Tuberculosis (TB) caused by Mycobacterium tuberculosis (Mtb) has become the world’s leading killer disease due to a single infectious agent which survives in the host macrophage for the indefinite period. Hence, it is necessary to enhance the efficacy of the clinically existing antitubercular agents or to discover new anti-antitubercular agents. Here, we report the synthesis, characterization and antimycobacterial activity of protein-drug conjugates. A carrier protein, Transferrin (Tf) was covalently conjugated to isoniazid (INH) utilizing hydrazone and amide linkers. The purity of the reactions was confirmed by SDS-PAGE while conjugation was confirmed by UV—visible spectrophotometry, MALDI-TOF analysis, and FTIR spectrophotometry. The in vitro antitubercular assay result showed that the inhibitory activity of the conjugates was 100 times effective against intracellular Mtb H37Rv and were devoid of cytotoxic effect at therapeutic concentration.
delivery system for tuberculosis therapy [12]. The use of plasma protein transferrin as a carrier moiety is widely known for anticancer drug delivery. One can also introduce it in the delivery of antitubercular drug because of expression of transferrin receptor (TfR) on infected macrophages along with simple, reproducible and well characterized covalent conjugation of protein with the drug.

Human plasma transferrin is a nonheme iron-binding glycoprotein [molecular mass 80 kDa] [13]. The primary role of Tf is to control the levels of free iron in body fluids by binding, sequestering and transporting Fe⁺⁺ ions [14–16]. Mtb infects macrophages and survives on host iron. Mtb acquire iron from the host cell by synthesizing siderophores (carboxymycobactin & mycobactin) and also from haem sources [17–19]. Besides, host iron carrier proteins Tf and lactoferrin are actively recruited by this pathogen to the phagosomal compartment for the acquisition of iron [20–23]. The TfR and specific receptor appear to be expressed on macrophages and mycobacteria for the binding of Tf on it [20]. A recent study shows that Mtb acquires iron from Tf by three mechanisms. The first is siderophores dependent withdrawal and intracellular delivery of iron. The second mechanism involves the delivery of iron using the high-affinity transporter IrtAB by a mycobactin-independent pathway. The third mechanism of transferrin-iron acquisition is independent of siderophores and consists of the internalization of transferrin by GAPDH receptor [24]. Thus expression of receptors, ability to internalize, and necessity of iron for the survival of mycobacteria make this protein an extensively accessible carrier for the receptor-mediated delivery of the drug into the infected macrophages.

The present work reports the synthesis of isoniazid-transferrin conjugates, its characterization and in vitro antimycobacterial activity. To the best of our knowledge, this is the first report on the conjugation of an antitubercular agent with human plasma protein transferrin. Our primary aim was to utilize simple and readily available linkers and to study its influence on Tf-INH conjugation.

2. Materials and methods

2.1. Materials

Transferrin (Holo) from human plasma (>95%, SDS-PAGE) purchased from Sigma Aldrich St. Louis, MO, USA. Isoniazid was obtained as a gift sample from Loba Chemie Ltd., glyoxylic acid monohydrate and glutaric anhydride, 2,4,6-trinitrobenzene-1-sulfonic acid (TNBS), 1-Ethyl-3-[3-dimethylaminopropyl] carbodiimide hydrochloride (EDC), H-hydroxy succinimide (NHS), Sodium chloride, potassium dihydrogen phosphate (KH₂PO₄), disodium hydrogen phosphate (Na₂HPO₄), potassium bromide were purchased from Sigma Aldrich St. Louis, MO, USA. Acetonitrile, dimethyl sulphoxide (DMSO) were obtained from Merck.

2.2. Synthesis of isonicotinohydrazonoacetic acid (IH) (2)

The aqueous solution of glyoxylic acid monohydrate (0.92 g) was prepared and INH (1.37 g) dissolved in acetonitrile/water (1:4 v/v) was added to it. After completion of the reaction, the precipitate was filtered and washed with water and dried. The spectroscopic data is summarized in supporting information.

2.3. Synthesis of isonicotinoylhydrazinyl-5-oxopentanoic acid (IG) (3)

INH (1.37 g) was dissolved in acetonitrile/water (1:4 v/v) and added to the aqueous solution of glutaric anhydride (1.14 g). After completion of the reaction, the precipitate was filtered and washed with water and dried. The spectroscopic data is summarized in supporting information.

2.4. Conjugation of isonicotinoylhydrazonoacetic acid (4) and isonicotinoylhydrazinyl-5-oxopentanoic acid (5) to transferrin

Conjugates of isonicotinohydrazonoacetic acid (IH) or isonicotinoylhydrazinyl-5-oxopentanoic acid (IG) with Tf at different molar ratios (TF: IH/IG; 1:20, 1:40, 1:60, 1:100) were prepared. IH/IG was added in PBS (pH 7.4) followed by EDC and NHS (1.5:3 mol). The solution was allowed to stir for 2 h for the activation of acid groups. After complete consumption of IH/IG, Tf [dissolved in PBS (pH 7.4)] was added dropwise in the reaction mixture with stirring. The reaction was carried out for 12 h at 30 °C. The mixture was then dialyzed against deionized water at 4 °C for 48 h using a dialysis membrane (MWCO: 12–14 kDa) to remove salt, unreacted drug intermediates, and urea by-products. Finally, the product was lyophilized. The purity of synthesized forms of conjugates was determined using SDS-PAGE on 10% polyacrylamide gel at 120 V. The samples were prepared under non-reducing condition. The staining of gel and visualization of bands was done with 0.25% Coomassie blue stain using Bio-Rad.

The available amino groups present on Tf before and after conjugation was determined by the 2,4,6-trinitrobenzene-1-sulfonic acid (TNBS) reagent [25–27]. Briefly, an amount of 4 ml of Tf or Tf conjugate (0.25 mg/ml) in the aqueous buffer of 0.1 N NaHCO₃ (pH 8.3) was mixed with 2 ml of 0.01% TNBS. The mixture was incubated at 37 °C for 4 h. Further, 1 ml of 1N HCl and 2 ml of 10% sodium dodecyl sulfate (SDS) were added to each sample and mixed. The absorbance of the sample was measured at 340 nm using a UV/Vis spectrophotometer (Shimadzu UV-2600). The calibration curve was plotted by using l-lysine as a standard (Fig. S4). The concentration of free primary amino groups on Tf and Tf conjugate was estimated from the calibration curve.

2.5. UV-visible spectroscopy

UV–visible spectrophotometric analysis of samples was performed for the confirmation of TF-INH conjugation on UV–visible spectrophotometer (Shimadzu UV-2600) in a 1 cm cell. The absorbance spectra of INH, Tf, Tf-IG, and Tf-IG were measured in phosphate buffer solution (pH 7.4), and absorbance maximum was measured.

2.6. MALDI-TOF measurement

Mass spectrometry (MS) was used to determine the molecular weight of prepared IG/IH-transferrin conjugate. The mass spectra of Tf and its conjugates were evaluated, and the difference between masses of both the compounds was used to determine the molar ratio of drug conjugated to a protein. Matrix-assisted laser desorption/ionization time of flight (MALDI-TOF) mass spectrometry (Ultraflex™, Bruker Daltonics, Bremen, Germany) was used in a linear mode to determine the molecular weight of the compounds. The solutions of transferrin and transferrin conjugates were prepared at 10 μg/ml of concentration. A saturated solution of matrix-sinapic acid (SA) in 50% acetonitrile and 0.05% trifluoroacetic acid was prepared, and protein or conjugate was mixed into it in a volume ratio of 1:1. Further, 0.5 μl of the sample was applied to a steel plate.

2.7. In vitro antibacterial activity

The antitubercular activity of INH, INH—H, INH—G, Tf-IG, and Tf-IG was determined using the Microplate Alamar Blue assay (MABA) [28]. MABA assay was performed in 96-well flat bottom plates. Mtb strains were cultured in 7H9-ADS medium and grown till the exponential phase (OD₆₀₀ of 0.6). Approximately 1 × 10⁵ bacteria
were taken per well in a total volume of 200 μl of the 7H9-ADS medium. Wells containing no Mtb were used for autofluorescence control. Additional controls consisted of wells containing cells and medium only. Plates were incubated for 5 days at 37 °C, 30 μl (0.02% wt/vol stock solution) Alamar Blue was added. Plates were re-incubated for color transformation (blue to pink). Fluorescence intensity was measured in a SpectraMax M3 plate reader (Molecular Device) in a top-reading mode with excitation at 530 nm and emission at 590 nm. Percentage inhibition was calculated based on the relative fluorescence units, and the minimum concentration that resulted in at least 90% inhibition was identified as MIC.

2.8. In vitro cytotoxicity study

The cytotoxicity of INH, TF-IH, and TF-IG was tested in THP-1 differentiated macrophages using MTT assay. Cells were allowed to adhere for 24 h in a 96-well plate at a density of 3000 cells/well prior to the assay. Further, cells were exposed to various concentrations of 0.9% saline (the control), INH, TF-IH, and TF-IG at various concentrations for 48 h at 37 °C and 5% CO₂ atmosphere, respectively. Culture medium was used as the blank group. After that, the MTT solution was added to each well and cells were incubated for 4 h. To dissolve the MTT formazan crystals, 200 μl of DMSO was added to each well. A microplate reader (Model 680, BIO-RAD, Hercules, CA) was used to measure the optical density of the formazan product at 570 nm. The drug concentration causing 50% inhibition (IC₅₀) was calculated.

2.9. Intracellular killing of Mtb

20000 THP-1 cells in 100 μl media (RPMI + 10% FBS) with 20 ng/ml phorbol 12-myristate 13-acetate (PMA) per well were seeded in flat with adhesive 96 well plate. After that, the plate was incubated for 24 h in 5% CO₂ incubator. After 24 h of incubation cells were washed with PBS and added fresh media and further incubated for 24–48 h, and after incubation cell were infected with H37Rv at multiplicity of infection (MOI) (20000 THP-1: 40000 bacilli) and left for 4 h, washed with PBS and were added fresh media with Amikacin – 0.2 mg/ml and incubated for 1 h, after that 3 times washing was done and cells were incubated with appropriate concentration of drugs, and finally cells were plated at two time points for determining the colony forming unit (CFU), first at 0 days before treatment and second after 48 h [29].

2.10. Stability study

The stability of hydrazone and amide bond containing INH conjugates was analyzed by HPLC sampling over 24 h. INH, TF-IH/IG (T3) were dissolved in PBS 4.0 and PBS 7.4 separately at 0.5 mg/ml concentration. 20 μl of sample was injected after 2, 6, 8, 12, 18, 24 h to 250 × 4.6 mm I.D., 5 μm particle, Phenomenex Luna C18 column. Gradient elution mode was applied using acetonitrile (A) and 15 mmol/L potassium dihydrogen phosphate buffer of pH adjusted to 4.0 ± 0.1 with orthophosphoric acid (B) as a mobile phase (A:B ratio 50:50 v/v).

2.11. Statistical analysis

Cytotoxicity study and intracellular killing assay were performed in triplicate, and the results are expressed as mean ± standard deviation (SD). Statistical analysis was done by performing Student’s t-test. A value of p < 0.05 was considered significant (n = 3).

3. Result and discussion

Transferin was conjugated to isonicotinoylhydrazono acetic acid (IH) forming hydrazone linkage and to 5-(2-sionicotinoylhydrazinyl)-5-oxopentanoic acid (IG) forming amide linkage using EDC-NHS coupling reaction. The –COOH groups of IH and IG were activated by EDC and NHS to react with primary amino groups of lysine of TF (Scheme 1).

The SDS-PAGE shows the single dark prominent band at ~79 kDa for all TF, TF-IH and TF-IG samples (Fig. 1). The absence of any smaller bands in SDS-PAGE confirms that there was no protein-protein coupling induced during the EDC-NHS coupling procedure. The absence of any smaller bands in the TF conjugates in SDS-PAGE suggests there was no degradation of the TF during the conjugation process.

The absorbance spectrum of free TF, TF-IH, TF-IG, and INH dissolved in PBS pH 7.4 is given in Fig. 2. The free TF showed an excitation wavelength at 279 nm while conjugated TF shifts its wavelength at 274 nm indicating a blue shift and confirms the conjugation on TF.

The concentration of surface lysine present in TF before and after conjugation was determined by TNBS assay. The gradual decrease in the free amino group with the increase in the molar ratio of TF-IH/IG confirms the covalent attachment of IH/IG (carboxyl group) to transferrin (amino group). The number of amino groups consumed on carrier TF after conjugation is given in Table 1.

The further confirmation of conjugation was done by determining the mass to-charge ratio of TF, and it’s conjugate (SS-7). The molecular weight of TF was found to be 78993.45 Da (SS). The number of molecules of IH/IG per TF molecule increases with the increase in the molar ratio of protein to the drug (Table 1). The changes in molecular weight were determined by observing the peak shift of the mass spectrum (SS-7). The molecular weight of free IH and free IG was 176.16 and 234.24 respectively. Based on the above findings, the approximate number of molecules of IH/IG per TF molecule was calculated (Table 1). Moreover, the highest ionization density was seen at a molar ratio of 1:60 (T3) in both the conjugation systems. It was also observed that IH is more reactive than IG, as the number of molecules of IH conjugated to TF was higher (Table 1).

The conjugates were also characterized by FTIR. An FTIR spectrum of IH, IG, TF-IH, and TF-IG is given in Fig. S1. The spectra of IH and IG showed broad C=O stretch (carboxylic acid band) at ~1720 cm⁻¹ and O–H stretch at ~3334 cm⁻¹. The peak at 1651 cm⁻¹ indicates C=O stretch of the amide. Spectra also showed N–H bend of pyridine at 1440 cm⁻¹. IG spectra showed C–H (alkane) bend at 1465 cm⁻¹ which is not present in IH spectra confirms the formation of isonicotinoylhydrazinyl-5-oxopentanoic acid. The spectra of TF-IH and TF-IG include the C=O stretching at 1635 cm⁻¹ in the amide I and N–H deformation at 1537 cm⁻¹ in amide II region which are characteristics of an intact α helix structure; indicate that the protein structure in the conjugates was intact. The peak at 1220 cm⁻¹ corresponds to the N–N bond in isoniazid which is not present in TF spectrum. The N–H bend of pyridine in TF-IH/TF-IG and C–H stretch of alkane in TF-IG shifted at 1395 cm⁻¹ and 2840 cm⁻¹ respectively because of conjugation.

3.1. In vitro antitubercular activity

The in vitro antitubercular activity of the INH, IH, IG, TF-IH/IG (T3) was carried out by the determination of the MIC₅₀ (the lowest concentration of the antibiotic at which 90% of the isolates was inhibited) using Mtb H37Rv strain. The MIC value of INH (0.25 μg/ml) was found to be close to the literature value (0.2–0.25 μg/ml) [30–32] (Table 2). The derivatization of INH showed significant antimycobacterial activity (6 μg/ml). The MIC value of TF-IH and TF-
IG conjugate was found to be 2.0 \( \mu \text{g/ml} \) and 4.0 \( \mu \text{g/ml} \) respectively (Table 2).

3.2. In vitro cytotoxicity study

From the therapeutic view, it is crucial that drugs should solely kill intracellular pathogen without displaying a cytotoxic effect on the host cell. In the present study, the cytotoxic study of prepared conjugates along with free drug was determined on THP-1 differentiated macrophages using MTT assay. It was found that the free drug and the synthesized conjugates had no cytotoxic effects on the THP-1 macrophages. (Table 2, Fig. 3).

3.3. Intracellular killing of Mtb

INH and its synthesized conjugates were cytotoxic above 25 \( \mu \text{g/ml} \) (Table 2). Hence to avoid false intracellular inhibition, the lowest concentration (MIC) 0.39 \( \mu \text{g/ml} \) was selected for further intracellular assay as at this concentration both INH, and its conjugates showed cell viability (Fig. 4).

Cells were treated with free INH and conjugates (T3); free INH significantly reduced the colony forming unit (CFU) as compared to the untreated control cells (Fig. 4). While in the case of conjugates, further reduction in CFU was observed as compared to that of free INH. Tf-IH showed slightly higher CFU than Tf-IH, this result may be attributed to the number of bound INH molecules to the transferrin (Tf-IH > Tf-IG) present in both the conjugates.

Hence, it is proved that both TF conjugates, TF-IH, and TF-IG meticulously inhibited the intracellular \textit{M. tuberculosis} bacterial burden.

3.4. Stability study

The pH-dependent stability of INH conjugates (T3) in phosphate buffer solution (pH 4.0 and 7.4) were analyzed by RP-HPLC sampling over 24 h. The results are summarized in Fig. 5. The TF-IH conjugate was found to be stable up to 12 h at pH 4.0. However, a further increase in incubation time increased the percent release of INH. While at pH 7.4, the release pattern was slow as compared to
pH 4.0. It may be due to the hydrolysis of hydrazone conjugate which occurs fast in acidic pH. In comparison with Tf-IH, less percent release of INH from Tf-IG was observed at both the pH even after 18 h. Thus stability data reflects the pH-dependent stability of both the linkers for a longer time.

Based on the above findings, it can be considered that Tf-INH conjugates take longer time to release the free drug. We can postulate the plausible mechanism for the cleavage of INH from the synthesized conjugates. Both the conjugates seem to be stable at blood pH (pH 7.4) for a longer time, which is suggestive of the enhanced residence of conjugates in circulation. Tf-IH releases drug faster due to hydrolysis of hydrazone conjugate at acidic pH 4.0 [33,34]. Thus after internalization of Tf conjugate in Mtb infected macrophage through Tf receptors, hydrazone linker may be cleaved in acidic endosome and lysosome via endocytosis, to trigger the release of INH. Tf-IG comparatively releases the drug slowly at pH 4.0, may be due to more stable amide linkage. While it may be assumed that, amide linkage in Tf-IG conjugate may be cleaved by lysosomal protease enzyme to release the free INH moiety in the cytosol [35].

4. Conclusion

In summary, we have synthesized novel isoniazid-transferrin conjugates and examined the influence of chemical linkages (hydrazone and amide linkers) on its antitubercular potential. The study revealed that higher numbers of hydrazone intermediates

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**Table 1**
The number of amino groups consumed on Tf and number of INH molecules per Tf molecule on conjugates determined by TNBS assay and MALDI-TOF spectrometric method at various Tf:IH/IG molar ratio.

<table>
<thead>
<tr>
<th>Conjugate</th>
<th>Tf:IH/IG molar ratio</th>
<th>Approximate no. of amino groups consumed on Tf</th>
<th>MALDI-TOF: Mass (Da)</th>
<th>No. of INH molecules per Tf molecule</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>1:0</td>
<td>0</td>
<td>78993.45</td>
<td>0</td>
</tr>
<tr>
<td>T1</td>
<td>1:20</td>
<td>4</td>
<td>79502.55</td>
<td>2.89 (3)</td>
</tr>
<tr>
<td>T2</td>
<td>1:40</td>
<td>6</td>
<td>79914.77</td>
<td>5.23 (5)</td>
</tr>
<tr>
<td>T3</td>
<td>1:60</td>
<td>8</td>
<td>80195.60</td>
<td>6.82 (7)</td>
</tr>
<tr>
<td>T4</td>
<td>1:100</td>
<td>8</td>
<td>80194.80</td>
<td>6.81 (7)</td>
</tr>
</tbody>
</table>

(-) no significant change.

**Table 2**

<table>
<thead>
<tr>
<th>Compound</th>
<th>MIC&lt;sub&gt;90&lt;/sub&gt; (µg/ml)</th>
<th>IC&lt;sub&gt;50&lt;/sub&gt; (µg/ml) (4days)</th>
</tr>
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<tbody>
<tr>
<td>INH</td>
<td>0.25</td>
<td>&gt;25</td>
</tr>
<tr>
<td>INH-H</td>
<td>6.0</td>
<td>-</td>
</tr>
<tr>
<td>INH-G</td>
<td>6.0</td>
<td>-</td>
</tr>
<tr>
<td>Tf-IH</td>
<td>2.0</td>
<td>&gt;25</td>
</tr>
<tr>
<td>Tf-IG</td>
<td>4.0</td>
<td>&gt;25</td>
</tr>
</tbody>
</table>

* a The half maximal inhibitory concentration (IC<sub>50</sub>).

* b MTT assay performed on THP-1 differentiated macrophages.
were conjugated to the TF suggesting that the hydrazone intermediate was more reactive towards amine functionality of TF than that of amide intermediate. Both the conjugates showed significant antimycobacterial activity in comparison with free INH and have pH dependent stability. While in the case of ex vivo study on infected THP-1 cells, TF-IH showed a substantial decrease in CFU at 0.39 μg/ml concentration in comparison with TF-IG and free INH. Hence, we can conclude that the conjugation of antitubercular agents to human plasma protein based carrier (Transferrin) is a promising approach to improve the efficacy and potency of a drug. Further, in-depth ex vivo and in vivo studies to identify the Mtb targeting efficiency of this novel conjugates and establishment of its delivery system are underway which will be communicated in the future.

Author contribution

All authors contributed equally.

Acknowledgments

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Appendix A. Supplementary data

Supplementary data to this article can be found online at https://doi.org/10.1016/j.ejmech.2019.111713.

Abbreviations

<table>
<thead>
<tr>
<th>Abbreviation</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mtb</td>
<td>Mycobacterium tuberculosis</td>
</tr>
<tr>
<td>TB</td>
<td>tuberculosis</td>
</tr>
<tr>
<td>TF</td>
<td>transferrin</td>
</tr>
<tr>
<td>MDR</td>
<td>multidrug resistant</td>
</tr>
<tr>
<td>XDR</td>
<td>extensively drug-resistant</td>
</tr>
<tr>
<td>HIV</td>
<td>human immunodeficiency virus</td>
</tr>
<tr>
<td>PAS</td>
<td>p-aminosalicylic acid</td>
</tr>
<tr>
<td>QD</td>
<td>quantum dots</td>
</tr>
<tr>
<td>TR</td>
<td>transferrin receptor</td>
</tr>
<tr>
<td>kDa</td>
<td>kiloDalton</td>
</tr>
<tr>
<td>SDS-PAGE</td>
<td>sodium dodecyl sulfate-polyacrylamide gel electrophoresis</td>
</tr>
<tr>
<td>EDC.HCl</td>
<td>1-ethyl-3-(3-dimethylaminopropyl)carbodiimide hydrochloride</td>
</tr>
<tr>
<td>NHS</td>
<td>N-hydroxy succinimide</td>
</tr>
<tr>
<td>ACN</td>
<td>acetonitrile</td>
</tr>
<tr>
<td>MALDI-TOF</td>
<td>Matrix-assisted laser desorption/ionization time of flight</td>
</tr>
<tr>
<td>PBS</td>
<td>phosphate buffer solution</td>
</tr>
<tr>
<td>MIC90</td>
<td>lowest concentration of drug that inhibits growth of more than 90% of the bacterial population</td>
</tr>
<tr>
<td>IC50</td>
<td>lowest concentration of drug that inhibits growth of more than 50% of the cell population</td>
</tr>
</tbody>
</table>

References


