4* gene included all five exons and intron-exon boundaries using Big Dye Terminator v3.1 cycle sequencing kit (Applied Biosystems, Foster City, CA) and separated on an ABI 3730xl DNA analyzer (Applied Biosystems). Segregation of each mutation was confirmed by screening all available family members. Effects of missense variants were predicted using PolyPhen-2 (<http://genetics.bwh.harvard.edu/pph2/>).

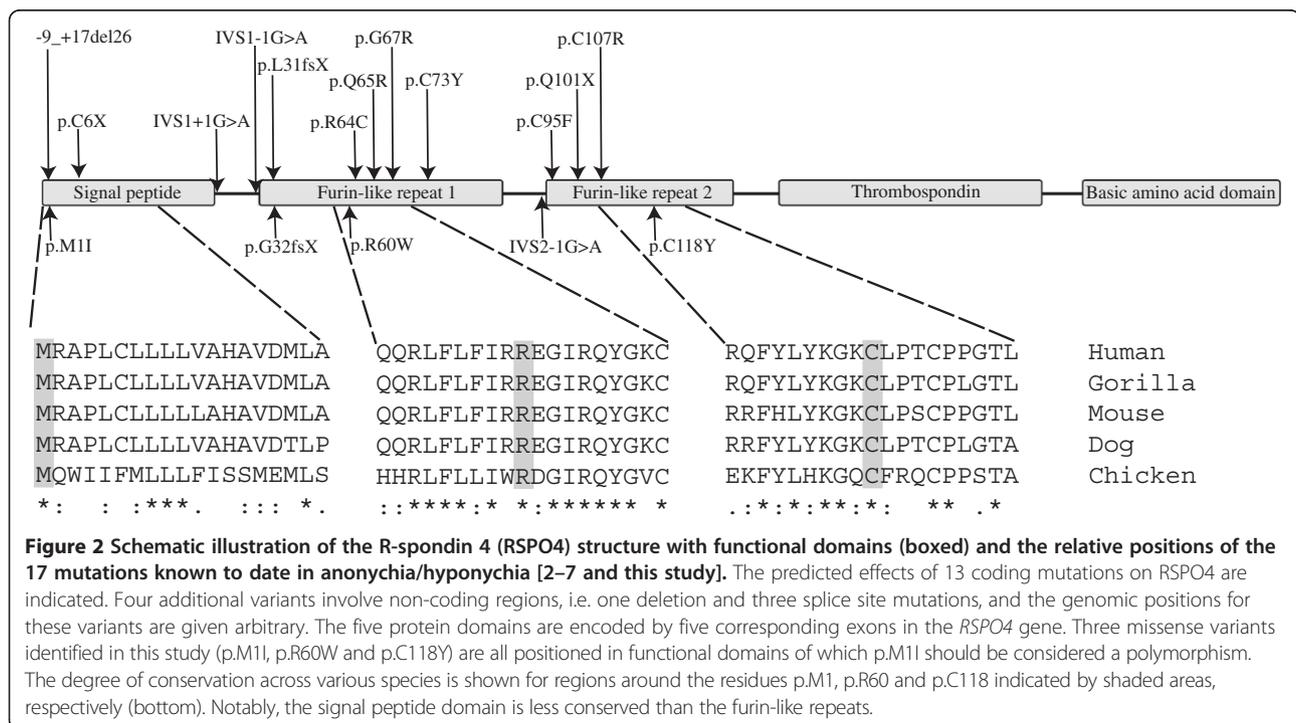
Results

Haplotype analysis and autozygosity mapping using microsatellite markers flanking *RSPO4* confirmed linkage to 20p13 in families 1 and 2 (Figure 1a-b). In family 3, the results suggested linkage (Figure 1c). Sequence analysis of the *RSPO4* gene in affected members of family 1 revealed a novel missense variant c.178C>T. The mutation was found in a homozygous state in all 5 affected individuals and in a heterozygous state in their parents (Figure 1a). The mutation is located in exon 2 and

results in a p.R60W substitution. The replacement for the neutral non-polar tryptophan is predicted to be damaging with a PolyPhen-2 score of 0.999. The mutation was excluded on 192 control chromosomes from 96 healthy Pakistani individuals. In family 2 we identified a missense variant c.353G>A resulting in a p.C118Y substitution in a homozygous state in all affected individuals (Figure 1b). The mutation is predicted to be damaging with a PolyPhen-2 score of 0.96. In family 3 we found the sequence variant c.3G>A resulting in p.M1I in five affected individuals as well as in five family members with normal finger- and toenails (Figure 1c). The amino-acid substitution is predicted as possibly pathogenic with a PolyPhen-2 score of 0.88 and both methionine and isoleucine belong to the group of neutral and non-polar amino acids. We screened 96 healthy Pakistani controls for this sequence variant and we identified one homozygous individual. This corresponds to an allele frequency of 1.1% for p.M1I. Non-coding variants in the *RSPO4* gene were excluded by sequencing the entire 5' and 3' UTRs as well as a predicted promoter region one kb upstream of the start codon in affected individuals.

Discussion

The majority of cases affected by isolated or non-syndromic congenital anonychia/hyponychia reported so far are caused by mutations in the *R-spondin 4* (*RSPO4*) gene [2-7]. *RSPO4* is expressed in mesenchymal cells from which the nails are derived [2] and the *RSPO4* protein act as a Frizzled (FZD) agonist in WNT signalling.



The importance of WNT signalling for nail development was further supported by the recent identification of *FZD6* mutations in autosomal recessive nail dysplasia [11].

In this study, we analysed three consanguineous pedigrees segregating hyponychia. In each family the affected members were found homozygous for a distinct *RSPO4* missense variant. In one family we found a novel mutation that results in a p.R60W substitution in the first furin-like domain associated with the disease. The finding is consistent with the suggestion that both furin-like domains are required for inducing β -catenin stabilization [5]. Furthermore, the positively charged arginine residue is highly conserved among different species (Figure 2). In a second family we identified a p.C118Y substitution in a homozygous state in affected individuals. This mutation has been reported previously in an independent Pakistani pedigree with hyponychia [2] suggesting a founder effect for the p.C118Y mutation in the Pakistani population. In a third family we identified a p.M1I variant in five affected individuals as well as in five members with a normal nail phenotype. Moreover, we also found the mutation in a homozygous state in one Pakistani control individual (1.1% allele frequency). Thus, the p.M1I variant, previously observed in two Pakistani siblings with onychia [5], is not sufficient to cause the disorder and may represent a rare polymorphism. We hypothesize that the *RSPO4* protein uses the second methionine codon at position 17 in the first exon for translational initiation. This will result in a shorter protein missing part of the signal peptide of importance for the secretory pathway [17]. However, the signal peptide is less conserved across species than the furin-like domains (Figure 2) and may not be required for normal nail development. This is supported by the fact that the furin-like domains of *RSPO4*, encoded after codon 17, are sufficient for inducing β -catenin stabilization [5].

With this report, the total number of *RSPO4* mutations associated with onychia/hyponychia is expanded to 17, all of which are located in the first three exons encoding a signal peptide and the highly conserved furin-like, cysteine-rich domains. Furthermore, we have identified a recurrent mutation in the Pakistani population and we suggest that the p.M1I variant could be considered a polymorphism as it is not sufficient to cause onychia/hyponychia.

Consent

Patients and their guardians provide written consent for publishing photographs and other material.

Conclusions

We have identified one novel and one recurrent missense mutation in the *RSO4* gene associated with congenital hyponychia. Furthermore, our findings indicate

that the *RSPO4* missense variant p.M1I is a polymorphism despite its previous association with onychia in two Pakistani siblings. We postulate that the common translational start for *RSPO4* may be replaced by the second methionine codon (p.M17). The combined results improve our current knowledge about *RSPO4* in nail development of both diagnostic and biological importance.

Abbreviations

RSPO4: R-spondin 4; NIBGE: National Institute for Biotechnology and Genetic Engineering; MIM: Mendelian inheritance in man; FZD: Frizzled; WNT10A: Wntless-type MMTV integration site family, member 10A; LMX1B: LIM homeobox transcription factor 1-beta; MSX1: Msh homeobox 1.

Competing interests

The authors declare that they have no competing interests.

Authors' contributions

TNK and JK carried out molecular genetic studies including autozygosity mapping and sequencing for two families and the controls and drafted the manuscript. SN, MJ, MT and NAM investigated the families, collected samples and extracted DNA. SMB and ND designed and supervised the study, analysed the data and finalized the manuscript. All authors read and approved the final manuscript.

Acknowledgements

We thank all family members who participated in this study. This work was supported by grants from the Swedish Research Council (K2010-66X-10829-17-3) and Swedish Links, Asia (348-2008-6069), The Swedish Institute, Uppsala University Hospital, Uppsala University and the Science for Life Laboratory. JK was supported by the Swedish Society for Medical Research.

Author details

¹Human Molecular Genetics Laboratory, Health Biotechnology Division, National Institute for Biotechnology and Genetic Engineering (NIBGE), Faisalabad 38000, Pakistan. ²Department of Immunology, Genetics and Pathology, The Rudbeck Laboratory and Science for Life Laboratory, Uppsala University, Uppsala 751 85, Sweden.

Received: 13 July 2012 Accepted: 10 December 2012

Published: 13 December 2012

References

1. Der Kaloustin VM, Kurban AK: *Genetic Diseases of the Skin*. Berlin: Springer Verlag; 1979.
2. Blaydon DC, Ishii Y, O'Toole EA, Unsworth HC, Teh M-T, Ruschendorf F, Sinclair C, Hopsu-Havu VK, Tidman N, Moss C, et al: **The gene encoding R-spondin 4 (RSPO4), a secreted protein implicated in Wnt signaling, is mutated in inherited onychia**. *Nat Genet* 2006, **38**:1245-1247.
3. Bergmann C, Senderek J, Anhuf D, Thiel CT, Ekici AB, Poblete-Gutiérrez P, van Steensel M, Seelow D, Nürnberg G, Schild HH, et al: **Mutations in the gene encoding the Wnt-signaling component R-spondin 4 (RSPO4) cause autosomal recessive onychia**. *Am J Hum Genet* 2006, **79**:1105-1109.
4. Bruchle NO, Frank J, Frank V, Senderek J, Akar A, Koc E, Rigopoulos D, van Steensel M, Zerres K, Bergmann C: **RSPO4 is the major gene in autosomal recessive onychia and mutations cluster in the furin-like cysteine-rich domains of the Wnt Signaling Ligand R-Spondin 4**. *J Invest Dermatol* 2007, **128**:791-796.
5. Ishii Y, Wajid M, Bazzi H, Fantauzzo KA, Barber AG, Blaydon DC, Nam J-S, Yoon JK, Kelsell DP, Christiano AM: **Mutations in R-Spondin 4 (RSPO4) Underlie Inherited Onychia**. *J Invest Dermatol* 2007, **128**:867-870.
6. Chishti MS, Kausar N, Rafiq MA, Amin M, Ahmad W: **A novel missense mutation in RSPO4 gene underlies autosomal recessive congenital onychia in a consanguineous Pakistani family**. *Br J Derm* 2008, **158**:621-623.
7. Wasif N, Ahmad W: **A Novel Nonsense Mutation in RSPO4 Gene Underlies Autosomal Recessive Congenital Onychia in a Pakistani Family**. *Ped Derm* 2012, [Epub ahead of print].

8. Kazanskaya O, Glinka A, del Barco Barrantes I, Stannek P, Niehrs C, Wu W: **R-Spondin2 is a secreted activator of Wnt/ β -catenin signaling and is required for xenopus myogenesis.** *Dev Cell* 2004, **7**:525–534.
9. Kim K-A, Zhao J, Andarmani S, Kakitani M, Oshima T, Binnerts ME, Abo A, Tomizuka K, Funk WD: **R-Spondin Proteins: a novel link to β -catenin activation.** *Cell Cycle* 2006, **5**:23–26.
10. Nam J-S, Turcotte TJ, Smith PF, Choi S, Yoon JK: **Mouse cristin/R-spondin family proteins are novel ligands for the frizzled 8 and LRP6 receptors and activate β -catenin-dependent gene expression.** *J Biol Chem* 2006, **281**:13247–13257.
11. Fröjmark A-S, Schuster J, Sobol M, Entesarian M, Kilander Michaela BC, Gabrikova D, Nawaz S, Baig Shahid M, Schulte G, Klar J, *et al*: **Mutations in frizzled 6 cause isolated autosomal-recessive nail dysplasia.** *Am J Hum Genet* 2011, **88**:852–860.
12. Adaimy L, Chouery E, Mégarbané H, Mroueh S, Delague V, Nicolas E, Belguith H, de Mazancourt P, Mégarbané A: **Mutation in WNT10A is associated with an autosomal recessive ectodermal dysplasia: the Odonto-Onycho-Dermal Dysplasia.** *Am J Hum Genet* 2007, **81**:821–828.
13. Nawaz S, Klar J, Wajid M, Aslam M, Tariq M, Schuster J, Baig SM, Dahl N: **WNT10A missense mutation associated with a complete Odonto-Onycho-Dermal Dysplasia syndrome.** *Eur J Hum Genet* 2009, **17**:1600–1605.
14. Bohring A, Stamm T, Spaich C, Haase C, Spree K, Hehr U, Hoffmann M, Ledig S, Sel S, Wieacker P, *et al*: **WNT10A mutations are a frequent cause of a broad spectrum of ectodermal dysplasias with sex-biased manifestation pattern in heterozygotes.** *Am J Hum Genet* 2009, **85**:97–105.
15. Dreyer SD, Zhou G, Baldini A, Winterpacht A, Zabel B, Cole W, Johnson RL, Lee B: **Mutations in LMX1B cause abnormal skeletal patterning and renal dysplasia in nail patella syndrome.** *Nat Genet* 1998, **19**:47–50.
16. Jumlongras D, Bei M, Stimson JM, Wang W-F, DePalma SR, Seidman CE, Felbor U, Maas R, Seidman JG, Olsen BR: **A nonsense mutation in MSX1 causes Witkop Syndrome.** *Am J Hum Genet* 2001, **69**:67–74.
17. de Lau W, Snel B, Clevers H: **The R-spondin protein family.** *Genome Biol* 2012, **13**:242.

doi:10.1186/1471-2350-13-120

Cite this article as: Khan *et al*: Novel missense mutation in the *RSPO4* gene in congenital hyponychia and evidence for a polymorphic initiation codon (p.M11). *BMC Medical Genetics* 2012 **13**:120.

Submit your next manuscript to BioMed Central and take full advantage of:

- Convenient online submission
- Thorough peer review
- No space constraints or color figure charges
- Immediate publication on acceptance
- Inclusion in PubMed, CAS, Scopus and Google Scholar
- Research which is freely available for redistribution

Submit your manuscript at
www.biomedcentral.com/submit

