The synthesis of gliflozins

INTRODUCTION

Gliflozins constitute a class of compounds that is useful as sodium glucose co-transporter-2 (SGLT2) inhibitors. The gliflozins have shown particular expediency in the treatment of diabetes 2. They accomplish this through blocking of sodium glucose transport proteins, which, in turn, inhibit the kidneys from resorbing glucose back into the blood stream. The excess, non-resorbed glucose is then eliminated with the urine with the net result being a dosage-regulated glucose level. It has been shown that a key feature of the gliflozins is their ability to distinguish between the inhibition of the SGLT1 transporter, a low-capacity, high-affinity transporter that is expressed in the gut, heart and kidney, and the SGLT2 transporter, a high-capacity, low-affinity transporter expressed mostly in the kidney. It is estimated that by the year 2025 nearly 400 million people will suffer from diabetes 2 (1). Due in large part to this growing population of diabetes sufferers, considerable efforts have been and continue to be taken in the gliflozin approach towards addressing the diabetes 2 affliction. A significant part of these efforts has centered on various derivatives of the gliflozin class, represented by the general structure 1. (Figure 1a) An excellent review of the structure activity relationship and some of the history of this class of drugs has appeared (2). This mini-review will look at some of the general approaches to the synthesis of the gliflozins with an emphasis on the glycosidation and the key reduction steps.

As early as the 19th century phlorizin was isolated and shown to inhibit both SGLT1 and SGLT2 and, thus, promote urinary excretion of glucose. Unfortunately, being a hemiacetal structure, phlorizin is subject to hydrolysis to phloretin and glucose thus excluding it as a candidate for oral administration as the SGLT inhibitory activity of the phloretin is significantly reduced (3). (Figure 1b) Nevertheless, this led to investigations of structurally similar, more hydrolytically stable derivatives of phlorizin as SGLT2 inhibitors (4,5).

Five of the gliflozins have now been approved for prescribed use. These are dapagliflozin 4, Farxica or Fonica, (Bristol Myers-Squibb/AstraZeneca), canagliflozin 5, Invokana, (Janssen Pharmaceuticals), ipragliflozin 6, Suglat, (Allersta Pharmaceuticals), empagliflozin 7, Jardiance, (Boehringer-Ingeheim/Eli Lilly) and ertugliflozin 8, Sitaglipton, (Pfizer/Merck) (Figure 1c).

The general structures of the gliflozins have in common a glucose sugar to which is attached an aromatic group in the β−position at the anomeric carbon 1. It will be noted that in addition to the glucose sugar moiety and the β−isomeric aryl substituent the aryl group is composed of a diarylmethylene structure. The differences in the structures are not large, vis the similarities in the structures of dapagliflozin, empagliflozin and ertugliflozin as well as those of canagliflozin and ipragliflozin.
SYNTHESIS OF GLIFLOZINS

The synthetic approaches to the gliflozins essentially consist of three general steps: 1) construction of the aryl substituent, 2) introduction of the aryl moiety onto the sugar or glucosylation of the aryl substituent, and 3) deprotection and modification of the arylated anomeric center of the sugar to the final desired product.

Synthesis of the aryl group

The synthetic approach to the gliflozins first involves the construction of the aryl substituent with a suitable handle to allow its attachment to the sugar. There are two classical entries into the diarylmethane units, namely, reaction of an organometallic reagent with an aldehyde to give the diaryl carbinol or a Friedel-Crafts acylation to provide a diaryl ketone. Both of these intermediates would then be followed by a reduction of the functionalized carbon to the requisite methylene. The former of these is exemplified in a synthetic route to 12 wherein aldehyde 9 was reacted with the Grignard reagent 10 to form the diarylmethanol 11, which was then reduced with triethylysilane to the diarylmethane 12 (6). It is worth noting that organosilanes have been shown to be excellent reagents for the reduction of benzylic alcohols to the corresponding alkanes (7,8) (Figure 2a).

A similar approach was employed in procuring the aryl substrate in a synthesis of ipragliflozin (9) (Figure 2b).

In an alternate approach to the diarylcarbinol precursor to the diarylmethane moiety Fürstner and Krause showed that the rhodium trichloride-catalyzed coupling of an aryl boronic acid with an aldehyde leads to the diaryl carbinol intermediates in high yields (10). Two of the 16 examples reported involved the preparation of an aryl bromide, indicating that the reaction can be carried out without competing aryl-aryl, Suzuki-type cross-coupling taking place. The resulting brominated diarylmethane could be converted to an organometallic reagent for reaction with the glucosonide leading to a gliflozin (Figure 2c).

Addition of the aryl group to the glucose

The second aspect in the synthesis of the gliflozins is that of glucosylation of the aryl moiety with the final stereochemistry being that wherein the aryl group occupies a β-position. One early approach to the gliflozins is illustrated for the synthesis of 17 as a general entry into the C1-aryl glucosonides.
The tribenzyl glycal 23 was converted to the gliflozin structures in a couple of ways. One of these is via the corresponding epoxide 24, which serves to place the 3-hydroxyl group on the sugar as well as provide a route for the introduction of the aryl unit (22). Reaction of 24 with 2-lithiofuran leads to the β−substituted benzyl-protected glucose 25. On the other hand reaction of the epoxide with the corresponding zinc reagent 26 results in the introduction of the aryl group in the α−position. This is based on the earlier work of Halcomb and Danishefsky (23) (Figure 5a).

The di-tert-butylsilylene-protected dihydropyran 27 was subjected to a Stille cross-coupling reaction with an aryl sulfonyl chloride to show that this approach will work to introduce an aryl group to the sugar. Introduction of the requisite 3-hydroxyl group in the α−position was then accomplished via hydroboration/oxidation to provide the β−arylated sugar derivative 29 (24) (Figure 5b).

Finally, the tetramethylidisiloxane, TMDS, silane reducing agent has been employed in the successful reduction of a hemiketal to the corresponding furan derivative as shown in the reduction of the protected precursor to canagliflozin, which is based on some earlier work by Kraus and coworkers who demonstrated the slanie reduction of sugar hemiketals (17-20).

A highly stereoselective introduction of the aryl moiety can be accomplished via the cross-coupling of an aryl zinc reagent, using the well-developed organozinc chemistry of Knochel and co-workers, with the bromo-functionalized glucose 21 (21). Thus, zinc reagent 21 was reacted with bromopyran 22 to introduce the aryl group in the β−position assisted by the neighboring group participation of the pivaloyl group. Other ester protecting groups for the hydroxyls on the sugar also worked, but the pivalate protection was shown to be preferred for reasons of selectivity and yield. Deprotection led to dapagliflozin 4 (Figure 4b).

Gong and Gagné have shown that the reaction of arylzinc reagents with suitably-protected glucosyl bromides under Ni(0) catalysis results in the C1-aryl glucoside with good β−selectivity (25) (Figure 5c).
Nanoparticles that deliver oligonucleotide drugs into cells

Therapeutic oligonucleotide analogs represent a new and promising family of drugs that act on nucleic acid targets such as RNA or DNA; however, their effectiveness has been limited due to difficulty crossing the cell membrane. A new delivery approach based on cell-penetrating peptide nanoparticles can efficiently transport charge-neutral oligonucleotide analogs into cells, as reported in Nucleic Acid Therapeutics, a peer-reviewed journal from Mary Ann Liebert, Inc., publishers.

In the article, "Peptide Nanoparticle Delivery of Charge-Neutral Splice-Switching Morpholino Oligonucleotides," Peter Järner and coauthors, Cambridge Biomedical Campus (U.K.), Karolinska University Hospital (Huddinge, Sweden), Stockholm University (Sweden), Alexandria University (Egypt), and University of Oxford (U.K.), note that while delivery systems exist to facilitate cell entry of negatively charged oligonucleotide drugs, these approaches are not effective for charge-neutral oligonucleotide analogs. The authors describe lipid-functionalized peptides that form a complex with charge-neutral morpholino oligonucleotides, enabling them to cross into cells and retain their biological activity. "The exploitation of phosphorodiamidate morpholinos represents an exciting approach to treating a number of therapeutic targets," says Executive Editor Graham C. Parker, PhD, The Carman and Ann Adams Department of Pediatrics, Wayne State University School of Medicine, Children's Hospital of Michigan, Detroit, MI. "This paper suggests an intriguing but practical approach to solving the lack of a convenient non-covalent delivery system."

Nucleic Acid Therapeutics journal