A hitherto unreported impurity in Terazosin – elucidation of the structure, synthesis and cytotoxicity

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Abstract: Analysis of several batches of the α₁-adrenergic blocking agent terazosin being used as a medication for treating benign prostatic hyperplasia and hypertension revealed the presence of a hitherto not reported impurity. The latter was isolated, and its structure was elucidated from NMR and Mass Spectrometry (MS) data and unambiguously confirmed by independent synthesis. This contamination, represented in 1-[(4-amino-6,7-dimethoxyquinazolin-2-yl)piperazin-1-yl]-pentane-1,2-dione 2 is likely to occur as the product of a side-reaction in the catalytic hydrogenation step during the synthesis of the drug. Biological screening showed this compound as not cytotoxic for several human tumor cell lines and non-malignant fibroblasts.

Keywords: Terazosin; reductive ring-opening; cytotoxicity; SRB assay.

1. Introduction

Drugs are developed, produced and marketed to be used as medications. They have to be delivered in utmost pure form. Their purification, however, is most often very challenging since there is an increased regulatory focus on impurities. A usual and generally accepted identification threshold is 0.1% for drugs whose maximum daily dose is less than or equal to 2.0 g while for those drugs whose daily dose exceeds 2.0 g the identification threshold is 0.05% 1-4. During a project dealing with the synthesis of postsynaptic α₁-adrenergic blocking agents, we came across the drug terazosin 1 (Fig. 1) 5-9.

Terazosin was first used in medical practice in 1987 to treat benign prostatic hyperplasia 10-12. A quarter of all 40-year-old men already have benign prostate hyperplasia with the number for 80-year-old men being above 80%. While in stage 1, a conservative "strategy of controlled waiting" is often supported by over-the-counter phytopharmaceuticals, but in stage 2 more effective drugs have to be used, such as α₁ receptor blockers including Terazosin.

![Figure 1. Structure of terazosin 1 and the hitherto unknown impurity 2](image-url)

A second indication for Terazosin, although less widespread, is the treatment of hypertension. It is estimated that about 25% of the world’s population has a too high blood pressure; by 2025 this number of patients is expected to rise to 29%, which means that about 1.5 billion people will be affected 13.

Recently, two other possible applications of Terazosin have become known. These include the treatment of hyperhidrosis (phase 4, NCT00449683) 14-16 and a study indicating possible re-purposing of 1 to treat Parkinson’s disease (PD) 8. PD is the second most common neurodegenerative disease being first
In 2016 alone, 3.2 million DALYs (disability-adjusted life years) - i.e., symptom-free years with a good quality of life were lost worldwide due to this disease. The disease recently claimed 200,000 deaths per year. Recently, 1 has been repurposed as anti-apoptotic drug targeting the human enzyme phosphoglycerate kinase 1.

2. Results and discussion

These data indicate that there may be an increased need for Terazosin in the future. Given the recent problem of critical impurities in several drugs, we examined some batches of Terazosin for impurities. It became apparent that individual batches contained traces (0.05-0.11%) of an impurity 2 not previously having been described in the literature. The structure of 2 was elucidated by the commercially obtained drugs which were extracted with methanol, and the extract was subjected to a repeated semi-preparative HPLC separation (RP18, CH₂CN/H₂O) to yield pure 2 being subject to an excellent elucidation of its structure by Mass spectrometry and NMR spectroscopy. Low-resolution ESI MS showed a quasi-molecular ion [M+H]+ m/z 388.27; this finding was supported by HRMS showing m/z 388.1977 and - as a consequence - a molecular formula of C₁₉H₂₄O₄N₅S. An ESI-MS fragmentation experiment gave fragments m/z 360.2 (Δm = 28, CO) and m/z = 290.2 (splitting off of the side chain from the piperazinyl residue).

Two carbonyl groups were detected in the ¹³C NMR spectrum at δ = 202.4 and 166.3 ppm, respectively. An intact piperazinyl moiety was established by the presence of the two N-CH₂-groups in the respective ¹H and ¹³C NMR spectra, and the two OMe groups were detected at δ = 56.3 and 55.9 ppm, respectively; their unambiguous assignment was made by 2D-NOESY NMR spectroscopy. The ¹H NMR spectrum also showed the presence of a n-propyl side chain. From these results as well as extra 2D NMR spectra, the structure of 2 was deduced as depicted in Fig 1.

A synthesis made independent confirmation of this proposal of 2 was designed and carried out (Scheme 1). Thus, commercially available 2-chloro-6,7-dimethoxyquinazoline-4-amine (3) was allowed to react with piperazine for 6 hours at reflux temperature in water to yield 68% of known compound 4. The latter compound was transformed into target amide 2 by its reaction with 2-oxopentanoic acid in the presence of HOBT, EDCI and TEA to yield 50% of 2. The sample obtained by synthesis was identical in each property (IR, UV-Vis, ¹H NMR, ¹³C NMR, MS) with compound 2 obtained from the extraction of the drug.

![Scheme 1](image)

**Scheme 1.** Synthesis of compound 2: a) piperazine, water, 100°C, 6 h, 68%; b) 2-oxopentanoic acid, DCM, HOBT, EDCI, TEA, 0°C → 12 h, 25°C, 50%

Terazosin 1 is synthesized from the reaction of 3 with 6 (Scheme 2); the latter compound can be obtained from the coupling of piperazine with 2-furoylchloride followed by hydrogenation in the presence of Raney nickel. Usually, hydrogenation of furans in the presence of metal catalysts leads to the formation of tetrahydrofurans in good yields.

However, there are some reports on ring-opening reactions under these conditions. Thus, 2-methylfuran forms 2-pentanone upon hydrogenation in the presence of Ru, Pt, Pd or a ZnO/Ni. Since these reactions usually give good yields when higher pressures and/or higher reaction temperatures are used, it can be assumed that - when applying low pressure of hydrogen and room temperature – this reaction only occurs as a minor side-reaction on a small scale. Thus, it seems highly probable that the impurity 2 observed in the end product is due to an already existing impurity 7 being formed during the synthesis of 6. Ring-opening reactions during the synthesis of 1 are known: For example, European
Pharmacopoeia lists two impurities, a 5-hydroxypentanoyl compound (impurity F) and 2-hydroxy-pentanoyl compound (impurity J). The latter is related to the 2-oxo-pentanoyl impurity 2 described here.

Terazosin, as well as the novel impurity 2 were tested for cytotoxic activity in sulforhodamine B assays employing several human tumor cells lines [A375 (melanoma), HT29 (colon carcinoma), MCF-7 (breast adenocarcinoma), A2780 (ovarian carcinoma), FaDu (hypopharynx carcinoma)] and non-malignant mouse fibroblasts (NIH 3T3). While 2 was not cytotoxic at all up to a concentration of EC50 = 30 μM (cut-off of the assay), terazosin was not cytotoxic for non-malignant mouse fibroblasts (NIH 3T3 EC50 > 30 μM) but slightly cytotoxic for HT29 (22.4±1.6 μM), MCF-7 (28.8±1.0 μM), A2780 (20.1±0.7 μM), A375 (26.6±1.7 μM) and FaDu cells (3.1±1.3 μM), respectively.

![Scheme 2. Reported synthesis of terazosin 1 and the proposed mechanism for the formation of impurity 2](image)

3. Conclusion

A hitherto unknown impurity in commercial samples of Terazosin was detected, isolated and its structure elucidated by spectroscopy and independent synthesis. This contamination, 1-[4-(amino-6,7-dimethoxyquinazolin-2-yl)-piperazin-1-yl]-pentane-1,2-dione 2 is likely to occur as the product of a side-reaction in the catalytic hydrogenation step during the synthesis of the drug. Biological screening showed this compound as not cytotoxic for several human tumor cell lines and non-malignant mouse fibroblasts.

4. Experimental

4.1. General

Melting points are uncorrected (Leica hot stage microscope). NMR spectra were recorded using the Varian spectrometers Gemini 2000 or Unity 500 (δ given in ppm, J in Hz); MS spectra were taken on a Finnigan MAT LCQ 7000 (electrospray, voltage 4.5 kV, sheath gas nitrogen) instrument. IR spectra were taken on a Perkin Elmer Spectrum Two (ATR) and UV/Vis on a Perkin Elmer Lambda 14 instrument. TLC was performed on silica gel (Merck 5554); elemental analyses were performed on a Vario EL (CHNS). The solvents were dried according to usual procedures. For HPLC-DAD investigations a LaChrom D-7000 HPLC-system (from Merck-Hitachi) was used at an operating temperature of 25°C using RP-18 Superspher column (125-2 mm, 4 mm) as the stationary phase.

6,6-Dimethoxy-2-(piperazin-1-yl)-quinazolin-4-amine 4

A mixture of 2-chloro-6,7-dimethoxyquinazolin-4-amine (3, 9.96 g, 41.56 mmol) and piperazine (35.8 g, 41.56 mmol) in water (50 mL) was stirred at 100 °C for 6 h. After cooling to room temperature, an aqueous solution of KOH (0.85 mol, 50 mL, 1.7 N) was slowly added, and the mixture was stirred for 1 h at room temperature (thereby the solution turned green, and a white precipitate was formed). The precipitate was filtered off, re-dissolved in water (100 mL) and extracted with a mixture of chloroform and isopropanol (100 mL, 5:1). The aqueous phase was extracted with a mixture of chloroform and isopropanol (2x100 mL, 5:1), and the combined organic phases were washed with brine (50 mL). The solvents were removed under diminished pressure. The residue was recrystallized from methanol to yield 4 (8.2 g, 68%) as a white solid; m.p. 223.8°C (from MeOH) (lit.:24 230-232°C); Rf (CHCl3: MeOH: NH4OH, 90:9:1) = 0.23.

1-[4-(Amino-6,7-dimethoxyquinazolin-2-yl)-piperazin-1-yl]-pentane-1,2-dione 2

To an ice-cold solution of 2-oxopentanoic acid (1.12 g, 9.6 mmol) in anhydrous DCM (250 mL), HOBT (1.51 g, 11.2 mmol), EDCI (3.68 g, 19.2 mmol), and TEA (3.47 mL, 24.0 mmol) were very
slowly added. After an additional stirring for 30 min at 25°C, 4 (2.31 g, 28.8 mmol) was added, and the mixture was stirred for 12 h at room temperature. The precipitate was filtered off, and the filtrate was washed with saturated NaHCO₃ solution (2x 50 mL). water (2x 50 mL) and brine (2x 50 mL). After drying (MgSO₄) and evaporation of the solvent under diminished pressure, the residual yellowish oil was purified by column chromatography (CHCl₃/MeOH/ NH₄OH, 98:1:8:0.2) yielding 2 (1.54 g, 50%) as an off-white solid; m.p. 175.4°C; Rf = 0.21; IR (ATR); ν = 2934w, 2876w, 2837w, 1709m, 1622s, 1575m, 1513m, 1480s, 1476s, 1472s, 1435s, 1380m, 1239m, 1208s, 1174m, 1130w, 997s, 840m cm⁻¹; UV-Vis (MeOH): λₘₐₓ (log ε) = 214 (4.23), 250 (4.70), 275 (4.13), 324 (3.71) nm; 

I H NMR (400 MHz, DMSO- d₆): δ = 7.41 (s, 1H, 3-H), 7.14 (s, 2H, 7-NH₂), 6.72 (s, 1H, 6-H), 3.81 (s, 3H, 18-H), 3.77 (s, 3H, 19-H₂), 3.77 - 3.70 (m, 4H, 9-H₂ + 12-H₂), 3.56 - 3.49 (m, 2H, 10-H₂), 3.39 - 3.32 (m, 2H, 11-H₂), 2.72 (t, J = 7.3 Hz, 2H, 15-H₂), 1.57 (h, J = 7.3 Hz, 2H, 16-H₂), 0.90 (t, J = 7.4 Hz, 3H, 17-H₃) ppm; 

I C NMR (101 MHz, DMSO-d₆): δ = 202.4 (C-14), 166.3 (C-13), 161.6 (C-7), 158.6 (C-8), 154.7 (C-2), 149.1 (C-5), 145.6 (C-1), 105.7 (C-6), 104.1 (C-3), 103.5 (C-4), 56.3 (C-19), 55.9 (C-18), 45.5 (C-11), 44.3 (C-12), 43.6 (C-9), 41.8 (C-15), 41.2 (C-10), 16.3 (C-16), 13.9 (C-17) ppm. 

MS (ESI, MeOH): calculated for C₁₉H₂₅N₅O₄: 387.19, found: m/z (%) = 388.27 (100%). [M+H]+; analysis calcd for C₁₉H₂₅N₅O₄ (387.44): C 58.90, H 6.50, N 18.08; found: C 58.75, H 6.71, N 17.76; HRMS: calcd. 388.1979, found: 388.1977

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