Comparison of in-vitro and in-vivo response to fetal hemoglobin production and γ-mRNA expression by hydroxyurea in Hemoglobinopathies

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BACKGROUND: Hydroxyurea, which induces Fetal hemoglobin (HbF) synthesis, is the only drug widely used in different hemoglobinopathies; however, the response is very variable. We compared the efficacy of hydroxyurea in-vitro in erythroid cultures and in-vivo in the same patients with different hemoglobinopathies to induce HbF production and enhance γ-messenger RNA expression.

MATERIALS AND METHODS: A total of 24-patients with different Hemoglobinopathies were given hydroxyurea and their response was studied in-vivo and in-vitro on mononuclear cells collected from them simultaneously.

RESULTS: A total of 57.7% of patients (responders) showed no further crisis or transfusion requirements after hydroxyurea therapy with a mean increase in fetal cells (F-cells) of 63.8 ± 59.1% and γ-mRNA expression of 205.5 ± 120.8%. In-vitro results also showed a mean increase in F-cells of 27.2 ± 24.7% and γ-mRNA expression of 119.6% ± 65.4% among the treated cells. Nearly 19.0% of the partial-responders reduced their transfusion requirements by 50% with a mean increase in F-cells of 61.2 ± 25.0% and 28.4 ± 25.3% and γ-mRNA-expression of 21.0% ± 1.4% and 80.0% ± 14.1% in-vitro and in-vivo respectively. The non-responders (15.3%) showed no change in their clinical status and there was no significant increase in F-cells levels and γ-mRNA expression in-vivo or in-vitro.

CONCLUSION: Thus, this method may help to predict the in-vivo response to hydroxyurea therapy; however, a much larger study is required.

Key words: 2 phase liquid erythroid cell culture, hydroxyurea, sickle cell disease, β-thalassemia syndromes, γ-mRNA expression

Introduction

Inherited disorders of hemoglobin, such as the structural Hemoglobinopathies, and the thalassemia syndromes are common, affecting around 7% of the global population with significant clinical conditions.[1] In India, the prevalence of sickle cell carriers ranges from 2% to 34% in different population groups[2,3] whereas that of β-thalassemia ranges from 1% to 17%.[4,5] On the other hand, Hemoglobin E (HbE) is generally found in the eastern and north eastern region of the country with a frequency of 3-10% and 4-51% respectively.[6]

Although, sickle cell disease has a milder clinical presentation than in Africans among tribal groups in India, many patients particularly from the non-tribal groups suffer from severe forms of the disease leading to repeated vaso-occlusive crisis, severe anemia, and end organ damage.[7,8] So far, mainly supportive measures have been used for decreasing the severity of symptoms. Few children with the severe forms of β-thalassemia receive regular blood transfusions and adequate iron chelation, the mainstay of treatment, due to the high-cost involved for optimum management.[9]

Thus, there was a need for an efficacious drug with low
toxicity, which could reduce the clinical symptoms of these Hemoglobinopathies in a cost-effective way.

Hydroxyurea reduces the frequency of vaso-occlusive crisis and blood transfusion requirements in sickle cell anemia patients[8,9] as well as the transfusion requirements in some β-thalassemia patients.[10,11] As there are no clear clinical or genetic factors, which can predict the response of these patients to hydroxyurea therapy, patients who are non-responders also get exposed to the drug. Thus, this study thus investigated whether the response to hydroxyurea in erythroid cultures of patients with different Hemoglobinopathies to increase fetal hemoglobin (HbF) and γ-gene expression in-vitro correlates simultaneously with the in-vivo response to hydroxyurea among the same patients.

Materials and Methods

Out of 24 patients (8 males and 16 females; age 5-30 years) with different Hemoglobinopathies (severe sickle cell anemia patients – 6; severe sickle-β-thalassemia patients – 2, these patients had more than 5 episodes of painful vaso-occlusive crisis and occasional blood transfusion requirements; β-thalassemia intermedia patients – 5, who presented after 2 years of age and required occasional blood transfusions initially however, later they became transfusion dependent; β-thalassemia major patients – 5, who were on regular blood transfusion; severe HbE-β-thalassemia patients – 6, who had 6-12 transfusion requirements annually) were included. Informed consent was taken from the patients and parents of pediatric patients and the Ethical Committee of the National Institute of Immunohematology approved these studies.

In-vivo studies

Hydroxyurea (Cytodrox, Cipla Ltd, Mumbai) was started at a dose of 10-15 mg/kg/day[8,10] after initial blood collection for in-vitro studies, hematological investigations like complete blood count, hemoglobin analysis on the Variant Hemoglobin Testing System (BioRad Laboratories, Inc., Hercules, CA, USA) and quantitative estimation of fetal cells (F-cells) by flow cytometry using a monoclonal HbF antibody (BD (Becton Dickinson and Company) Immunocytometry Systems, San Jose, CA, USA).[12] Molecular analysis such as confirmation of the HbS (sickle hemoglobin) and HbE mutations and characterization of β-thalassemia mutations were carried out as described earlier.[13] Xmn I polymorphism analysis was carried out by PCR (polymerase chain reaction) and restriction enzyme digestion.[14] Absolute quantification of gamma (γ)-globin mRNA transcripts was carried out by real time PCR using Taqman probes and the 7900 High Temperature Fast Real-Time PCR System (Applied Biosystems, NJ, USA) as described earlier.[15] The expression of the housekeeping gene β-actin was used as control in each sample. Patients were assessed clinically and laboratory investigations carried out monthly for 24 months to evaluate their status after starting hydroxyurea therapy. The numbers of remaining capsules of hydroxyurea were counted during each follow-up to evaluate therapeutic compliance and the hematological status was monitored to rule-out cytopenia. At the end of 24 months the patients’ response to hydroxyurea therapy was evaluated by the reduction in clinical severity of the disease in case of sickle cell disease patients and reduction or cessation of transfusion requirements in β-thalassemia and HbE-β-thalassemia patients. After starting hydroxyurea therapy, the β-thalassemia patients were transfused only if their hemoglobin dropped below 7.5 g/dl. Partial responders of hydroxyurea therapy among the β-thalassemia were those patients whose transfusion requirement reduced by 50% whereas non-responders were those patients who did not show any reduction in transfusion requirements after 1 year of hydroxyurea therapy.

In-vitro studies

In-vitro studies were performed using the two phase liquid culture technique.[16,17] Mononuclear cells, collected from peripheral blood of all patients before starting hydroxyurea were cultured in phase-I medium containing serum free StemSpan medium (StemCell Technologies Inc.), 50 ng/ml of stem cell factor (SCF), 25 ng/ml of interleukin-3 (IL3), 0.01% bovine serum albumin (BSA) and Cyclosporin A (1 μg/ml) (Sigma-Aldrich Co. USA) for 7 days at 37°C and 5% CO₂. Non-adherent cells after day-7 were further cultured in phase-II medium consisting of StemSpan medium, 2 U/ml of human recombinant erythropoietin,
50 ng/ml SCF and 10⁻⁷ M Dexamethazone (Sigma-Aldrich Co. USA). The culture was then bifurcated with a cell concentration of less than 1 × 10⁶/ml and hydroxyurea (1.5 mM/1 × 10⁶ cells) was added to one of the cultures between days 6 and 8 of phase-II. After 10-12 days the cells were collected for F-cells estimation and γ-mRNA expression as described above.

**Flow cytometry**

For F-cells estimation, the cultured cells were washed twice and blocked with 2% BSA and stained with cell surface markers like anti-human-CD45 (leukocyte common antigen) tagged with perCP (Peridinin Chlorophyll Protein Complex) (10 μl) and anti-human-CD71 (transferrin receptor) tagged with Phycoerythrin (10 μl) and incubated in the dark for 30 min. After washing, the fluorescently labeled cells were fixed, permeabilized and stained with anti-HbF-FITC (Fluorescein isothiocyanate) as mentioned earlier. Cultured cells, which were CD45 negative were gated (10,000 cells) and used for flow cytometric analysis to quantitate the number of F-cells.

**Statistical analysis**

Student t-test was used to compare the findings of *in-vitro* studies with and without hydroxyurea and of patients before and after hydroxyurea therapy. A *P* < 0.01 was considered statistically significant.

**Results**

**Clinical and hematological response to hydroxyurea therapy in-vivo**

After hydroxyurea therapy, 8 patients, which include 6 sickle cell anemia patients and 2 sickle-β-thalassemia patients had no further episodes of vaso-occlusive crisis, need for blood transfusions, infections, stroke or acute chest syndrome and they had a feeling of general well-being after therapy. The mean hematological parameters of these 8 patients also showed a significant increase in Hb, MCV (Mean corpuscular volume), HbF, and F-cells (*P* < 0.001) [Table 1] after hydroxyurea therapy.

**Response to hydroxyurea therapy in-vitro**

The erythroid cells of the patients were cultured for an average of 18.5 ± 4 days. Erythroblasts increased dramatically from day 1 to 4 with a decrease in cell volume from days 5 to 7. Orthochromatic normoblasts
representing the late erythroblastic cells stage rarely lead to enucleation under the culture conditions. Hydroxyurea added on day 7 ± 1 in one of the duplicate sets of culture to evaluate the F-cells response was titrated and a dose of 1.5 mM/1x10^6 cells was found to be optimum for the cells. A concentration above 1.5 mM reduced the cell number and the expression of CD71. Figure 1 shows the flow cytometric data of cultured cells showing co-positivity for F-cells and CD71 before and after addition of hydroxyurea.

The percent increase in F-cells, F-cells + CD71 and γ-mRNA expression of individual patient's cultured erythroid cells after treating with hydroxyurea is shown in Table 2. The difference in F-cells + CD71 and γ-mRNA expression seen among the responders and partial-responders was not statistically significant [Table 3]. 4 non-responders in-vivo, did not show any increase in F-cells in their cultured cells treated with hydroxyurea. Only 1 of the 4 non-responders (β-thalassemia major patient) showed an increase in F-cells + CD71 by 1% after hydroxyurea treatment, whereas the other 3 patients did not show any increase. No increase was found in the γ-mRNA expression carried out in 2 of the 4 non-responders' cultured erythroid cells after hydroxyurea treatment.

discussion

Many pharmacological agents have been studied to enhance the production of HbF in various in-vivo and in-vitro systems. Use of these drugs on non-human primates are a pre-requisite for clinical trials in patients, however, they are too expensive and time consuming for large scale screening of these compounds. Hence, attempts have been made to mimic the in-vivo condition in-vitro by various experimental systems using either immortalized cell lines or isolation and culture of erythroid progenitor cells from the bone marrow or peripheral blood. Fibach et al. in 1989 established primary erythroid cultures by isolating erythroid progenitor cells from the bone marrow, cord blood or peripheral blood. Fibach et al. in 1989 established primary erythroid cultures by isolating erythroid progenitor cells from the bone marrow, cord blood or peripheral blood, which when grown in-vitro resembles more closely the in-vivo situation.[18] Fibach in 1998 studied many pharmacological inducers of HbF like hydroxyurea and 5-azacytidine using peripheral blood of sickle cell anemia patients with this technique.[16] Treatment with hydroxyurea resulted in reduction in cell number, increase in MCV, MCH and HbF. It was further suggested that this technique could be used to predict the responders of hydroxyurea.
In Table 2: Comparison of the percent increase in F-cells, CD71+F-cells and γ-mRNA expression of cultured erythroid cells (in-vitro) with the in-vivo percent increase in F-cells and mRNA expression in few patients before and after hydroxyurea therapy, the data is presented in a tabular format. The table includes columns for Patient Type, Mutation, Xmn I Polymorphism, Alpha Thalassemia, In-vitro % increase in F-cells, CD71+F-cells, % increase in γ-mRNA, In-vivo % increase in F-cells, % increase in γ-mRNA, and Clinical response. The table details the percent increase in F-cells and γ-mRNA expression for various mutations and patient responses to hydroxyurea therapy.

In 2002, Wojda et al. found that HbA and HbF content were regulated with the rate of proliferation during adult erythropoiesis and that there was no evidence for a HbF dominant population or switching during differentiation in adult cells. Wojda et al. in 2003 showed significant SCF mediated increase in HbF using progenitor cells derived from donor cells. Mithramycin and rapamycin were shown to induce HbF production by up regulating γ-globin mRNA production in normal and thalassemia human erythroid precursor cells using the two phase liquid culture technique. Human erythroid progenitor cells isolated from 18 HbE-β-thalassemia patients...
patients with different Hemoglobinopathies gives an idea about the parameters that may predict the response to hydroxyurea therapy among these patients.

Hydroxyurea was found to be beneficial both among adults and children with severe form of sickle cell disease\[8,26-28\] and was further used in the treatment of patients with β-thalassemia and HbE-β-thalassemia where a good response was seen among a few β-thalassemia intermedia and HbE-β-thalassemia patients in whom many genetic and epigenetic factors are believed to play a role in predicting the response to hydroxyurea therapy.\[10,11,25,29-32\] Watanapokasin et al. 2005 showed that 13 β-thalassemia/HbE-β-thalassemia patients treated with hydroxyurea in-vivo had HbF synthesis and γ-globin-mRNA that correlates with the in-vivo results suggesting that in-vitro testing may predict the in-vivo response to hydroxyurea. Bianchi et al. 2007 used 3 inducers of HbF from the biological materials, which included angelicin linear psoralens, resveratrol and rapamycin. A good correlation was found between in-vitro and in-vivo response to HbF production.\[33\]

In our study, a significantly higher expression of γ-mRNA ($P < 0.01$) was seen among the cultures of responders treated with hydroxyurea as compared to γ-mRNA expression seen among partial-responders. The non-responders did not show any increase in F-cells or γ-mRNA expression in-vitro, however, they showed a 16.0 ± 6.1% increase in F-cells but only a 2.9 ± 0.8% increase in γ-mRNA expression in-vivo after therapy. Thus γ-mRNA expression was a better indicator of the response to hydroxyurea therapy both in-vivo and in-vitro as compared to the F-cells level as the F-cells showed an increase in most of the patients and their cultured erythrocytes treated with hydroxyurea, however, the percentage increase in F-cells levels differed among the responders and the non-responders.

### Table 3: Percent increase in F-cells, CD71+ F-cells and γ-mRNA expression in-vitro and in-vivo among the responders, partial-responders and non-responders of hydroxyurea therapy

<table>
<thead>
<tr>
<th>Response</th>
<th>In-vitro</th>
<th></th>
<th>In-vivo</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>% increase in F-cells mean±SD</td>
<td>% increase in CD71+F-cells mean±SD</td>
<td>% increase in γ-mRNA expression mean±SD</td>
<td>% increase in F-cells mean±SD</td>
</tr>
<tr>
<td>Responders</td>
<td>14.2±11.0</td>
<td>27.2±24.7</td>
<td>119.6±65.4</td>
<td>63.8±59.1</td>
</tr>
<tr>
<td>Partial-responders</td>
<td>5.5±5.3</td>
<td>28.2±25.3</td>
<td>80.0±14.1</td>
<td>61.2±25.0</td>
</tr>
<tr>
<td>Non-responders</td>
<td>No increase</td>
<td>1.0</td>
<td>No increase</td>
<td>16.0±6.1</td>
</tr>
<tr>
<td>Responders versus partial-responders</td>
<td>$P&lt;0.01$</td>
<td>$P&lt;0.1$</td>
<td>$P&lt;0.1$</td>
<td>$P&lt;0.1$</td>
</tr>
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F-cells: Fetal cells

![Figure 1: Flow cytometric estimation of CD71 and fetal cells on cultured erythroid cells before and after addition of hydroxyurea](image)

In this study, the response to hydroxyurea is compared in-vivo as well as in-vitro. Our earlier study on the use of hydroxyurea among sickle cell disease patients in India has shown the drug to be very effective in reducing clinical crisis,\[11\] however, only 60% of the β-thalassemia intermedia and 50% of HbE-β-thalassemia patients fully benefited from the drug.\[12,25\] The in-vitro study on the effect of hydroxyurea on the cultured erythrocytes of
This method of erythroid cell culture may be useful to compare the in-vivo response of hydroxyurea with in-vitro using γ-mRNA expression and the number of F-cells. However, a much larger study is required to confirm these results. This culture system could be further used to identify other pharmacological inducers of HbF production, which could be useful for non-responders of hydroxyurea and to study expression of other genes that may be involved in HbF production.

References


Source of Support: Nil, Conflict of Interest: None declared.