

Cite this: *RSC Adv.*, 2018, 8, 40760

## Carbohydrate-conjugated 4-(1,3,2-dithiarsolan-2-yl)aniline as a cytotoxic agent against colorectal cancer

Boqiao Fu,<sup>a</sup> Xiaolin Wang,<sup>b</sup> Yingjie Li,<sup>b</sup> Jingying Hu,<sup>c</sup> Dai Lu,<sup>d</sup> Wei Li,<sup>a</sup> Kewang Zheng<sup>a</sup> and Caiqin Qin<sup>\*a</sup>

Arsenic trioxide (As<sub>2</sub>O<sub>3</sub>) has been approved for the treatment of acute promyelocytic leukemia (APL); however, its use in the treatment of solid tumors is limited due to its pharmacokinetic properties. Organic arsenic compounds provide better options for pharmaceutical optimization. *p*-Aminophenyl arsenoxide (*p*-APAO), an organic arsenic compound, was found to interact with the promyelocytic leukemia–retinoic acid receptor alpha (PML–RARα) fusion protein in a similar manner to arsenic trioxide. Analogs of *p*-APAO such as 4-(1,3,2-dithiarsolan-2-yl)aniline (*p*-APDTAs) were recently found to show improved cytotoxicity toward several solid tumor cell lines with lower toxicity to normal cells. Here, we synthesized a carbohydrate-conjugated 4-(1,3,2-dithiarsolan-2-yl)aniline (*p*-APDTAs) and showed that it exhibited reduced cytotoxicity to normal cells, suggesting a feasible approach to improve the therapeutic index of arsenic-containing compounds as chemotherapeutic agents.

Received 21st September 2018

Accepted 24th November 2018

DOI: 10.1039/c8ra07860b

rsc.li/rsc-advances

Arsenic trioxide (As<sub>2</sub>O<sub>3</sub>) has been used for various purposes in ancient China and Greece for more than 2400 years.<sup>1,2</sup> Use of arsenic-containing compounds to treat hematologic disorders such as leukemia, Hodgkin's disease, and pernicious anemia has been reported since the 1700s.<sup>3</sup> In the 1970s, a group of physicians in China reported that a formulation of arsenic trioxide induced complete remission in a small group of patients with acute promyelocytic leukemia (APL).<sup>1</sup> Later on, the follow-up clinical trials by American investigators confirmed the clinical efficacy of arsenic trioxide in the management of APL. Subsequently, arsenic trioxide (Trisenox<sup>TM</sup>) was approved by the U.S. Food and Drug Administration (FDA) in 2000 for the treatment of relapsed or refractory APL.<sup>4</sup>

Investigations to decipher the mechanism of action of arsenic trioxide revealed that the drug is able to promote the catabolic degradation of an oncogenic fusion protein that derives the proliferation of APL cells and has been found in over 98% of cases of human APL.<sup>5</sup> The oncogenic fusion protein is the result of chromosomal rearrangements that juxtapose the

*promyelocytic leukemia (PML) gene* and the retinoic acid receptor alpha (*RARα*) gene.<sup>6</sup> Arsenic trioxide is capable of covalently modifying a cysteine-rich region of the PML moiety of the PML–RARα fusion protein, leading to proteasome-dependent degradation of the oncogenic PML–RARα fusion protein.<sup>7,8</sup> Further studies have shown that arsenic trioxide can induce apoptosis of hematologic cancer cells and solid tumor cells *via* a myriad of mechanisms such as disruption of mitochondrial functions and cellular redox processes, activation of different caspases, and downregulation of Bcl-2 expression.<sup>9,10</sup>

Despite the remarkable success of arsenic trioxide in the treatment of APL, the inorganic arsenic compound has certain limitations as a chemotherapeutic such as the systemic toxicity associated with a high amount of arsenic compounds in the blood, and poor pharmacokinetic properties.<sup>2,11</sup> Although arsenic trioxide has shown efficacy for solid tumors in many preclinical studies, the therapeutic effects of low-dose arsenic oxide for solid tumors have not been clinically proven yet. This may be attributed to the rapid renal clearance of arsenic trioxide metabolites. This likely results in an insufficient amount of arsenic oxide at the tumor sites.<sup>2</sup>

Along with the inorganic arsenic trioxide, organic arsenic compounds such as *p*-aminophenylarsine oxide (*p*-APAO)<sup>5,12,13</sup> and others<sup>3,14–16</sup> were investigated either as preclinical and clinical experimental drugs or as molecular probes in cancer cells. Arsenic sulfide and its derivatives were reported with potent antitumor activities to solid tumor cell lines such as HCT 116.<sup>17,18</sup> In comparison with inorganic arsenic compounds, organic arsenic compounds offer certain advantages. They are

<sup>a</sup>Hubei Provincial Collaborative Innovation Center of Biomass Resources Transformation and Utilization, College of Chemistry and Materials Science, Hubei Engineering University, Hubei, 432000, P. R. China. E-mail: fuboqiao@126.com

<sup>b</sup>Guangdong Institute of Gastroenterology, Guangdong Provincial Key Laboratory of Colorectal and Pelvic Floor Diseases, The Sixth Affiliated Hospital, Sun Yat-sen University, Guangzhou, Guangdong, 510655, China

<sup>c</sup>State Key Laboratory of Oncogenes and Related Genes, Shanghai Cancer Institute, Renji Hospital, Shanghai Jiaotong University School of Medicine, Shanghai, 200032, China

<sup>d</sup>Department of Pharmaceutical Sciences, Rangel College of Pharmacy, Texas A&M University, Texas, 78363, USA



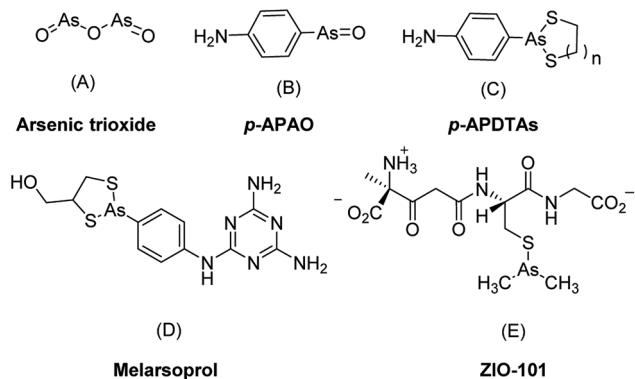


Fig. 1 Representative arsenic-containing compounds showing anti-cancer activities.

generally more chemically stable than inorganic arsenic compounds and are easier for structural optimization to improve drug-like properties such as pharmacokinetic properties, therapeutic efficacy, and target selectivity *etc.* Some of the organic arsenic compounds, such as melarsoprol (Fig. 1D) and ZIO-101 (Fig. 1E) are more efficacious than arsenic trioxide to induce apoptosis in cancer cells.<sup>3</sup> Interestingly, a biotin-conjugated *p*-APAO was found capable to bind to PML-RAR $\alpha$  in a manner similar to arsenic trioxide (As<sub>2</sub>O<sub>3</sub>).<sup>5,13</sup> These suggested that compounds containing a *p*-APAO moiety may possess anticancer activities As<sub>2</sub>O<sub>3</sub>. However, *p*-APAO is a substituted phenyl arsonoxide, which can be easily oxidized to pentavalent arsenic (As<sup>V</sup>) compound and has higher tendency to form polymers.<sup>15,19</sup> Recently, it was reported that *p*-APAO analogues 4-(1,3,2-dithiarsolan-2-yl)aniline (*p*-APDTAs, Fig. 1C) obtained by protecting the arsonoxide moiety of *p*-APAO with dithiols are more efficacious than *p*-APAO in inducing apoptosis in several cancer cell lines. Furthermore, the *p*-APDTAs showed less cytotoxicity toward normal cells than *p*-APAO.<sup>20</sup> Collectively, these triggered our interests to employ *p*-APDTAs (Fig. 1C) rather than the *p*-APAO (Fig. 1B) as the arsenic-containing moiety for developing novel arsenic-containing compounds as anticancer agents.

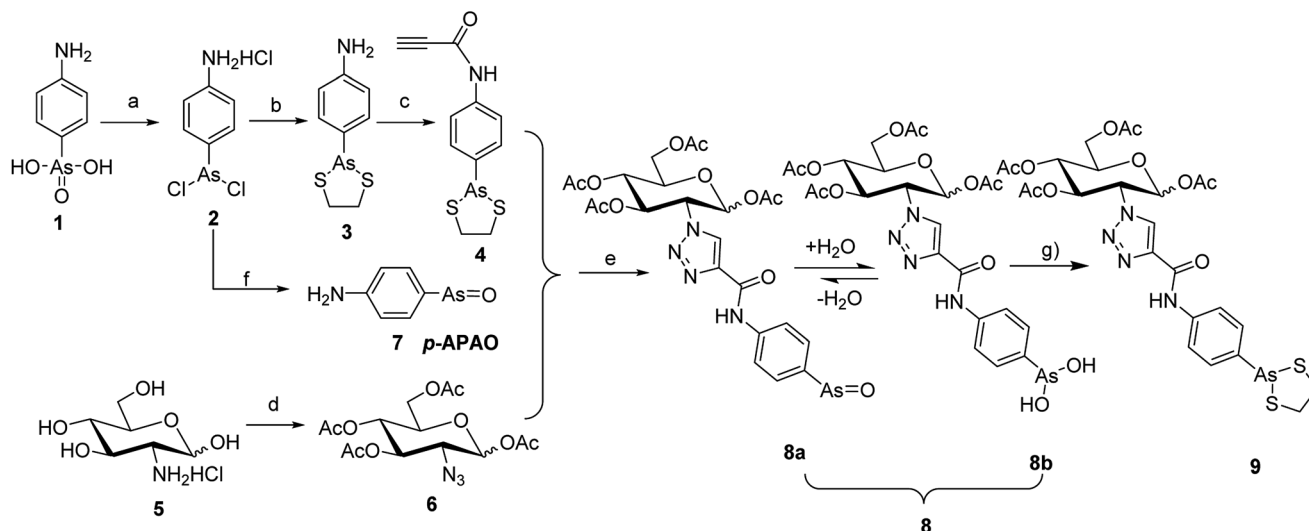
In our earlier investigation of detoxification of *p*-APAO with various carbohydrates, we found that 1,3,4,6-tetra-*O*-acetyl- $\beta$ -D-glucosamine (OADG) is capable of reducing the toxicity of *p*-APAO toward the normal organelle modeled by *S. cerevisiae*.<sup>21</sup> The natural product Jusbetonin is an indolo[3,2-*b*]quinoline alkaloid glycoside containing a  $\beta$ -D-glucose moiety. In a structural optimization, the replacement of its  $\beta$ -D-glucose with OADG was shown to enhance the cytotoxicity against breast cancer cell line MDA-231.<sup>22</sup> Based on these results, we hypothesized that introducing OADG into organic arsenic compounds such as *p*-APDTAs may preserve the anticancer effects of organic arsenic compounds while the cytotoxicity toward normal cells can be reduced. Hence, we designed and synthesized OADG-conjugated *p*-APAO analogue (8) and *p*-APDTAs analogue (9). We assessed the cytotoxicity of the synthesized conjugates toward the solid tumor cells from three colorectal cancer cell lines HCT116, DLD1, and RKO and the cytotoxicity toward normal cells using epithelial NCM460 cells as the model.<sup>23</sup>

Click chemistry has been frequently applied in conjugation of bioactive molecules with various pharmaceutical agents and biomolecules for preclinical and clinical applications.<sup>24</sup> Its typical process involves azide-alkyne [3 + 2] dipolar cycloaddition to form a 1,4-regioselectivity 1,2,3-triazole-based linker under the catalysis of Cu<sup>I</sup>. The products from click chemistry has been proven to be superior in satisfying many criteria in drug development and biomedical research such as excellent biocompatibility, selectivity, yield, and stereospecificity.<sup>24</sup> Many known 1,2,3-triazoles have various biological activities such as anti-HIV, anticancer, and antibacterial activities.<sup>25–27</sup> In our study, the target compound was synthesized through the click reaction between a carbohydrate (*i.e.*, OADG) moiety functionalized with an azido group and a *p*-APDTAs moiety functionalized by a propargyl group. The synthetic chemistry employed in this work is illustrated in Scheme 1 and 2.

To synthesize the alkyn-containing reactant (4) for click reaction, commercially available *p*-arsanilic acid 1 was first converted into 4-aminophenyldichloroarsine (2) according to reported protocols.<sup>20,28,29</sup> The product 2 was easily transformed into the intermediate 3 through reaction with 1,2-ethanedithiol under aqueous sodium carbonate. Compound 3 reacted with propionic acid in the presence of DCC to provide the click chemistry reactant 4. To synthesize the other reactant of the click reaction (*i.e.*, azide 6), which bears the 1,3,4,6-tetra-*O*-acetyl- $\beta$ -D-glucosamine (OADG) moiety, the amino group of commercially available 2-amino-2-deoxy-D-glucose (5) was converted to azido functionality by reaction with trifluoromethanesulfonyl azide, followed by acetylation of its 1,3,4,6 hydroxyl groups to yield the key building block 6.<sup>30,31</sup> Coupling of 4 with 6 under typical conditions of click reaction produced the desired conjugate 8, which exists as a mixture of equilibrium of arsenic oxide 8a and its hydrated analog arsonous acid 8b. The presence of the arsenic oxide 8a and arsonous acid 8b was confirmed by NMR and mass spectrometry analysis.<sup>32</sup> To obtain the *p*-APDTAs analogue (9), the mixture of arsenic compounds 8a and 8b was treated with 1,2-ethanedithiol in anhydrous methanol to provide the target compound. The reference compound *p*-APAO (7) was prepared as previously described,<sup>28</sup> and used to compare with other derivatives of dithiarsolanes in the cytotoxicity assays. Because of formation of a triazole moiety during the linkage by click reaction, we synthesized the carbohydrate-triazole-containing compounds 11 and 15 to study the biological influence of the triazole moiety in the structure of 9 (Scheme 2).

It was reported that derivatives of arsenic trioxide (As<sub>2</sub>O<sub>3</sub>) slightly inhibited the proliferation of human colon cancer cell line HCT116.<sup>17,34</sup> To determine the antitumor activities of the synthesized compounds in colorectal cancer, three common colorectal cancer cell lines (*i.e.*, HCT116, DLD1, and RKO)<sup>33</sup> were selected for cytotoxicity assessment of the synthesized compounds (3, 4, *p*-APAO, 8, 9, 11, 15) and the standard chemotherapeutic for colorectal cancer 5-FU was used as a positive control drug.<sup>35</sup> Their cytotoxicity in intestinal epithelial cell line NCM460, which is a model of normal cells, were also evaluated. The half maximal inhibitory concentrations (IC<sub>50</sub>) of the synthesized compounds are shown in Table 1



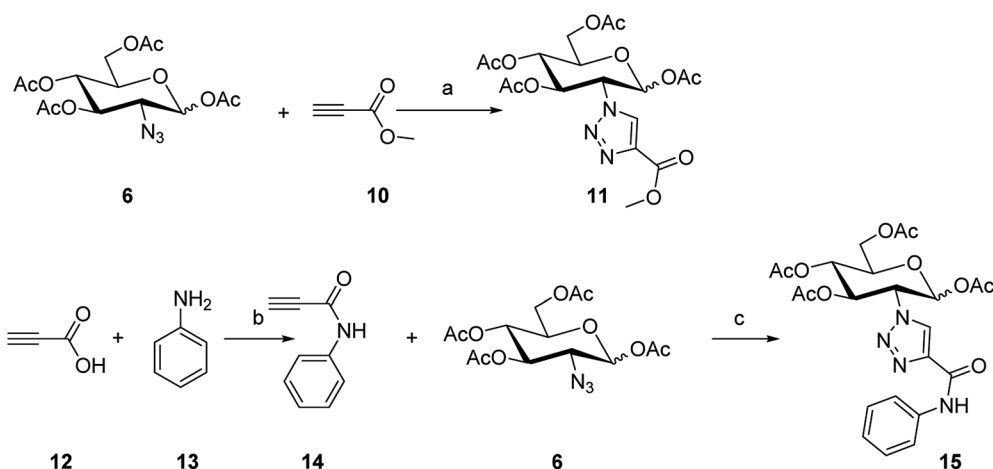


**Scheme 1** Synthesis routes of compounds **7** and **9**. Reaction conditions: (a)  $\text{SO}_2$ , KI, HCl, MeOH; (b) ethane-1,2-dithiol,  $\text{NaHCO}_3$ , MeOH, 82%; (c) propionic acid, dicyclohexyl carbodiimide,  $\text{CH}_2\text{Cl}_2$ , r.t., 64%; (d) (i)  $\text{TfN}_3$ ,  $\text{CuSO}_4 \cdot 5\text{H}_2\text{O}$ ,  $\text{CH}_3\text{CN}$ ; (ii)  $\text{Ac}_2\text{O}$ , pyridine; (e) sodium L-ascorbate,  $\text{CuSO}_4 \cdot 5\text{H}_2\text{O}$ , microwave,  $100^\circ\text{C}$ , 38.3%; (f)  $\text{NH}_3 \cdot \text{H}_2\text{O}$ ,  $\text{N}_2$ , 53.9%; (g) ethane-1,2-dithiol, MeOH, 69%.

and the dose dependent inhibition of proliferation by compound **9** and **5-FU** are illustrated in Fig. 2.

Similar to **5-FU**, arsenic containing compounds **3**, **4**, **8**, **9** and **p-APAO** all exhibited anti-proliferation activity in the three colorectal cancer cell lines. Particularly, arsenic compound **4** showed significantly enhanced activity cross the three colorectal cancer cell lines in comparison with **5-FU**. However, these compounds showed higher cytotoxicity in normal cells except the *p*-APDTAs analogue (**9**), which exhibited potent activity against HCT116 cells with  $\text{IC}_{50}$  value of  $1.29 \pm 0.21 \mu\text{M}$ . It also showed strong activities against DLD1 and RKO cell lines with  $\text{IC}_{50}$  values of  $13.96 \pm 2.05 \mu\text{M}$  and  $12.55 \pm 1.90 \mu\text{M}$ , respectively. Compound **9** showed stronger cytotoxicity than **5-FU** in HCT-116 and DLD1 cells, but not in RKO cells. Compare to **5-FU**, compound **9** displayed similar cytotoxicity towards normal colorectal endothelial cells NCM460, but with over 1.8 fold increase of cytotoxicity to colorectal cancer HCT-116 cells.

The results from compounds **11** and **15** indicated that the *p*-APDTAs moiety is critical for the anticancer effects of compound **9**. The absence of the arsolane group significantly reduced antitumor activity as observed from compound **15**. Similarly, the triazole-containing compound **11**, which is derived from 1,3,4,6-tetra-*O*-acetyl- $\beta$ -D-glucosamine (OADG), was not responsible for the anticancer effects of compound **9**. The compound **8** is the OADG-conjugated derivative of *p*-APDTAs and the compound **9** is the OADG-conjugated derivative of *p*-APDTAs. In comparison with their non-conjugated counterpart (*i.e.* **8** vs. *p*-APDTAs, and **9** vs. **3**), the OADG-conjugated organic arsenic compounds **8** and **9** showed significantly reduced toxicity towards normal colorectal cells while their anti-proliferation activities were generally maintained. We also found that deacetylation of compound **9** with sodium methoxide and then neutralization with acid resin couldn't give deacetylation compound. Actually dithiol protective group on



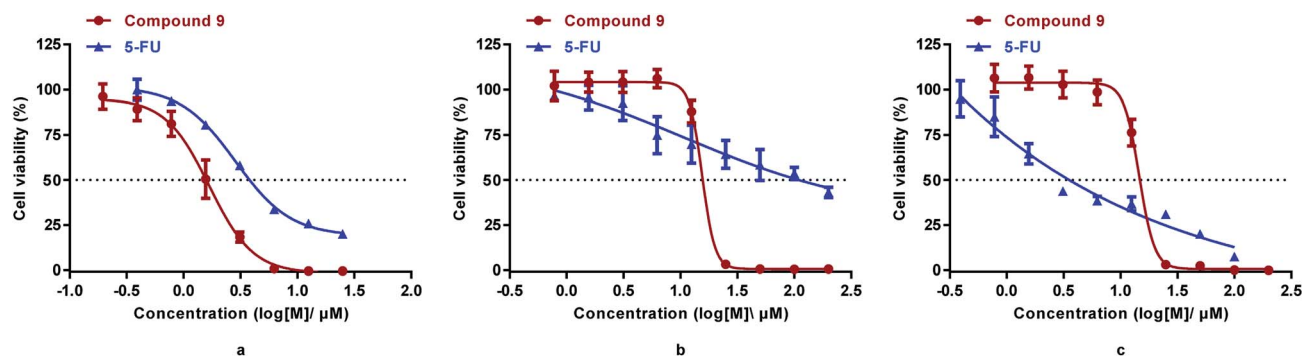
**Scheme 2** Synthesis routes of compounds **11** and **15**. Reaction conditions: (a) sodium L-ascorbate,  $\text{CuSO}_4 \cdot 5\text{H}_2\text{O}$ , microwave,  $100^\circ\text{C}$ , 61.1%; (b) propionic acid, dicyclohexyl carbodiimide,  $\text{CH}_2\text{Cl}_2$ , r.t., 70%; (c) sodium L-ascorbate,  $\text{CuSO}_4 \cdot 5\text{H}_2\text{O}$ , microwave,  $100^\circ\text{C}$ , 32.5%.



**Table 1** Cytotoxicity of carbohydrate-conjugated and non-conjugated dithiarsolanes in colorectal cancer cells and normal cells

IC <sub>50</sub> (μM) ± SD (μM)								
Compounds	5-FU	<i>p</i> -APAO	3	4	8	9	11	15
NCM460 <sup>a</sup>	44.57 ± 4.69	3.67 ± 0.10	11.11 ± 0.82	3.20 ± 0.78	16.15 ± 2.21	50.75 ± 0.97	ND <sup>b</sup>	ND <sup>b</sup>
HCT-116	3.62 ± 1.22	5.03 ± 0.66	1.86 ± 0.34	0.53 ± 0.20	17.69 ± 1.44	1.29 ± 0.21	>50	43.39 ± 4.17
DLD1	>50	2.17 ± 0.21	6.90 ± 1.81	1.41 ± 0.59	14.96 ± 2.72	13.96 ± 2.05	>50	20.28 ± 2.30
RKO	2.09 ± 0.87	2.99 ± 0.04	3.74 ± 0.88	0.79 ± 0.05	4.57 ± 0.17	12.55 ± 1.90	20.03 ± 3.38	>50
Selectivity index <sup>c</sup>	12.31	0.73	5.97	6.04	0.91	39.34	—	—

<sup>a</sup> NCM460 cell is used as the model of normal cell. <sup>b</sup> ND = not determined. <sup>c</sup> Selectivity index: the IC<sub>50</sub> against HCT116 over the IC<sub>50</sub> against NCM460.

**Fig. 2** Cytotoxicity of compound 9 and 5-FU in HCT-116 (a), DLD1(b), and RKO (c) cells measured by the MTS assay (mean ± SD).

the arsenic compound **9** was also deprotected under acid resin.<sup>36,37</sup> When we used dithiol to protect arsenic acid of deacetylation carbohydrate-conjugated arsenic compound again, all the naked hydroxyl groups could be protected. So it is very difficult to get the deacetyl compound from **9**. The usefulness of 1,3,4,6-tetra-*O*-acetyl-β-D-glucosamine (OADG) in anti-cancer molecules were also demonstrated by others. The acetyl protective groups were deemed beneficial in comparison with the unprotected β-D-glucosamine bearing analogues.<sup>22</sup>

## Conclusions

In this investigation, it was found that conjugation of cytotoxic arsenic-containing compound *p*-APDTAs with β-D-glucose **9** derived **1**, **3**, **4**, 6-*O*-acetyl-β-D-glucosamine (OADG) can significantly reduce the toxicity to normal colorectal endothelial cells while the cytotoxicity towards colorectal cancer cells can be maintained. The conjugation linker introduced by click chemistry did not abolish the anticancer activity of the resultant compound, which showed anticancer activity comparable to the standard chemotherapeutic 5-fluorouracil (5-FU) in colorectal cancer models. These results suggested a new approach for the discovery and development of arsenic-containing compounds as novel chemotherapeutics for the treatment of solid tumors.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

This work was supported by Natural Science Foundation of Hubei Province, China (2014CFB570), the National Natural Science Foundation of China (Grant No. 31371750 and 81502022) and the Fundamental Research Fund for the Sun Yat-sen University (Grant No. 16ykpy35).

## References

- 1 S. Waxman and K. C. Anderson, *Oncologist*, 2001, **6**, 3–10.
- 2 E. P. Swindell, P. L. Hankins, H. Chen, Đ. U. Miodragović and T. V. O'Halloran, *Inorg. Chem.*, 2013, **52**, 12292–12304.
- 3 P. J. Dilda and P. J. Hogg, *Cancer Treat. Rev.*, 2007, **33**, 542–564.
- 4 K. H. Antman, *Oncologist*, 2001, **6**, 1–2.
- 5 X.-W. Zhang, X.-J. Yan, Z.-R. Zhou, F.-F. Yang, Z.-Y. Wu, H.-B. Sun, W.-X. Liang, A.-X. Song, V. Lallemand-Breitenbach and M. Jeanne, *Science*, 2010, **328**, 240–243.
- 6 G.-B. Zhou, J. Zhang, Z.-Y. Wang, S.-J. Chen and Z. Chen, *Philos. Trans. R. Soc., B*, 2007, **362**, 959–971.
- 7 A. Emadi and S. D. Gore, *Blood Rev.*, 2010, **24**, 191–199.
- 8 J. Zhu and V. Lallemand-Breitenbach, *Oncogene*, 2001, **20**, 7257.
- 9 A. M. Evens, M. S. Tallman and R. B. Gartenhaus, *Leuk. Res.*, 2004, **28**, 891–900.
- 10 K. Alimoghaddam, *Int. J. Hematol. Oncol. Stem Cell Res.*, 2014, **8**, 44.



- 11 I. Khairul, Q. Q. Wang, Y. H. Jiang, C. Wang and H. Naranmandura, *Oncotarget*, 2017, **8**, 23905.
- 12 H.-N. Zhang, L. Yang, J.-Y. Ling, D. M. Czajkowsky, J.-F. Wang, X.-W. Zhang, Y.-M. Zhou, F. Ge, M.-K. Yang and Q. Xiong, *Proc. Natl. Acad. Sci. U. S. A.*, 2015, **112**, 15084–15089.
- 13 X. Zhang, F. Yang, J.-Y. Shim, K. L. Kirk, D. E. Anderson and X. Chen, *Cancer Lett.*, 2007, **255**, 95–106.
- 14 E. Hikita, M. Arai, S. Tanaka, K. Onda, H. Utsumi, B. Yuan, H. Toyoda and T. Hirano, *Anticancer Res.*, 2011, **31**, 4169–4178.
- 15 N. Wimmer, J. A. Robinson, N. Gopisetty-Venkata, S. J. Roberts-Thomson, G. R. Monteith and I. Toth, *Med. Chem.*, 2006, **2**, 79–87.
- 16 X.-Y. Fan, X.-Y. Chen, Y.-J. Liu, H.-M. Zhong, F.-L. Jiang and Y. Liu, *Sci. Rep.*, 2016, **6**, 29865.
- 17 L. Zhang, Y. Tong, X. Zhang, M. Pan and S. Chen, *Drug Des., Dev. Ther.*, 2015, **9**, 5851–5862.
- 18 W. Ding, Y. Tong, X. Zhang, M. Pan and S. Chen, *Sci. Rep.*, 2016, **6**, 19793.
- 19 C. Huang, Q. Yin, J. Meng, W. Zhu, Y. Yang, X. Qian and Y. Xu, *Chem.-Eur. J.*, 2013, **19**, 7739–7747.
- 20 Y. Liu, D. Duan, J. Yao, B. Zhang, S. Peng, H. Ma, Y. Song and J. Fang, *J. Med. Chem.*, 2014, **57**, 5203–5211.
- 21 B. Fu, L. Lv, C. Xia, K. Zheng, L. Zhu, W. Li, Y. Yan and C. Qin, *J. Med. Chem. Toxicol.*, 2017, **2**, 28–33.
- 22 S. Liu, J. Meng, W. Zhang, S. Wan and T. Jiang, *J. Heterocycl. Chem.*, 2016, **53**, 1194–1199.
- 23 S. L. Organ, J. Hai, N. Radulovich, C. B. Marshall, L. Leung, T. Sasazuki, S. Shirasawa, C. Zhu, R. Navab, M. Ikura and M. Tsao, *PLoS One*, 2014, **9**, e86103.
- 24 K. Nwe and M. W. Brechbiel, *Cancer Biother. Radiopharm.*, 2009, **24**, 289–302.
- 25 R. Alvarez, S. Velazquez, A. San-Felix, S. Aquaro, E. De Clercq, C.-F. Perno, A. Karlsson, J. Balzarini and M. J. Camarasa, *J. Med. Chem.*, 1994, **37**, 4185–4194.
- 26 (a) S. Velazquez, R. Alvarez, C. Perez, F. Gago, C. De, J. Balzarini and M. J. Camarasa, *Antiviral Chem. Chemother.*, 1998, **9**, 481–489; (b) M. J. Genin, D. A. Allwine, D. J. Anderson, M. R. Barbachyn, D. E. Emmert, S. A. Garmon, D. R. Graber, K. C. Grega, J. B. Hester, D. K. Hutchinson, J. Morris, R. J. Reischer, C. W. Ford, G. E. Zurenko, J. C. Hamel, R. D. Schaadt, D. Stapert and B. H. Yagi, *J. Med. Chem.*, 2000, **43**, 953–970.
- 27 J. Heredia-Moya and K. L. Kirk, *Bioorg. Med. Chem.*, 2008, **16**, 5743–5746.
- 28 K. J. Stevenson, G. Hale and R. N. Perham, *Biochemistry*, 1978, **17**, 2189–2192.
- 29 J. Liu, M. M. Numa, H. Liu, S.-J. Huang, P. Sears, A. R. Shikhman and C.-H. Wong, *J. Org. Chem.*, 2004, **69**, 6273–6283.
- 30 Q.-H. Fan, N.-T. Ni, Q. Li, L.-H. Zhang and X.-S. Ye, *Org. Lett.*, 2006, **8**, 1007–1009.
- 31 B. Fu, C. Qin, C. Xia and X. Yang, Synthesis Method of arsenic-sugar with anticancer biological activity and its application, Chinese Patent, ZL201510437907.9, 2015.
- 32 H. R. Lee, H.-J. Cheong, S.-J. Kim, N.-S. Lee, H.-S. Park and J.-H. Won, *Oncol. Rep.*, 2008, **20**, 41–47.
- 33 G. Malich, B. Markovic and C. Winder, *Toxicology*, 1997, **124**, 179–192.
- 34 D. U. Miodragović, J. A. Quentzel, J. W. K. C. L. Stern, R. W. Ahn, I. Kandela, A. Mazar and T. V. O'Halloran, *Angew. Chem., Int. Ed.*, 2013, **52**, 10749–10752.
- 35 B. Pardini, R. Kumar, A. Naccarati, J. Novotny, R. B. Prasad, A. Forsti, K. Hemminki, P. Vodicka and J. Lorenzo Bermejo, *Br. J. Clin. Pharmacol.*, 2011, **72**, 162–163.
- 36 C. Lin, T. Sugai, R. L. Halcomb, Y. Ichikawa and C. Wong, *J. Am. Chem. Soc.*, 1992, **114**, 10138–10145.
- 37 S. D. Rychnovsky and R. C. Hoye, *J. Am. Chem. Soc.*, 1994, **116**, 1753–1765.

