Pyridine Moiety: Recent Advances in Cancer Treatment

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Sahu et al.: A Review: An Insight on Recent Advancement of Pyridine Moiety in Treatment of Cancer

Pyridine is an imperative pharmacophore, a privileged scaffold and an exceptional heterocyclic system in the field of drug discovery which provides many opportunities in study/explore this moiety as an anticancer agent by acting on various receptors of utmost importance. Several pyridine derivatives are reported to inhibit tubulin polymerization, androgen receptors, human carbonic anhydrase, kinase, topoisomerase enzyme and many other targets for controlling and curing global health issue of cancer. Now a days in combination with other moieties researchers are focusing for development of pyridine new entities for the treatment of cancer. This review throws light on recent biological expansions of pyridine along with their structure activity relationships/molecular docking to deliver association between various synthesized newer derivatives and receptor sites.

Key words: Pyridine moiety, anticancer, heterocyclic compound, cancer treatment

Cancer is a global health issue that affects a vast portion of the human population and is regarded as the uncontrolled growth of cells in the body, which is one of the most challenging and complex diseases to treat[1,2]. It is considered as another major cause of the death after cardio vascular disorders worldwide[3]. In 2018, world health organization (WHO) reported that 18.1 million people around the globe had cancer and 9.6 million deceased from the disease. By 2040, these records will nearly twice, with the highest increase in low and middle income countries, where greater than two thirds of the world’s tumors will occur[4]. Cancer is the cause of almost 30 % of all hasty deaths from non-communicable diseases among adults aged 30 to 69. Among all, the most frequently diagnosed cancer is lung (11.6 % of all cases), followed by female breast (11.6 %) and colorectal cancers (10.2 %). Lung cancer is the foremost cause of death from cancer (18.4 % of all deaths), followed by colorectal (9.2 %) and stomach cancers (8.2 %). The most usual cause of cancer is tobacco usage, which accounts for 25 % of all cancer deaths around globally. Hence cancer is a serious issue hampering human health. So, as to tackle them new various anticancer agents are being developed ever now and then. Still their toxicity profile has restricted their clinical use as anticancer agents[5]. Therefore, further exploration of new chemo therapeutic agents with great efficacy and least adverse effects is critically essential for medicinal chemists.

Anticancer drugs are categorized into several groups based on their mechanism of action that is biosynthetic blocker of nucleic acid, transcription interferer, structural and functional of Deoxyribonucleic acid (DNA) interferer, interferer of protein synthesis and function, influencer of hormone homeostasis; or modulator of immune system[6]. To decrease toxic effects of antitumor agents towards normal cells, specific targeted drug therapy is chosen which is considered to hit tumor cells only[7]. As specified by the USA national cancer institute drug repository, 227 endorsed antitumor treatments are at present avowed to treat roughly 40 types of malignancies[6,8]. Because of great demand of antitumor medications (tons/y) investigators are going on in the way to develop new and effective treatment for tumor[8]. For this reason some researchers or scholars looked into the effect produced by various compounds containing pyridine moiety.

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Pyridine compounds (fig. 1) are well defined by the presence of a six membered heterocyclic ring with the chemical formula $C_5H_4N$, comprising of five carbon atoms and one nitrogen atom. In many aspects it can be correlated to well recognized and fundamental aromatic benzene molecule, with one C-H group changed by a nitrogen atom. It was first isolated from bone oil and coal tar and characterized by Anderson in 1846. The cyclic nature of pyridine was identified by Dewar and Körner in 1869. It plays an important role by catalyzing both biological and chemical systems. It has a conjugated arrangement of six π electrons precisely as benzene has, that are delocalized about the heterocyclic ring. The molecule is planar in nature and follows Huckel standards for aromaticity. Additionally, pyridines are a class of both synthetically and naturally occurring heterocyclic compounds and its nucleus is a well-studied six membered heterocyclic moiety with a wide range of biological and therapeutic applications. Moreover, the current interest in the development of new anticancer agents can be partially ascribed to the increasing appearance of multidrug resistance and adverse side effects are a serious threat to public health. Therefore, the development of new and efficacious medications is a very significant goal and most of the researcher’s efforts in this field are directed towards the design of new agents. It is reported that some important anticancer drugs possess a pyridine nucleus. Thus, this study gives promising compounds possessing a pyridine nucleus that can be investigated for future in vivo and clinically oriented studies.

Some of the physical properties and marketed products of pyridine containing antitumor drugs are mentioned in Table 1 and Table 2, whereas in Table 3 patents on pyridine are mentioned respectively.

### ANTICANCER PROFILE OF PYRIDINE BASED DERIVATIVES

Recently reported some synthetic new compounds that have a promising and potential activity against cancer cells and has pyridine moiety as essential part in those structure are mentioned here along with their structure activity relationship (SAR) or molecular docking or both studies and acting targets or inhibitory target.

#### Migration and Ras related C3 botulinum toxin substrate 1 (RAC1) inhibitors:

A cancer cell finishes a series of stages including migration from the primary cancer, surrounding tissues invasion and intravasation through basement membranes, survival during circulation and arrest at a distant target organ. The Ras-related C3 botulinum toxin substrate 1 (RAC1) is a member of the Ras-related C3 botulinum toxin substrate (RAC) subfamily of guanosine triphosphate (GTPase). It has been involved in the regulation of cellular migration and invasion in breast tumor cells. Rac1 is activated by Guanine nucleotide exchange factors (GEFs) that are regulated by many of cell surface receptors. Hence, therapeutic approaches that inhibit binding of GEFs to Rac1 are a rational means to migration inhibition of cancer cells.

Vlaar et al. explored a carbozole derivatives in order to develop new anti-migratory agents to prevent the malignance cell spreading by inhibiting migration and RAC1. Compounds 1a and 1b (fig. 2) inhibit migration activity of metastatic cell line MDA-MB-231 (Breast carcinoma cell line) by 32 % and 34 %, respectively. Furthermore compound 1b was shown to inhibit activation of the Ras homologous (Rho) GTPase Rac1 by 55 % at 0.25 μM in both MDA-MB-231 and MDA-MB-435 (Breast carcinoma cell line) cell lines.

#### Indoleamine 2,3-dioxygenase 1 inhibitors:

The metabolism of Tryptophan (Trp) occurs through two independent pathways: the kynurenine pathway (KP) that comprises of the oxidative ring opening of the

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**TABLE 1: A PHYSICAL PROPERTY OF THE PYRIDINE**

<table>
<thead>
<tr>
<th>Formula</th>
<th>Molecular weight</th>
<th>Density</th>
<th>Melting point</th>
<th>Boiling point</th>
<th>Refractive</th>
<th>Index</th>
<th>Solubility</th>
</tr>
</thead>
<tbody>
<tr>
<td>$C_5H_4N$</td>
<td>79.102 Da</td>
<td>0.9819 g/ml</td>
<td>−41.6º</td>
<td>115.2º</td>
<td>1.5093</td>
<td>Miscible in all proportions with water and most common organic solvents</td>
<td></td>
</tr>
<tr>
<td>S. No.</td>
<td>Drug</td>
<td>Brand and company</td>
<td>Primary target</td>
<td>Food and drug administration (FDA) approval</td>
<td>Structure</td>
<td></td>
<td></td>
</tr>
<tr>
<td>-------</td>
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<td>---------------</td>
<td>------------------------------------------</td>
<td>-----------</td>
<td></td>
<td></td>
</tr>
<tr>
<td>1.</td>
<td>Apalutamide</td>
<td>ERLEADA, Janssen Biotech Inc.</td>
<td>Androgen receptor inhibitor</td>
<td>In September, 2019</td>
<td><img src="image1" alt="Structure" /></td>
<td></td>
<td></td>
</tr>
<tr>
<td>2.</td>
<td>Pexidartinib</td>
<td>TURALIO, Daiichi Sankyo</td>
<td>Kinase inhibitor</td>
<td>In August, 2019</td>
<td><img src="image2" alt="Structure" /></td>
<td></td>
<td></td>
</tr>
<tr>
<td>3.</td>
<td>Alpelisib</td>
<td>PIQRAY, Novartis Pharmaceuticals Corporation</td>
<td>Kinase inhibitor</td>
<td>In May, 2019</td>
<td><img src="image3" alt="Structure" /></td>
<td></td>
<td></td>
</tr>
<tr>
<td>4.</td>
<td>Venetoclax</td>
<td>VENCLEXTA, AbbVie Inc. and Genentech Inc.</td>
<td>BCL-2 inhibitor</td>
<td>In May, 2019</td>
<td><img src="image4" alt="Structure" /></td>
<td></td>
<td></td>
</tr>
<tr>
<td>5.</td>
<td>Ivosidenib</td>
<td>TIBSOVO, Agios Pharmaceuticals Inc.</td>
<td>IDH1 inhibitor</td>
<td>In May, 2019</td>
<td><img src="image5" alt="Structure" /></td>
<td></td>
<td></td>
</tr>
<tr>
<td>6.</td>
<td>Lorlatinib</td>
<td>LORBRENA, Pfizer Inc.</td>
<td>Kinase inhibitor</td>
<td>In November, 2018</td>
<td><img src="image6" alt="Structure" /></td>
<td></td>
<td></td>
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<tr>
<td>7.</td>
<td>Netarsudil</td>
<td>RHOPRESSA, Aerie Pharmaceuticals</td>
<td>Rho kinase inhibitor</td>
<td>In December, 2017</td>
<td><img src="image7" alt="Structure" /></td>
<td></td>
<td></td>
</tr>
<tr>
<td>8.</td>
<td>Acalabrutinib</td>
<td>CALQUENCE, AstraZeneca</td>
<td>Kinase inhibitor</td>
<td>In October, 2017</td>
<td><img src="image8" alt="Structure" /></td>
<td></td>
<td></td>
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</table>
9. Abemaciclib VERZENIO, Eli Lilly and Company Kinase inhibitor In September, 2017

10. Enasidenib IDHIFA, Celgene Corp. IDH2 inhibitor In August, 2017


### TABLE 3: RECENTLY PATENTED DRUGS THAT CONTAIN PYRIDINE IN THEIR PRIMARY STRUCTURE HAVE ANTICANCER ACTIVITY

<table>
<thead>
<tr>
<th>S.No.</th>
<th>Patent no.</th>
<th>Patent date</th>
<th>Inventors</th>
<th>Description</th>
</tr>
</thead>
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<tr>
<td>2.</td>
<td>US20200062754A1 [25]</td>
<td>February 27, 2020</td>
<td>Xiong Cai, Changeng Qian, Haixiao Zhai</td>
<td>Fused amino pyridine as hsp90 inhibitors</td>
</tr>
</tbody>
</table>
  Zhang Sanqi, Fan Ye, Cao Yongxiao, Shen Ying, Yang Xueyan, Xin Minxing, Cao Lei
  2- (camptothecin-10-oxyl) acetamide compound and application thereof

  Alfredo C. Castro, David T. Jonaitis
  Noriyasu Haginoya, Takashi Suzuki, Miho Hayakawa, Masahiro Ota, Tomoharu Tsukada, Katsuhiko Kobayashi, Yosuke Ando, Takeshi Jimbo, Koichi Nakamura
  Polymorphic compounds and uses thereof

    Zhang Sanqi, Fan Ye, Cao Yongxiao, Shen Ying, Yang Xueyan, Xin Minxing, Cao Lei
    2- (camptothecin-10-oxyl) acetamide compound and application thereof

11. US20200010420A1 [34] January 09, 2020
    Zhang Sanqi, Fan Ye, Cao Yongxiao, Shen Ying, Yang Xueyan, Xin Minxing, Cao Lei
    Pyridone derivatives having tetrahydropyranymethyl groups

    Zhang Sanqi, Fan Ye, Cao Yongxiao, Shen Ying, Yang Xueyan, Xin Minxing, Cao Lei
    Pyridyl inhibitors of hedgehog signalling

    Zhang Sanqi, Fan Ye, Cao Yongxiao, Shen Ying, Yang Xueyan, Xin Minxing, Cao Lei
    Sulfonamide derivatives for the treatment of abnormal cell growth

    Zhang Sanqi, Fan Ye, Cao Yongxiao, Shen Ying, Yang Xueyan, Xin Minxing, Cao Lei
    Biaryl compositions and methods for modulating a kinase cascade

15. US20190343814A1 [38] November 07, 2019
    Zhang Sanqi, Fan Ye, Cao Yongxiao, Shen Ying, Yang Xueyan, Xin Minxing, Cao Lei
    Heterocyclic compounds and uses thereof

    Zhang Sanqi, Fan Ye, Cao Yongxiao, Shen Ying, Yang Xueyan, Xin Minxing, Cao Lei
    3-(benzoimidazol-2-yl)-indazole inhibitors of the Wnt signaling pathway and therapeutic uses thereof

17. US2019035785A1 [40] November 07, 2019
    Zhang Sanqi, Fan Ye, Cao Yongxiao, Shen Ying, Yang Xueyan, Xin Minxing, Cao Lei
    Pyrazolopyridine compounds and uses thereof

    Zhang Sanqi, Fan Ye, Cao Yongxiao, Shen Ying, Yang Xueyan, Xin Minxing, Cao Lei
    Tricyclic compounds as modulators of TNF-α synthesis and as PDE4 inhibitors

    Zhang Sanqi, Fan Ye, Cao Yongxiao, Shen Ying, Yang Xueyan, Xin Minxing, Cao Lei
    Quinoline derivative

    Zhang Sanqi, Fan Ye, Cao Yongxiao, Shen Ying, Yang Xueyan, Xin Minxing, Cao Lei
    Crystalline form of r)-3-((4-(2-(2-methyltetrazol-5-yl) pyridin-5-yl)-3-fluorophenyl)-5-hydroxymethyl oxazolidin-2-one dihydrogen phosphate

21. RU2682245C1 [44] March 18, 2019
    Zhang Sanqi, Fan Ye, Cao Yongxiao, Shen Ying, Yang Xueyan, Xin Minxing, Cao Lei
    Indazolyl Wnt signal pathway inhibitors and their therapeutic applications (s,e)-3-((6-aminopyridin-3-yl)-n-((5-(4-(3-fluoro-3-methylpyrrolidine-1-carbonyl) phenyl)-7-(4-fluorophenyl) benzofuran-2-yl)methyl) acrylamide for the treatment of cancer

    Zhang Sanqi, Fan Ye, Cao Yongxiao, Shen Ying, Yang Xueyan, Xin Minxing, Cao Lei
    Polysaccharide and nucleic acid formulations containing viscosity-lowering agents
Sunil Kumar KC, David Mark Wallace, Jianguo Cao, Chandramouli Chiruta, John Hood
Marion Hitchcock, Anne Mengel, Vera Pütter, Gerhard Siemeister, Antje Margret Wengner, Hans Briem, Knut Eis, Volker Schulze, Amaury Ernesto Fernandez-Montalvan, Stefan Prechtli, Simon Holton, Jorg Fanghanel, Philip Lienau, Cornelia Preusse, Mark Jean Gnoth
3-(1H-pyrrolo[2,3-B]pyridin-2-yl)-1H-pyrazolo[3,4-C]pyridines and therapeutic uses thereof

Marion Hitchcock, Anne Mengel, Vera Pütter, Gerhard Siemeister, Antje Margret Wengner, Hans Briem, Knut Eis, Volker Schulze, Amaury Ernesto Fernandez-Montalvan, Stefan Prechtli, Simon Holton, Jorg Fanghanel, Philip Lienau, Cornelia Preusse, Mark Jean Gnoth
Substituted benzylindazoles for use as BUB1 kinase inhibitors in the treatment of hyperproliferative diseases

Brian A. Sherer, Nadia Brugger Benjamin Bader, Wilhelm Bone, Hans Briem, Uwe Eberspacher, Knut Eis, Joanna Grudzinska-Goebel, Marcus Koppitz, Julien Lefranc, Philip Lienau, Ulrich Lucking, Dieter Moosmayer, Hans Schick, Gerhard Siemeister, Franz Von Nussbaum, Antje Margret Wengner, Lars Wortmann
TLR7/8 antagonists and uses thereof

Benjamin Bader, Wilhelm Bone, Hans Briem, Uwe Eberspacher, Knut Eis, Joanna Grudzinska-Goebel, Marcus Koppitz, Julien Lefranc, Philip Lienau, Ulrich Lucking, Dieter Moosmayer, Hans Schick, Gerhard Siemeister, Franz Von Nussbaum, Antje Margret Wengner, Lars Wortmann
2-(morpholin-4-yl)-l,7-naphthyridines

This group in between the pyridine & morpholine rings, increase growth inhibitory activity

Ortho-substituted compounds more easily adapt the U-shaped conformation that was calculated for binding of EHop-016 to Rac1.

Formed the peptide bond between Val36 and Asp38 residues

These carbazole group occupied the hydrophobic pocket created by Val36 and Ala59

Fig. 2: Docked structure and SAR of carbozole derivatives

pyrrole moiety of the indole as the initiating step and the melatonin pathway that first introduces a hydroxyl group to the indole followed by subsequent biochemical alterations of the hydroxyl and the carboxyl functional groups to produce the hormone melatonin and the neurotransmitter serotonin. The rate limiting step of the KP is catalyzed by the heme-containing Indoleamine 2,3-dioxygenase (IDO) and tryptophan 2,3-dioxygenase, the product N-formyl kynurenine from which is rapidly converted to kynurenine. In mammalians, there are two known isoforms of IDO as IDO1 and IDO2. Among them, IDO2 is the least recognized and its functional role is yet to be fully defined. IDO1 is well defined and recognizes a number of indole type compounds like L/D-Trp and melatonin. IDO1 is up regulated in immune cells by the proinflammatory cytokine Interferon-γ (IFN-γ), which works as a mechanism to prevent overactive immunological responses, hence it prevents damages to host’s tissues and organs[57-60].

Kong et al.[61] designed and synthesized a series of 1H indole-4,7-dione derivatives and evaluated their inhibitory activity in IDO1 and in IFN-γ stimulated Hela (cervical cancer cell line) cells. The SAR and enzyme kinetics experiments revealed that N alkyl substituted side chain of 5-(pyridin-3-yl)-1H-indole-4,7-dione (compound 2) has most promising inhibitory activity (50 % inhibitory concentration value or half maximal inhibitory concentration (IC 50) of 0.16 ±0.02 μM in enzymatic assay) against IDO1 enzyme (fig. 3). Moreover, it has been indicated that this
type of IDO1 inhibitors might be a kind of reversible competitive inhibitors.

Yang et al.\cite{62} synthesized and evaluated two series of naphthoquinone derivatives, naphthoindolizine and indolizinoquinoline-5,12-dione derivatives for their IDO1 inhibitory activity. Many of the compounds showed significant inhibition potency and high selectivity for IDO1 over tryptophan 2,3-dioxygenase and the SAR was also summarized. Among all, the compound 3a (fig. 4) showed promising and most potent with IC\textsubscript{50} value 0.23 μM against IDO1 enzyme. However, compound 3b was also identified as promising lead compound with IC\textsubscript{50} value 0.372 μM against HeLa cell. Furthermore, molecular docking study revealed that it has good interaction with protein and can be useful for further design of new IDO1 inhibitors as antitumor agents.

**Human carbonic anhydrase IX/XII inhibitors:**

Carbonic anhydrase (CAs) is an important target for cancer treatment because of its limited aspects in the normal tissues and predominant expression in varieties of cancer cells\cite{63}. CAs is omnipresent metallo enzymes which catalyze the reversible hydration of carbon dioxide into protons and bicarbonate. Among the seven CA families, αCA are only enzyme subfamily found in human and further categorized in 15 varied human isoforms playing critical roles in a multitude of physiological functions and pathological processes for example respiration and transport of CO\textsubscript{2} and bicarbonate between metabolizing tissues and lungs, homeostasis of pH and CO\textsubscript{2}, secretion of electrolytes in various tissues and organs, biosynthetic reactions (i.e., gluconecogenesis, lipogenesis and ureagenesis), bone resorption, calcification and tumor growth\cite{64,65}. Human CA I and II are cytosolic proteins with high enzymatic efficiency that are constitutively expressed in all tissues. Conversely, human CA IX and XII are membrane related proteins that can be over expressed in the hypoxic cancer environment, with CA IX not considerably present in the majority of healthy tissues\cite{64,66}. Furthermore, it contributes to tumor progression by stimulating tumor cell migration, adhesion and invasion\cite{67}. Now a days CAs has been recognized as a significant biomarker of hypoxia and its over expression is frequently linked with a poor responsiveness to the classical radio and chemotherapies\cite{68}.

Uslu et al.\cite{69} prepared a series of 2-arylbenzimidazole derivatives containing sulfonamide functionality as well as carboxylic acid, hydroxamic acid, carboxamide and boronic acid functionalities, which act as human CA inhibitors. The sulfonamide containing benzimidazole compound 4 derivatives (fig. 5) revealed intriguing inhibitory activity against cancer associated CA IX and XII with inhibitory constant (K\text{\textit{i}}) values in the range of 0.0052-0.0293 μM and 0.0099 -0.0417 μM, respectively. Notably, compound 4 was the most potent and selective CA IX (K\text{\textit{i}}=0.0066 μM) and XII (K\text{\textit{i}}=0.0099 μM) inhibitor with an important selectivity ratio over cytosolic CA I and II isoforms in the range of 0.0034-0.0252 μM.

Ansari et al.\cite{70} design, synthesis and evaluate a series of pyridine-thiazolidinone derivatives in order to obtain human CA IX inhibitors. The binding affinity of the compounds was measured using fluorescence binding studies and enzyme inhibition activity by esterase assay of CA IX. It was observed that compound 5a and 5b
This hydrophilic thiourea group, showed the highest inhibition activity.

Three H-bonds formed with amino acid L234, G236, and G263 by these side chain interaction by pyridine moiety with amino acid F163.

This Electron donating substituent groups exhibited stronger inhibitory activity than other groups.

This 5-sulfonamide group exhibit highest inhibitory activity and selectivity against CA IX/XII.

This pyridine 2-yl moiety increases selectivity towards CA IX (25.2-fold) and CA XII 16.8, respectively

This ring stabilized by interaction with His96

One H-bond formed with Thr200

This di-hydroxyl moiety group exhibited excellent inhibitory activity against CA IX with 14 times selectivity and IC50 = 1.84 µM

This moeity interact via H-bond with His94, His96, Thr199, Thr200

This nitro substituted benzene exhibited high inhibitory activity against CAIX with 9 times selectivity and IC50 = 1.61 µM

This moeity interact via H-bond with His94, His96, Thr199, Thr200

This ring stabilized by interaction with His96

Fig. 4: Docked structure and SAR of naphthoindolizine (3a) and indolizinoquinoline (3b) 5,12-dione derivatives

Fig. 5: Structure of 5-sulfonamide benzimidazole (4) with SAR

Fig. 6: Docked structure and SAR of pyridine-thiazolidinone derivatives
(fig. 6) significantly inhibit the CA IX activity with the 
IC$_{50}$ values 1.61 µM and 1.84 µM, the binding affinity 
for CA IX was significantly great with their equilibrium 
dissociation constant (K$_d$) values 11.21 µM and 
2.32 µM, respectively. All the compounds were further 
screened in vitro for anticancer activity and found that 
compound 5a and 5b exhibit substantial antitumor 
activity against HepG-2 (human liver cancer cell line) 
and MCF-7 (Breast carcinoma cell line) cell lines.

**Mutant isocitrate dehydrogenase (IDH) inhibitors:**

Isocitrate dehydrogenase (IDH) enzyme family, 
includes IDH1, IDH2 and IDH3, are key metabolic 
enzymes which can convert isocitrate to α-ketoglutarate 
(α-KG)[71,72]. Recently IDH mutations have been 
recognized in somatic tumor associated numerous 
hematologic and solid tumors, such as glioma, acute 
myeloid leukemia (AML), cholangiocarcinoma cancer, 
malignant chondrosarcomas 4 and others[73-75]. IDH 
mutations frequently occur at R132 in IDH1 or R140 
and R172 in IDH2, which are situated in the catalytic 
pocket of these enzymes. Considerably, these mutations 
let IDH enzymes to gain an exceptional activity, 
which catalytically alters α-KG to an onco-metabolite 
2-hydroxyglutarate. This onco-metabolite is intensely 
connected to impaired hematopoietic differentiation 
and stimulates leukemia due to its ability of overall 
DNA hyper methylation[76]. Pharmacological barrier 
of mutant IDH1 enzyme efficiently inhibits colony 
formation of IDH mutated patient derived AML cells 
but not that of normal CD34+ (cluster of differentiation 
34 which is a glycosylated transmembrane protein) 
bone marrow cells[77]. By itself, mutant IDH has now 
become a therapeutic specified target of high interest in 
the field of anticancer drug discovery.

Liu et al.[78] synthesized and evaluated a series of 3-aryl-
4-indolylmaleimides in order to get IDH1/R132H 
inhibitors. New structure was acquired through high-
throughput screening and structure-based optimization 
and most compounds displayed great inhibitory effects 
and were highly selective against IDH1/R132H. 
Biological evaluation of the activities and function 
at cellular level revealed that compounds 6a, 6b and 
6c (fig. 7) were the most potent with IC$_{50}$ values of 
0.05 μM, 0.06 μM and 0.06 μM, respectively against 
IDH1/R132H in U87MG (Glioblastoma cell line) cells 
and great selectivity against IDH1/WT (wild type) and 
IDH2/WT. Furthermore, molecular modeling study

<table>
<thead>
<tr>
<th>Compound</th>
<th>R$_1$</th>
<th>R$_2$</th>
<th>R$_3$</th>
</tr>
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<tbody>
<tr>
<td>6a</td>
<td>-H</td>
<td>-CH$_3$</td>
<td>(H$_2$C)$_3$N</td>
</tr>
<tr>
<td>6b</td>
<td>-CH$_3$</td>
<td>-CH$_3$</td>
<td>(H$_2$C)$_3$N</td>
</tr>
<tr>
<td>6c</td>
<td>-Br</td>
<td>-CH$_3$</td>
<td>(H$_2$C)$_3$N</td>
</tr>
</tbody>
</table>

Fig. 7: Docked structure and SAR of 3-aryl-4-indolylmaleimide derivative
provided information that can be useful for the design of new IDH1/R132H inhibitors as anticancer agents. Hu et al. [79] designed, synthesized and biologically evaluated a series of 3-(7-azainodyl)-4-indolylmaleimides for their IDH1/R132H inhibitory activities. Many compounds showed favorable inhibitory effects and were highly selective against the IDH1/R132H. Evaluation of the biological activities at the cellular level showed that compounds 7a (IC50=0.40 µM), 7b (IC50=0.36 µM), 7c (IC50=0.16 µM), 7d (IC50=0.28 µM) and 7e (IC50=0.34 µM) effectively reduce the production of 2-hydroxyglutaric acid in U87MG cells in a dose independent manner, which associates well with their inhibitory activity toward IDH1/R132H. Preliminary SAR and molecular modeling studies (fig. 8) provided that these findings may offer new insights into the development of new IDH1/R132H inhibitors.

**Androgen receptor (AR) inhibitor:**

The Androgen receptor (AR) belongs to the subfamily of steroid receptors, which includes the glucocorticoid and progesterone receptors. They all belong to the nuclear hormone receptor superfamily of ligand activated transcription factors. The AR comprises of a carboxyl (C-) terminal ligand binding domain (LBD), a central DNA binding domain (DBD) and the amino N terminal domain (NTD) protecting the major transactivation function [80-82] and it possesses an extremely conserved DBD, a moderately conserved LBD and a slight conserved NTD [83,84]. All three these domains are important for receptor functions. The biological functions of AR are initiated by binding with 5-dihydrotestosterone/testosterone, which leads to the AR conformational change and its translocation from the cytosol to the nucleus to stimulate transcriptional regulation of specific genes that further modulated by various AR coregulators [85,86]. The AR continues play a stimulator role in several different prostate cancer cell types during progression of prostate cancer (PCa) and metastasis [87]. In the beginning of the treatment, the insufficiency of androgens leads to cancer regression. However, eventually during hormone therapy, PCa develops into a castration resistant stage but remains AR dependent [88,89]. Inactivation of the AR also in castration resistant PCa seems to be therefore a key target of treatment. Current therapeutics include steroidal and non-steroidal anti androgens targeting typically LBD of AR [90-92]. Although other modulators targeting another
domain are speedily investigated and shows beneficial preclinical profiles\cite{84,93}.

Shi et al.\cite{92} developed and synthesized a series of steroidal pyridines derivative through the base promoted three component reaction and preliminarily evaluated for their anti-proliferative activity against cancer cell lines. SARs studies displayed that the 4th position of the pyridine ring were chosen over the phenyl rings for the activity. Among these compounds, 8 (fig. 9) possessing an extra pyridine ring at the p position demonstrated good growth inhibition against cancer cells, especially for PC-3 (prostate cancer cell line) cells with an IC\textsubscript{50} value of 1.55 µM. For the AR+ sensitive PCA cell line LNCaP (prostate cancer cell line), compound 8 inhibited growth with an IC\textsubscript{50} value of 8.48 µM, slightly less potent than the anti-prostate cancer drug abiraterone (IC\textsubscript{50}=3.29 µM). Further mechanistic studies showed that it inhibited colony formation, migration and evasion of PC-3 cells in a concentration dependent manner as well induced apoptosis of PC-3 cells possibly via the mitochondria associated apoptotic pathways.

**Aldo keto reductase (AKR) inhibitors:**

The Aldo keto reductase (AKRs) is a distinct superfamily of proteins that catalyze the reduction of carbonyl groups as well steroid double bonds in the presence of NADPH\cite{94}. It comprises various genes, among them AKR1C1-AKR1C4 genes are situated on chromosome 10 p15-p14 and include of 12 exons. The AKR1C4 and AKR1C3 are almost exclusively involve in the liver and PCa, AKR1C1 and AKR1C2 are most distinguished in the mammary glands includes breast cancer, endometrial cancer, colorectal cancer.

The AKR1C3 (also known as prostaglandin-F synthase) protein catalyzes the conversion of prostaglandins (PGF) H2 and D2 into Prostaglandin F2 alpha (PGF2α) and 9α, 11β-PGF2α respectively. It has the highest catalytic effectiveness than the AKR1C enzymes to interconvert testosterone with 14-androstene 3,17-dione. AKR1C2, is also known as bile acid binding protein and preferentially reduces Dihydrotestosterone (DHT) to the weak metabolite 5α-androstane-3α,17β-diol (3α-diol) without conversion of 3α-diol to DHT in the PC-3 cell line\cite{95-97}.

Savic et al.\cite{98} synthesized and validated new A ring pyridine fused androstanes in 17a-homo-17-oxa (D-homo lactone), 17α-picolyl or 17(E)-picolinylidene series for potential antitumor activity in vitro using human tumor cell lines and recombinant targets of steroidal antitumor drugs. SAR revealed that pyridine fusion to position 3, 4 of the A ring intensely enhance affinity of 17α-picolyl compounds for cytochrome (CYP) 17 while conferring selective anti-proliferative activity against PC-3 cells. Similarly, pyridine fusion to the A ring of steroidal D-homo lactones led to recognition of new inhibitors of AKR1C3. Among them, compound 9 (fig. 10) found to be most promising with IC\textsubscript{50} 9.13 and 11.77 µM in MDA-MB-231 and HeLa cell respectively. Additionally molecular docking also suggested that compound 9 has higher binding affinity toward AKR1C3 (9.8 kcal/mol) similar to androstenedione.

**Kinase inhibitor:**

Kinase inhibitors have played an increasingly key role in the treatment of tumor and other diseases. Currently, more than 25 oncology medications that target kinases have been approved and many drugs therapeutics are in various stages of clinical evaluation\cite{99}. Kinases are enzymes that transfer a phosphate group to a protein whereas phosphatases remove a phosphate group from protein\cite{100}. Almost 538 known kinases are encoded in the human genome and these kinases uphold cellular activities.

This extra pyridine ring over phenyl ring enhance the inhibitory activity

\[ \begin{align*}
\text{H}_{3}
\text{C}_{2}
\text{CH}_{2}
\text{CO} \\
\text{N}
\text{N}
\text{H}_{3}
\text{C}
\text{H}_{3}
\end{align*} \]

**Fig. 9: Structure of steroidal pyridine derivative with SAR**
function by turning protein function on, whereas corresponding phosphatases reverse this action\cite{101,102}. Recent advances demonstrated that molecular mechanisms underlying cancer cell signaling have elucidated a critical role for kinases in the carcinogenesis and metastases of various types of cancer\cite{103}. Since most protein kinases promote cell survival, proliferation and migration, when constitutively overexpressed or active, they are also linked with oncogenesis\cite{104}. Over the past decades, multiple human malignancies have been identified to be associated with modulation and dysfunction of protein and lipid kinases and deactivated phosphatases on account of chromosomal reshuffling and genetic mutations\cite{105,106}. Thus, some example of kinase inhibitors mentioned below that have been recently developed or reported as potential inhibitors of kinase.

Nawaz et al.\cite{107} designed, synthesized and evaluated pyrazoline linked carboxamide derivatives for potential epidermal growth factor receptor (EGFR) kinase inhibitors, apoptotic, cardiomyopathy toxicity and other antitumor activity. Among all the synthesized compounds, 10a and 10b shown (fig. 11) good broad spectrum \textit{in vitro} anticancer activity arrayed with 3-pyridine and the 4-pyridine group respectively, against the A549 (Lung adenocarcinoma cell line) (IC\textsubscript{50} 10.3±1.07 and 4.6±0.57 µM) and HCT-116 (Colon cancer cell line) (IC\textsubscript{50} 12.9±1.76 and 6.5±0.60 µM) cancer cell lines by using doxorubicin as the standard drug. Additionally, compounds 10a and 10b were exhibited to induce apoptosis in A549 cancer cells as evidenced by 4’,6-diamidino-2-phenylindole staining and apoptosis assay, which further strengthened \textit{in vitro} anticancer findings.

Yu et al.\cite{108} designed, synthesized and biologically evaluated a series of imidazo[1,2-a]pyridine derivatives for potential phosphoinositide 3-kinase (PI3K) and mammalian target of rapamycin (mTOR) dual inhibitors as an effective targeting pathway for tumor therapy. Among the synthesized compounds, compound 11 was proved to be a potent PI3K/mTOR dual inhibitor with exceptional kinase selectivity, modest plasma clearance and acceptable oral bioavailability, with IC\textsubscript{50} 0.01, 0.052, 0.04, 0.071, 0.0052, 15.53 µM against various HCT-116, HT-29 (Colon cancer cell line), MCF-7, PC-3, LOVO (Colon cancer cell line) and HUVEC (Human umbilical vein endothelial cells) cell lines respectively. Besides, compound 11 displayed significant inhibition with IC\textsubscript{50} 0.20 0.58 1.20 0.50 21 nM against PI3K\textsubscript{α}, PI3K\textsubscript{β}, PI3K\textsubscript{γ}, PI3K\textsubscript{δ} and mTOR respectively. The docking results also revealed the inhibitory activity of compound 11 via forming various interactive bonds with PI3K\textsubscript{α} and mTOR (fig. 12).

Ullah et al.\cite{109} synthesized and evaluated a series of pyridine-pyrazole-benzothiourea and pyridine-pyrazole-benzenesulfonamide scaffold for their inhibitory effect on human nucleotide Ectonucleotide pyrophosphatase/phosphodiesterase (ENPP) 1 and ENPP3 isoenzymes. Among them, compound 12a was the most potent inhibitor of ENPP3 (IC\textsubscript{50} =0.21 µM) and compound 12b was much selective to ENPP1 with IC\textsubscript{50} value of 0.40 µM and found promising cytotoxic against HeLa, MCF-7 and 1321N1 (Astrocytoma cell line) cell lines. However, 12b displayed preferential cytotoxicity against MCF-7 (IC\textsubscript{50} =16.05 µM), which is similar to the potency of cisplatin. Additionally, docking results showed considerable binding interactions with active sites of ENPP isoenzymes (fig. 13).

\textbf{B-cell lymphoma 2 (Bcl-2) inhibitors:}
Apoptosis is a form of planned cell death that is activated in response to developmental cues or cellular stress. This selective cell suicide shows a crucial role in various physiological and pathological processes including growth, immunity and disease by the elimination of damaged or unnecessary cells helps to ensure organs health\textsuperscript{[110]}. B-cell lymphoma 2 (Bcl-2) is a family of proteins responsible as key regulators of cell death that can either suppress (pro survival) or promote (pro apoptotic) apoptosis\textsuperscript{[111,112]}. The pro survival subfamily includes Bcl-2, B-cell lymphoma extra-large (Bcl-XL), Bcl-w (anti-apoptotic protein), anti-apoptotic member of the Bcl-2 (A1) and myeloid cell leukemia-1 (MCL-1) whereas the pro apoptotic subfamily is classified into the multi domain group (BCL2-associated X Protein (Bax), BCL-2-antagonist/killer (Bak) and BCL-2 related ovarian killer (Bok)) and the BCL-2 Homology 3 (BH3) only group\textsuperscript{[113]}.

Favorable results are being reported with the usage of inhibitors of BCL-2 and other related molecules, especially with BH3 mimetics\textsuperscript{[114,115]}. Various studies suggested that apoptosis blockage is a crucial oncogenic mechanism in lymphoid malignancies and that BCL-2 overexpression is a common finding in leukemia and lymphomas, numerous antagonists of anti-apoptotic BCL-2 have been investigated and developed for the treatment of hematological neoplasms\textsuperscript{[116,117]}.

Santosh et al.\textsuperscript{[118]} synthesized, characterized and evaluated a series of oxadiazole and hydroxypyrazoline derivatives for promising anticancer inhibitors in which compounds 13a and 13b (fig. 14) exhibited remarkable...
growth inhibition of MDA-MB-231 (IC$_{50}$: 10.2 ±0.02 and 29.50±1.26 µM) and HT-29 (IC$_{50}$: 25.91±1.12 and 20.32±1.23 µM) cell lines and flow cytometric analysis also revealed that 13a arrests both cells lines at G0/G1 phase while 13b induced G0/G1 arrest only in the HT-29 cells. Furthermore, molecular interaction studies also revealed that 13a and 13b exhibited its capacity of being a plausible Bcl-2 and CDK2 (cell division protein kinase 2) inhibitor respectively.

Sabour et al.[119] designed, synthesized and evaluated a new series of 3-cyanopyridine derivatives for their targeting survivin and Bcl-2/Bax. Among them, the compound 14 showed most active cytotoxic and 17.92 times more potent than 5-fluoro uracil drug. Cell cycle analysis of the most potent 14 (fig. 15) revealed that cell cycle arrest at the G2/M phase with an increase in pre G1 apoptotic cells and was subjected to apoptosis studies to estimate their apoptotic potency in which it decreases the expression of the apoptosis suppressor Bcl-2 and increases in the level of apoptosis inducer Bax. The in vitro cytotoxicity of the synthesized compound 14 was evaluated against a panel of cell lines: PC-3, HepG-2 and MDA-MB-231 with IC$_{50}$ value 0.42±0.05, 1.22±0.06 and 0.66±0.03 µM respectively and in normal cell: WI-38 (Normal human fetal lung fibroblast cell line) with IC$_{50}$ value 193.15±0.09 µM.

**Topoisomerase (Topo) inhibitors:**

Human DNA topoisomerase (Topo) is a nuclear enzyme and key component of the cell which can explain all topological issues associated with several vital cellular processes like DNA replication, transcription, repair, recombination, chromatin assembly and chromosome segregation[120,121]. It comprises of two subtypes Topo I and Topo II, since the discovery of Topo by J. C. Wang in 1971, have been vitally prominent molecular targets
for the development of antitumor drugs\cite{122,123}. Human Topo I make DNA single strand cleavage at a time while Topo II which is in the presence of Magnesium (Mg) (II) and adenosine triphosphate (ATP) hydrolysis, cleaves both strands of the DNA double helix to complete their catalytic functions\cite{124,125}. Furthermore, Topo II has two isoforms: Topo IIα and Topo IIβ, both plays an essential role in chromosome segregation and proper organization\cite{126} and these isoforms have been considered as more imperative molecular target than Topo I for designing of antitumor agents\cite{127}.

Lee et al.\cite{128} designed, synthesized and biological evaluated a series of thiochromeno[2,3-c]quinolin-12-one derivatives for their Topo inhibition. Among them, compound 15 showed full inhibitory activities against Topo I and Topo IIα with IC\textsubscript{50} values of 1.14±0.04, 0.22±0.04, 0.80±0.21, 0.53±0.31 and 0.53±0.08 μM in MDA-MB-231, MDA-MB-468 (Breast carcinoma cell line), MCF-7, HCT-116 and H1299 (Lung carcinoma cell line) cells respectively. Additionally recognized compound 15 as a most potent dual Topo inhibitor with low toxicity to normal cells and SAR study fig. 16) also revealed that the terminal amino group of N-2-aminoethylamino or N-3-aminopropylamino at the 6th position and 8,10-di-halogen substituents on thiochromeno[2,3-c]quinolin-12-one are crucial for the Topo inhibitions and tumor-killing activities.

Jin et al.\cite{129} designed, synthesized and evaluated a series of new quinoline and quinolinium iodide derivatives to discover potential antitumor and antibacterial agents. The compound 16 was found to be the most potent derivative with IC\textsubscript{50} values of 4.45±0.88, 4.74±0.42, 14.54±1.96 and 32.12±3.66 μM against A549, Hela, SGC-7901 (Gastric cancer cell line) and L-02 (Liver cancer cell line) cells respectively, stronger than the positive control by 5-fluoro uracil and Methotrexate. Furthermore, compound 16 also had the most potent bacterial inhibitory activity. Additionally, the docking result suggested there were numerous crucial interactions between the 4-position of compound 16 and human Topo I, among which two main H-bonds were formed between the amino (NH) groups or aryl-O-aryl interacts with the catalytic amino acid residue of Dystroglycan (DG) 112 and Lysine (Lys) 425, respectively (fig. 17).

**Tubulin polymerization inhibitors:**

Microtubules are involved in numerous vital cellular functions in various eukaryotic cells such as cell growth,
motility, division, intracellular trafficking and the ability to adapt several shapes to interact with the surrounding environment\cite{130,131}. Microtubule targeting agents are a major class of tumor chemotherapeutic drugs\cite{132,133}. Those agents bind to heterodimers of α and β tubulin which are the building blocks of cellular microtubules. There is a growing evidence displaying that interfering with microtubules normal dynamic equilibrium can engage the spindle roadblock and the arrest of cell cycle progression leading to cell death\cite{134-136}.

Jian et al.\cite{137} were synthesized and biologically evaluated a new pyrazolo[3,4-b]pyridine bridged analogues of combretastatin A-4 possessing 3,4,5-trimethoxylphenyl groups for their anti-proliferative and tubulin polymerization inhibitory activities. Among all these analogs, the most active analogue 17 was found to induce MCF-7, MDA-MB-231, HeLa and Kyse150 (Esophageal squamous cancer cell line) cells with \textit{in vitro} cytotoxicity (IC$_{50}$±standard deviation (SD)) as 27.22±2.31, 27.04±6.42, 18.08±1.48 and 62.82±2.52 μM, respectively. Moreover, molecular modeling studies showed that derivative 17 most possibly occupies the colchicine site of tubulin (fig. 18).

Mirzaei et al.\cite{138} synthesized and evaluated a series of quinoline-chalcone hybrids to discover a promising tubulin inhibitors and anticancer activity. Among all

\[ \text{PDB ID: 1TL8 (Topo-I) and 2XCT (Topo-II)} \]

\[ \text{Alkyliodine substituted at the 1-position in pyridine with the 4-position N-phenyl substituent possessed the highest dual activity against all cancer cell and strains tested} \]

\[ \text{H-bond interaction with Lys425} \]

\[ \text{H-bond interaction with base DG112} \]

\[ \text{H-bond interactions with TPC11} \]

\[ \text{Cation-□ interactions with Arg1122 and Arg458} \]

\[ \text{Fig. 17: Docked structure and SAR of quinoline derivative} \]

\[ \text{PDB ID: 5lyj} \]

\[ \text{This group established two H-bonds with } \text{□Asd258 (2.5 Å) and } \text{□Cet259 (2.6 Å)} \]

\[ \text{Oxygen of this trimethoxyphenyl formed a strong H-bond with } \text{□Cys241 (2.1Å)} \]

\[ \text{Fig. 18: Docked structure of pyrazolo [3,4-b] pyridine-bridged analog} \]
quinolones, compound 18 revealed as the most potent anti-proliferative activity with IC\textsubscript{50} values 2.32±0.84, 2.615±0.63, 4.96±1.46, 2.32±0.46 and 4.44±1.28 against A2780 (Ovarian carcinoma cell line), A2780/RCIS (Ovarian carcinoma cell line), MCF-7, MCF-7/MX (Breast carcinoma) and HUVEC cells respectively. It is also recognized as tubulin inhibitors and induced more arrest at G2/M phase in four cancer cell lines compared to other synthesized compounds. Furthermore, SAR study, molecular dynamics simulation and molecular docking studies of compound 18 into the colchicine binding site of tubulin showed the possible interaction with the active site of tubulin (fig. 19).

**Cytochromes P450 (CYPs) inhibition:**

Cytochromes P450 (CYPs) is a superfamily of hemo proteins involved in many oxidation and reduction reactions on both endogenic and xenobiotic compounds\textsuperscript{[130]}. In general the CYP isoenzyme superfamily consists of 57 CYP genes and 58 pseudogenes arranged into 18 families and 43 subfamilies\textsuperscript{[140]}. For instance, the CYP1 family is included of 3 members: CYP1A1 (cytochrome P450 Family 1 Subfamily A Member 1), CYP1A2 (cytochrome P450 Family 1 Subfamily A Member 2) and CYP1B1 (cytochrome P450 Family 1 Subfamily B Member 1). The latter is primarily expressed in extra hepatic mesodermal cells including steroidogenic tissues such as ovaries, testes and adrenal glands and in steroid responsive tissues such as breast, uterus and prostate\textsuperscript{[141]}. In human eighteen (18) CYP gene families, including over 50 enzymes are found\textsuperscript{[139]}. Despite the importance of other CYP isoforms like CYP3A4 and CYP2D6 are responsible in the metabolism of 50 % of clinical drugs\textsuperscript{[142]}.

Mohamed et al.\textsuperscript{[144]} synthesized and evaluated a new class of hetero steroids derivatives for their anticancer activities. It was observed that compounds 20a (IC\textsubscript{50}: 38.99, 28.88 and 21.98 μM) and 20b (IC\textsubscript{50}: 41.19, 31.99 and 23.91 μM) showed the highest cytotoxic effects against all tested HepG-2, Huh-7 (Liver cancer cell line) and A549 cell lines. Furthermore, molecular simulation also revealed the activity of the tested compounds against four different proteins (CDK2, CYP19 (aromatase), janus Kinase 2 and Bcl2) which are vastly implicated in tumor regulation and progression. Compound 20a and 20b found that were indicated by lowest binding energy (compound 20a=−11.3, −9.2, −11.3 and −9.8 kcal/mol and compound 20b=−10.5, −11.8, −12.0 and −9.7 kcal/mol) compared to reference ligand (fig. 21).

**Miscellaneous:**

Vadukoot et al.\textsuperscript{[145]} reported the synthesis, SAR and biological evaluation of a series of 1H-pyrrolo[2,3-b] pyridine-2-carboxamide derivatives as selective and potent phosphodiesterase type 4B (PDE4B) inhibitors. Among all, compound 21 is a PDE4B preferring PDB ID: 4O2B

**Fig. 19: Docked structure and SAR of quinoline-chalcone derivative**

The benzoyl group in 6 or 8 position showed the most cytotoxic activity

This trimethoxy phenyl group exhibited strong inhibitory activity

Two H-bonds formed by the carbonyl group of chalcone moiety with residues Asn\textsubscript{101} and Lys\textsubscript{254}
inhibitor with IC$_{50}$ value 0.14 and 0.88 μM for PDE4B and PDE4D respectively and displayed acceptable in vitro absorption, distribution, metabolism and excretion (ADME) and significantly also inhibited tumour necrosis factor alpha (TNF-α) release from macrophages showed to pro inflammatory stimuli. In a radio ligand binding assay of PDE isoforms for percentage inhibition at 10 μM, compound 21 was selective against all tested isoforms with <50 % inhibition, except PDE3B with about 87 % inhibiton (fig. 22).

Lin et al.$^{[146]}$ designed, synthesized and evaluated a new C-X-C chemokine receptor type 4 (CXCR4) antagonists based on an aminoquinoline template. A different potent CXCR4 antagonists were recognized, exemplified by compound 22, which revealed excellent binding affinity with CXCR4 receptor (IC$_{50}$=57 nM) and inhibited C-X-C motif chemokine 12 (CXCL12) induced cytosolic calcium released (IC$_{50}$=0.24 nM). Moreover, compound 22 potently inhibited CXCL12/ CXCR4 mediated cell migration in a transwell invasion assay. SAR study also revealed it as potent inhibitor, as mentioned in fig. 23.

Xi et al.$^{[147]}$ designed, synthesized and evaluated a series of new 3-(thiophen-2-ylthio)pyridine derivatives to discover promising multi target antitumor agents. Distinctively compound 23 (fig. 24) with IC$_{50}$ values for HepG-2 is 2.98±1.11 μM and WSU-DLCL2 (B-cell lymphoma cell line): 4.34±0.84 μM, demonstrated good inhibitory activities against Fibroblast growth factor receptor 2 (FGFR2), FGFR3, EGFR, Janus kinase and receptor originated from Nantes (RON). Additionally, the cell cycle analysis showed that compound 23 could arrest HepG-2 cells in the G1/G0 phase.

Gao et al.$^{[148]}$ designed, synthesized and evaluated a series of 6-substituted pyrrolo[2,3-b]pyrimidine derivatives as non-classical antifolate (dihydrofolate reductase (DHFR)) and potential anticancer inhibitors. In preliminary anti-proliferation assay of all compounds revealed submicromolar to nanomolar inhibitory effects against KB (keratin forming tumor cell line) tumor cells, whereas compounds 24a-24c also exhibited nanomolar anti proliferative activities towards SW620 (Colon cancer cell line) and A549 cells. In particular,
Compounds 24a-24c were considerably more potent with IC_{50} value for compound 24a: 0.66, 0.43 and 0.22, compound 24b: 0.78, 0.76 and 0.81 and compound 24c: 0.47, 1.52 and 0.56, against KB, SW620 and A549 cells line, respectively. The growth inhibition induced cell cycle arrest at G1 phase with S phase suppression. Moreover, molecular modeling studies also suggested two binding modes of the target compounds with DHFR (fig. 25).

Mizojiri et al.\textsuperscript{[149]} designed, synthesized and biologically evaluated a series 1H-pyrrolo[2,3-b]pyridine-3-carboxamide derivatives for their Acetyl-coenzyme-A carboxylase1 (ACC1) inhibitory activity. Initially identified 1-methyl-1H-pyrrolo[3,2-b] pyridine-3-carboxamide derivative as a potent ACC1 inhibitor, because of it having pharmacokinetic and physicochemical issues, this led to discovery of 1-isopropyl-1H-pyrrolo[3,2-b]pyridine-3-carboxamide derivative compound 25 as a promising ACC1 inhibitor.
which showed potent ACC1 inhibition as well as sufficient cellular potency with IC_{50} value of 150 nM against HCT-116 cells. SAR study also revealed it as potent and promising ACC1 inhibitor, as mentioned in fig. 26.

**CONCLUSION**

This review study brought out pyridine as an advantageous and privileged moiety in the field of pharmaceutical sciences with its broad range of anticancer profile including numerous targets to throw enormous scientific knowledge for development and designing of newer pyridine derivatives or analogs. The thoroughgoing declaration of SAR/molecular docking will surely help the scientific society to produce effective and potent drugs with excellent pharmacological activity.

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**Conflict of Interests:**

The authors declared no conflict of interest.

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