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Synthesis, characterization, hypoglycemic and aldose reductase inhibition activity of arylsulfonylspiro[fluorene-9,5'-imidazolidine]-2',4'-diones

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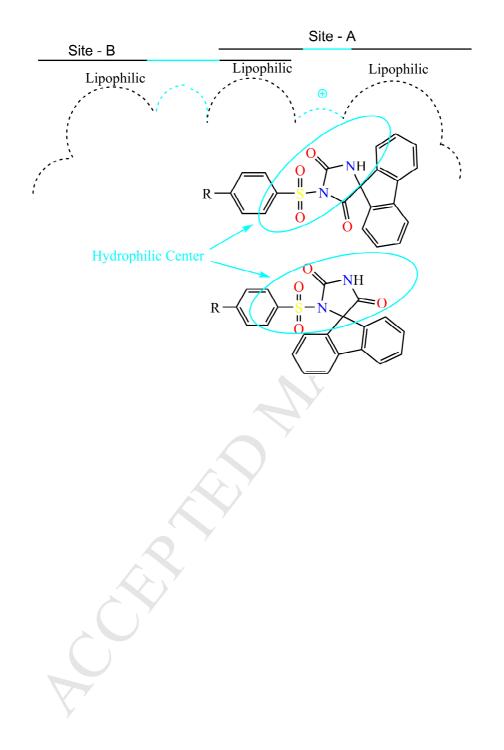
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Three of the synthesized sulfonylcyclic urea derivatives were found more potent *in-vivo* hypoglycemic agents than glibenclamide. In *in-vitro* ARI assay, the compounds 3a-g were found more active ALR1 inhibitors than the standard.



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