We are IntechOpen, the world’s leading publisher of Open Access books
Built by scientists, for scientists

3,700 Open access books available
108,500 International authors and editors
1.7 M Downloads

154 Countries delivered to

Our authors are among the

TOP 1% most cited scientists
12.2% Contributors from top 500 universities

WEB OF SCIENCE™
Selection of our books indexed in the Book Citation Index
in Web of Science™ Core Collection (BKCI)

Interested in publishing with us?
Contact book.department@intechopen.com

Numbers displayed above are based on latest data collected.
For more information visit www.intechopen.com
Chapter 10

Apoptotic Molecular Advances in Breast Cancer Management

Pontsho Moela and Lesetja R. Motadi

Abstract
Breast cancer is the most common cancer type amongst women, accounting for most female cancer deaths second to cervical cancer worldwide. It is, therefore, highly crucial to understand the molecular biology and explore other pathways involved in carcinogenesis in order to select appropriate treatment not only for breast cancer but for other cancers as well. Cancer progression is favoured by DNA damage and in most cases a consequent disruption of the apoptotic pathway, thus leading to uncontrolled cell proliferation. Therefore, current therapeutic strategies aim at targeting the apoptotic pathways in order to combat cancer. In this manuscript, we discuss the ways in which evasion of apoptosis during carcinogenesis occurs and the types of current therapeutic strategies as well as promising future approaches against breast cancer.

Keywords: Breast cancer, apoptosis, small molecules, p53, RBBP6

1. Introduction
The human body is composed of trillions of cells that behave and function to provide structure of the body, convert nutrients into energy and carry out specialised functions [1, 3]. Growing, dividing, differentiating and dying are the cells’ behavioural mechanisms to maintain tissue homeostasis [3]. However, molecular disturbances that disrupt this balance may potentially lead to disease. Such molecular disturbances include mutations, among others, during which any change to the DNA sequence might result in abnormality in the cell or tissue [4]. With a population of more than a trillion cells, the human body is prone to mutations that may give one cell a selective advantage of growing and dividing more vigorously to become a growing mutant clone [4, 5]. Such mutations, in which a mutant clone of cells grows and divides out of control at an expense of neighbouring wild-type cell populations, serve as a prerequisite for the development of cancer [3].

© 2015 The Author(s). Licensee InTech. This chapter is distributed under the terms of the Creative Commons Attribution License (http://creativecommons.org/licenses/by/3.0), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.
Cancer is defined as uncontrolled cell proliferation that leads to the formation of abnormal cells and invasion of other adjacent tissues [1, 3–5]. The migration of cells from the origin of tumour to another part of the body is referred to as metastasis. Tumours can either be malignant or benign. While malignant tumours have the ability to invade surrounding tissue, benign tumours cannot invade other tissues and are therefore not as life-threatening [3]. Efficient treatment against malignant tumours is therefore necessary in cancer management. In this chapter, we discuss current anticancer strategies that are targeted on the apoptosis pathway in breast cancer management.

In order to understand breast cancer, it is necessary to understand the normal anatomy of the female breast [3, 20]. The female breast is made up of milk-producing glands called lobules which are connected to ducts that transport milk from the glands to the nipples. The ducts and lobules are surrounded by connective tissue, fatty tissue, blood vessels and lymphatic vessels. In most cases, breast cancer starts in cells surrounding the ducts or the lobules [23]. Metastatic breast cancer is as a result of migration of cancerous cells from ducts and/or lobules via lymphatic vessels to the lymph nodes of the lymphatic system [3, 20, 23].

Breast cancer is the most common cancer type amongst women accounting for many cancer deaths, second to cervical cancer. Risk factors of breast cancer are divided into non-modifiable and modifiable factors [39]. Advanced age, female gender, menarche before the age of 12, menopause after the age of 45, genetic mutations and family history are the major non-modifiable risk factors associated with breast cancer [6, 11, 26, 46, 56, 57]. Breast cancer risk factors that can be controlled include hormone replacement therapy, oral contraceptives, pregnancy, breast feeding and high breast density [31]. Behavioural and life-style risk factors associated with the development of breast cancer include poor diet, i.e. high fat, low vegetable/fruit, low fibre and high in simple carbohydrates; overweight and obesity; and decreasing physical activity [29, 39].

Nearly 80% of human breast cancers are hormone-positive (estrogen and progesterone), followed by human epidermal growth factor receptor 2 (HER2)-positive, then vascular endothelial growth factor (VEGF)-positive breast tumours [8, 9]. Targeting estrogen receptor (ER) pathway, VEGF and HER2 are the long-established breast cancer therapeutic approaches responsible for the improvements of breast cancer prevention and treatment. However, resistance to these endocrine and cell-growth-inhibiting treatments is the main drawback that reduces the benefits of these novel treatment approaches [8, 9, 20, 23]. It is therefore highly crucial to understand the molecular biology and explore other pathways involved in carcinogenesis in order to select appropriate treatment not only for breast cancer but for other cancers as well. In this chapter we discuss different ways of targeting apoptosis in breast cancer management.

2. Targeting apoptosis in breast cancer treatment

During the process of breast cancer progression, normal cells transform into malignant types as a result of genetic alterations [12]. This leads to dysregulation of cellular processes such as
angiogenesis, cell cycle and apoptosis [17]. Therefore, current therapeutic strategies aim at targeting these pathways, more especially apoptosis, in order to combat cancer [18]. Apoptosis is a form of programmed cell death in which cells are programmed to die if found to be cellular damaged [21, 25]. Apoptosis is made up of two major pathways called the death receptor pathway and the mitochondrial pathway, which are both propagated by a caspase cascade that ultimately leads to apoptosis induction [27, 34]. Evasion of apoptosis during carcinogenesis occurs by three distinct mechanisms: disrupted signalling of death receptors, loss of caspase activity as well as impaired balance between anti-apoptotic and pro-apoptotic proteins [14, 42, 50, 59]. Targeting the caspase cascade, Bcl-2 family proteins as well as other factors associated with apoptosis signalling have thus become the major strategy in anticancer therapeutics (table 1).

<table>
<thead>
<tr>
<th>Reagent</th>
<th>Target</th>
<th>Technology</th>
<th>Function</th>
<th>Status</th>
</tr>
</thead>
<tbody>
<tr>
<td>Apoptin</td>
<td>Caspases in the extrinsic pathway</td>
<td>Vector-based</td>
<td>Caspase 3 and 8 activation</td>
<td>Preclinical</td>
</tr>
<tr>
<td></td>
<td></td>
<td>(adenoviral and virus vectors)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Flavipirodol, gossypol,</td>
<td>Anti-Bcl-2 family proteins</td>
<td>Small molecule</td>
<td>Inhibit Bcl-2 family proteins by reducing their expression</td>
<td>Phase I/II</td>
</tr>
<tr>
<td>depsipeptide, ABT-737,</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>ABT-264, fenretinide, HA</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>14-1, GX15-070</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>ABT 737</td>
<td>Anti-apoptotic proteins</td>
<td>Small molecule</td>
<td>Inhibit expression of anti-apoptotic proteins such as Bcl-xL, Bcl-2 and Bcl-W</td>
<td>Phase I</td>
</tr>
<tr>
<td>Oblimersen Sodium</td>
<td>Anti-Bcl-2 targeted drug</td>
<td>Antisense</td>
<td>Bcl-2 antisense increases survival rates in chronic myeloid leukaemia patients when combined with chemotherapy</td>
<td>Phase II</td>
</tr>
<tr>
<td>ONXY-015 drug</td>
<td>p53-based gene therapy</td>
<td>Adenoviral</td>
<td>Genetically engineered adenovirus that has been modified to infect and lyse p53-deficient cells</td>
<td>Phase III</td>
</tr>
<tr>
<td>CD8+ cytotoxic T-lymphocytes (CTLs)</td>
<td>Tumour associated antigens (mutant p53)</td>
<td>Vaccine</td>
<td>Recognize TAA-derived peptides that are processed and presented on the tumours cell surface in association with MHC class I molecules, leading to killing of tumour cells</td>
<td>Phase I</td>
</tr>
</tbody>
</table>
Table 1. Apoptosis-based anticancer drugs in development

<table>
<thead>
<tr>
<th>Reagent</th>
<th>Target</th>
<th>Technology</th>
<th>Function</th>
<th>Status</th>
</tr>
</thead>
<tbody>
<tr>
<td>Phikan083, CP-31398</td>
<td>p53-targeted</td>
<td>Small molecule</td>
<td>Restores p53 function by intercalating with p53-bound DNA and destabilising the p53-DNA interaction</td>
<td>Phase I/II</td>
</tr>
<tr>
<td>Tenovins, Nutlins, MI-219</td>
<td>p53-MDM2 interaction</td>
<td>Small Molecule</td>
<td>Interrupt the p53-MDM2 interaction to prevent inactivation of p53 by MDM2</td>
<td>Phase I/II</td>
</tr>
<tr>
<td>siMDM2, siE6/7, siBBP6</td>
<td>p53-MDM2, p53-E6, p53-RRBP6 interaction</td>
<td>Liposomal encapsulated synthetic with its negative siRNA</td>
<td>Interrupt p53 interaction Research</td>
<td>Research</td>
</tr>
</tbody>
</table>

2.1. Caspase-targeted therapy

Pathogenic as they are, disruptions in the apoptotic pathway provide compelling possible strategies for the treatment of breast cancer and other related types of cancers [59]. Therapeutic agents designed to re-establish the normal functioning of the apoptotic signalling pathways have the potential to get rid of over 50% of human cancers including breast cancer [34]. Novel drug discoveries in recent years have led to promising advances in the treatment of breast cancer as well as other cancers. For example, the caspase-targeting therapies that use small molecules to act as caspase activators have been identified [24, 32]. These small molecule caspase activators are pro-apoptotic due to their characteristic arginine-glycine-aspartate motif that enables them to directly convert non-active procaspase-3 into active caspase-3 thus leading to apoptosis induction. Apoptotin is a caspase-based drug therapy that has the ability to induce caspase activity thus increasing apoptosis induction [32]. MCF-7 breast cancer cells completely lack the expression of caspase-3 due to frame-shift mutation in exon 3 of the caspase-3 gene [13]. As a result, caspase-based gene therapy that relies on caspase-3 gene delivery techniques in order to up-regulate caspase-3 expression in caspase 3-deficient breast cancers has been invented. In human liver tumorigenesis, caspase-3 gene therapy led to a significant increase in apoptosis and shrinkage in tumour size when combined with other chemotherapeutic drugs [13, 32]. Caspase-8 expression has also been found to be impaired due to hypermethylation in several cancer cells. In small cell lung carcinomas, demethylation treatments have been shown to sensitise these cancer cells to drug-induced apoptosis [32, 53].

2.2. Anti-Bcl-2 therapy

The mitochondrial pathway is down-regulated by the anti-apoptotic Bcl-2 family proteins [19, 22, 40, 43, 60]. Drug-based therapy using anti-Bcl-2 small molecules has led to a significant
induction of apoptosis in several cancers. Flavipirodol, gossypol, depsipeptide, ABT-737, ABT-264, fenretinide, HA 14-1 and GX15-070 are some of the small molecules that inhibit BCl-2 by reducing their expression [41, 45, 59]. Small molecules with the ability to mimic pro-apoptotic or anti-apoptotic BH3-only Bcl-2 family proteins in order to induce apoptosis have also been designed [2, 41]. This class of drugs that imitate BH3-only pro-apoptotic and anti-apoptotic Bcl-2 family proteins is referred to as BH3-only mimetic drugs [2, 35]. ABT 737 is one example of the BH3-only mimetics that has been shown to inhibit expression of anti-apoptotic proteins such as Bcl-xl, Bcl-2 and Bcl-W; and is showing promising results in clinical trials [2, 59]. The first anti-Bcl-2 targeted drug to enter clinical trials in leukemic patients is known as oblimersen sodium [41, 59]. This Bcl-2 antisense has been shown to increase survival rates in chronic myeloid leukaemia patients when combined with chemotherapy [41, 59].

2.3. p53-based gene therapy

The loss of p53 function is a common feature in almost all human cancer including breast cancer [37, 43, 47]. Because of this, there is a lot of interest in targeting p53 for anticancer therapeutic drugs [7, 10, 16, 55]. The first biological approach which is now widely used in targeting p53 is gene delivery of wild-type p53 into tumour cells using adenoviral or retroviral techniques [28, 48]. p53-based gene therapy is however not effective on its own in killing cancer cells and for this reason combinational therapies involving other modes of treatments in the presence of p53 therapy are being investigated [10, 55, 59].

For example, it was discovered that concurrent treatment using adenoviral-mediated wild-type p53 injection with ionising therapy significantly reduces tumour size in cancers of prostate, brain and spine as well as head and neck [28, 59]. Elimination of p53-defective cells using synthetic viruses designed to infect and kill cancer cells is another breakthrough in p53-based gene therapy [28, 48, 59]. One example is the ONYX-015 drug, which is a genetically engineered adenovirus that has been modified to infect and lyse p53-deficient cells [28]. Genetic alterations that take place in p53 during tumorigenesis can trigger the immune responses in both T- and B-cells [10]. This provides yet another interesting platform for p53-based anticancer therapy, and a number of p53-based vaccines are currently undergoing clinical trials [10, 59].

2.4. Small molecule approach in p53-based drug therapy

In comparison to large biological drugs that are present with complex structures, small molecular drugs are organic compounds designed to be extremely low in molecular weight and are made up of well-defined chemical structures that enable them to pass through the cell membrane when taken orally. A further advantage of small molecule drugs over biologics is that they are stable, mostly non-immunogenic and it is easy to characterise their molecular composition and heterogeneity. The mode of action for small molecules relies on their binding to specific biopolymers such as proteins and nucleic acids and act as effectors to alter function or activity of the specific biopolymer.
In cancer, small molecules are used to restore mutated proteins back to their wild-type forms and induce activity of proteins responsible for elimination of tumorigenic cells [30]. In p53-based drug therapy, several small molecules that can restore the function of mutated p53 have been investigated. One example of a small molecule drug known is CP-31398; which has been shown to restore p53 function by intercalating with p53-bound DNA and destabilising the p53-DNA interaction [30]. Another small molecule called Phikan083, which is a derivative of carbazole, has been identified as one of the small molecules that has the ability to restore mutant p53 too. The most advanced of these small molecules are those that act by interrupting the p53-MDM2 interaction which is responsible for the inactivation of wild-type p53 [51, 52, 54]. These include the nutlins, tenovins and the MI-219 [52]. MDM2 acts as a negative regulator of p53 by binding to and inactivating the function of p53. This activity results in the loss of p53-mediated apoptosis in cancer cells, thus promoting carcinogenesis. While the MI-219 small molecular drugs are responsible for the destabilisation of the MDM2-p53 interactions in order to selectively induce apoptosis and inhibit apoptosis, nutlins disrupt the MSM2-p53 complex and selectively induce senescence [51, 52, 54].

2.5. siRNA-based p53 therapy

There are certain cancers with no mutations in p53 but in which non-mutated p53 might be down-regulated by certain p53 negative regulators [30]. In these cancers, development of specific siRNAs for silencing of the negative regulatory genes is often used to activate p53 [10, 30, 55]. MDM2 E3 ligase and the viral E6 protein are two extensively studied negative regulators of p53 that are associated with cancer progression [51, 52, 54].

Under normal cellular conditions, p53 tumour suppressor gene is kept under tight regulation by the MDM2-p53 auto-regulatory feedback loop [36, 51, 52]. In response to stress stimuli such as DNA damage or radiation, activated p53 interacts with genes responsible for the induction of cell cycle arrest or apoptosis (figure 1) [36]. During cancer development, the interaction between p53 and MDM2 mediates p53 interaction with the ring finger domain of the MDM2 ubiquitin ligase for degradation of the p53 tumour suppressor protein [54]. This event compromises the occurrence of cell cycle arrest and p53-mediated apoptosis and facilitates abnormal cell proliferation. The use of MDM2-specific siRNA to disrupt the p53-MDM2 interaction in breast cancer cells has been shown to induce apoptosis, inhibit cell proliferation and lead to decreased tumour size [36, 51, 52, 54].

The E6 viral protein is another thoroughly studied p53 negative regulator in HPV (human papillomavirus)-related cancers such as anogenital, cervical, head and neck cancers. During HPV infection, E6 protein expression increases in order to facilitate HPV replication and viral integration into the host cell. The E6 protein achieves this outcome by using its E3 ligase Hect domain to bind to and degrade the cellular tumour suppressor proteins p53 and pRB, thus abrogating the host cells’ potential to initiate cell cycle arrest and apoptosis. Therapeutic strategies to disrupt E6-p53 interactions in the form of antisense and siRNA application specific to E6 viral protein have received the most attention in HPV-related cancer therapeutics [10, 30, 55].

A third ubiquitous protein suspected to be yet another negative regulator of active p53 especially in breast cancer progression is known as retinoblastoma binding protein 6 (RBBP6)
[36]. RBBP6 is a 250kDa protein that has been shown to interact with and possibly lead to the degradation of p53 tumour suppressor gene since it possesses an E3 ligase activity [44, 49]. Its mRNA codes for a p53-binding domain as well as other domains known as DWNN domain, zinc finger domain and a ring finger domain, which are responsible for the ubiquitous nature of RBBP6 [44]. Besides the p53 domain, which is only present in human RBBP6, the above-mentioned domains are conserved in about all eukaryotic organisms such as humans, plants, protozoa, fungi, microsporidia and the single-celled parasite Encephalitozoon cuniculi [44]. RBBP6 is a spliced-associated protein and therefore exists in different other homologues known as PACT and P2P-R [44, 58].

A critical insight into the role played by RBBP6 in certain cancers via p53 has been elucidated [15]. Transfection of lung cancer cells with siRBBP6 led to a decrease in RBBP6 expression whereas sip53 transfection led to an increase in RBBP6 expression and, according to this study, RBBP6 may be involved in the degradation of p53 thereby enhancing abnormal cell prolifer-
ation [38]. In one study, it was demonstrated that down-regulation of the PACT homologue of RBBP6 in mice induces embryonic lethality with a consequent accumulation of p53 and a widespread apoptosis [33]. In addition to identifying PACT as a negative regulator of p53, further discoveries suggest that PACT knockdown enhances p53-Hdm2 interaction thus reducing p53 poly-ubiquitination by RBBP6 [33].

In recent studies, it was found that silencing RBBP6 gene in MCF-7 and CAMA-1 cells led to p53 up-regulation and sensitised the breast cancer cells to apoptosis induction [36]. Concurrent treatment of these cells with apoptosis-inducing agents, camptothecin or staurosporine, further increased apoptosis induction [36]. Furthermore, up-regulation of bax as a result of the co-treatment provided early insights into the possible mechanism behind the observed apoptosis [36]. Taken together, it is suspected that RBBP6 silencing may be responsible for the identified p53 up-regulation in breast cancer and other cancers and that the observed apoptosis is more likely p53-dependent; however, further in vivo investigations would validate these observations.

3. Conclusions

Taken all together, it is evident that anticancer therapeutics primarily depend on apoptosis pathway activation in breast cancer and several other cancers. However, a few milestones still need to be reached with regard to this novel anti-tumour molecular approach. For example, most of the experimentally studied apoptosis-inducing regimens in breast cancer cells have not reached the clinical stages. Another important factor that needs to be addressed in apoptosis-targeted therapy is to determine whether the observed cytotoxicity of breast cancer cells in experimental settings is comparable in clinical settings. Moreover, understanding tumour biology of individual cancer patients can help select therapeutic interventions that are highly specific to a presented tumour. Nonetheless, the link between apoptosis and tumorigenesis has been thoroughly investigated in breast cancer and has led to lots of promising strategies that attempt to eradicate cancer cells by targeting the apoptosis signalling pathway.

Author details

Pontsho Moela¹ and Lesetja R. Motadi²*

*Address all correspondence to: lesetja2007i@webmail.co.za

1 School of Molecular and Cell Biology, University of the Witwatersrand, Johannesburg, South Africa

2 Department of Biochemistry, North-West University (Mafikeng Campus), Potchefstroom, Sout, Africa
References


cytotoxic T lymphocytes by recombinant adenoviral vector based vaccination in mice, but not man. *Gene Ther*; 9(13):833-843


