Special Issue on Gene Defects and Genetic Syndromes

Call for Papers

A syndrome is a set of medical signs and symptoms that are correlated with each other and, often, with a specific disease. In some instances a syndrome is so closely correlated with a pathogenesis or etiology that the words syndrome, disease, and disorder end up being used interchangeably for them. This is especially true of genetically caused syndromes. For example, Down syndrome, Wolf–Hirschhorn syndrome, and Andersen syndrome are disorders with known pathogeneses, so each is more than just a set of signs and symptoms, despite the syndrome nomenclature.

In this special issue, we intend to invite front-line researchers and authors to submit original researches and review articles on exploring gene defects and genetic syndromes. Potential topics include, but are not limited to:

- Autosomal disorder
- X-linked and Y-linked disorders
- Albinism
- Down's syndrome
- Huntington's disease
- Marfan syndrome
- Roberts syndrome
- Mitochondrial disease

Authors should read over the journal’s Authors’ Guidelines carefully before submission. Prospective authors should submit an electronic copy of their complete manuscript through the journal at Paper Submission System.

Please kindly specify the “Special Issue” under your manuscript title. The research field “Special Issue – Gene Defects and Genetic Syndromes” should be selected during your submission.

Special Issue timetable:

<table>
<thead>
<tr>
<th>Submission Deadline</th>
<th>September 20th, 2016</th>
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<tr>
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Guest Editor:

For further questions or inquiries
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