Balasingam, Nirujah

2017

Elucidating the role of human obg like ATPase1 (hOLA1) in apoptosis

Department of Chemistry and Biochemistry

https://hdl.handle.net/10133/5173

Downloaded from OPUS, University of Lethbridge Research Repository
ELUCIDATING THE ROLE OF HUMAN OBG LIKE ATPASE 1 (HOLA1) IN APOPTOSIS

NIRUJAH BALASINGAM
Bachelor of Science, University of Mysore, 2014

A Thesis
Submitted to the School of Graduate Studies
of the University of Lethbridge
in Partial Fulfilment of the
Requirements for the Degree

MASTER OF SCIENCE

Department of Chemistry and Biochemistry
University of Lethbridge
LETHBRIDGE, ALBERTA, CANADA

© Nirujah Balasingam, 2017
ELUCIDATING THE ROLE OF HUMAN OBG LIKE ATPASE 1 (HOLA1) IN APOPTOSIS

NIRUJAH BALASINGAM

Date of Defense: December 07, 2017

Dr. N. Thakor                                Assistant Professor    Ph.D.
Supervisor

Dr. H. J. Wieden                             Professor              Ph.D.
Co-Supervisor

Dr. M. Roussel                               Professor              Ph.D.
Thesis Examination Committee Member

Dr. T. Russell                               Associate Professor     Ph.D.
Thesis Examination Committee Member

Dr. A. Gorrell                               Associate Professor     Ph.D.
External examiner
University of Northern British Columbia
British Columbia

Dr. P. G. Hayes                              Professor              Ph.D.
Chair, Thesis Examination Committee
This thesis is dedicated to my beloved family,

I am grateful for my Parents and Siblings for the endless love and faith in me.

Special thanks to my brother for being my pillar of strength, a constant source of knowledge and inspiration for me.

Thank you, Anna,

I am also grateful to my sister-in-law for the support and guidance

& finally, the joy of my life,

My cute little Nephews and Niece

Without you all I am nothing.
Abstract

hOLA1 is a purine nucleotide binding protein that belongs to the P-loop GTPase family. hOLA1 suppresses protein synthesis by limiting ternary complex formation thus promoting the integrated stress response (ISR). ISR determines cell fate by activating or inhibiting pro- and anti-apoptotic proteins depending on the type of the stress condition. ISR facilitates cell survival in response to mild stress and promotes apoptosis in response to chronic stress. Depletion of hOLA1 leads to increased cell survival and diminished ISR. This study investigates the role of hOLA1 in apoptosis. My hypothesis was that diminished ISR in hOLA1-depleted cells leads to upregulation of anti-apoptotic proteins thus inhibiting apoptosis. Indeed, this study shows that hOLA1 depletion upregulates anti-apoptotic proteins such as cIAP1, cIAP2, and Bcl-xL, thus inhibiting apoptosis. The inhibition of apoptosis was studied by assessing caspase activation, cleaved poly (ADP-ribose) polymerase (PARP) level, and measuring apoptosis by propidium iodide (PI) staining and flow cytometry.
Acknowledgment

First, I would like to thank my supervisor Dr. Nehal Thakor for his constant guidance and support throughout this journey. Next, special thanks to my co-supervisor Dr. Hans-Joachim Wieden for the collaboration.

I wish to thank my committee members Dr. Marc Roussel and Dr. Tony Russell for the valuable comments and guidance. Finally, I would like to thank my lab members for the great support during my master’s program.
# Table of Contents

List of Tables ........................................................................................................... vii  
List of Figures ......................................................................................................... viii  
Abbreviations .......................................................................................................... ix  

Chapter 1: Introduction  
1.1 Overview of eukaryotic gene expression ......................................................... 1  
1.1.1 Cap-dependent translation initiation ......................................................... 1  
1.1.2 Cap-independent translation ........................................................................ 5  
1.1.3 Regulation of translation and integrated stress response ......................... 6  
1.2 Activation and termination of the ISR ............................................................ 8  
1.3 Apoptosis and programmed cell death ............................................................ 11  
1.3.1 Mitochondrial or intrinsic apoptotic pathway ........................................... 12  
1.3.2 Extrinsic pathway ....................................................................................... 13  
1.3.3 Regulation of apoptosis .............................................................................. 15  
1.4 Role of translational control in cancer ........................................................... 17  
1.5 Human obg like ATPase 1 (hOLA1) ............................................................... 18  
1.5.1 hOLA1 as a regulator of protein synthesis and ISR ..................................... 19  
1.5.2 hOLA1 as a cell cycle regulator Regulation of apoptosis ......................... 20  
1.6 Aim of the project ........................................................................................... 22  

Chapter 2: Materials and methods  
2.1 Cell lines ........................................................................................................ 23  
2.2 Transfection .................................................................................................... 23  
2.3 Western blotting .............................................................................................. 24  
2.4 alamarBlue™ assay ....................................................................................... 26  
2.5 Propidium iodide (PI) staining and flow cytometry ....................................... 26  
2.6 Immunoprecipitation and mass spectrometry ............................................. 27  
2.7 Western blot quantification ............................................................................ 29  

Chapter 3: Results  
3.1 hOLA1 depletion leads to upregulation of key anti-apoptotic proteins .......... 30  
3.2 hOLA1 depletion leads to decreased caspase activation in the MDA-MB231 cell line ........................................................................................................... 41  
3.3 hOLA1 depletion does not have considerable effects on the inhibition of cell death ........................................................................................................ 45  
3.4 Measurement of apoptosis in hOLA1-depleted cells ..................................... 48  
3.5 Co-immunoprecipitation and mass spectrometry to identify the protein(s) interacting with hOLA1 ................................................................. 51  

Chapter 4: Discussion and future perspective  
4.1 Discussion ....................................................................................................... 55  
4.2 Future perspectives ........................................................................................ 60  

References ............................................................................................................. 62
List of Tables

Table 2.1: Antibodies used in this study.................................................................25
Table 3.1: Mass spectrometry data analysis of sample I (MW ~45 kDa)......................53
Table 3.2: Possible interacting partner of hOLA1......................................................54
List of Figures

Figure 1.1: Schematic representation of cap-dependent translation initiation .................. 4
Figure 1.2: Schematic diagram of apoptotic pathway ................................................. 15
Figure 1.3: hOLA1 as regulator of protein synthesis, possible mechanism suggested by Chen, H et al ................................................................. 20
Figure 3.1: Optimization of the concentration of si-hOLA1 for the depletion of hOLA1 ... 32
Figure 3.2: Optimization of si-hOLA1 transfection duration time for the effective depletion ............................................................... 33
Figure 3.3: hOLA1 depletion leads to diminished ISR ............................................... 34
Figure 3.4: hOLA1 depletion leads to increased level of cIAP1 in MDA-MB231 and HEK-293T cell lines .................................................. 36
Figure 3.5: hOLA1 depletion leads to increased level of cIAP2 in MDA-MB231 and HEK-293T cell lines .................................................. 36
Figure 3.6: hOLA1 depletion does not have a considerable effect on the level of cFLIP_L in MDA-MB231 and HEK-293T cell lines ........................................ 39
Figure 3.7: hOLA1 depletion leads to increased levels of Bcl-xL in MDA-MB231 and HEK-293T cell lines .................................................. 40
Figure 3.8: hOLA1 depletion does not have a considerable effect on the levels of XIAP ................................................................. 41
Figure 3.9: hOLA1 depletion leads to decreased caspase activation ................................. 45
Figure 3.11: hOLA1 depletion and apoptotic induction do not have considerable effects on the inhibition of cell death in MDA-MB231 cell line ............................................. 47
Figure 3.11: hOLA1 depletion and apoptotic induction do not have considerable effects on the inhibition of cell death in HEK-293T cell line ............................................. 48
Figure 3.12: hOLA1 depletion leads to decreased apoptosis .......................................... 50
Figure 3.13: Co-immunoprecipitation to identify the protein(s) interacting with hOLA1 ............................................................................. 53
List of Abbreviations

ATF4 - Activating transcription factor-4
Bcl-xL - B-cell lymphoma-extra large
cFLIPL - Cellular FLICE-like inhibitory protein
CHOP - C/EBP homologous protein
cIAP1 - Cellular inhibitor of apoptosis 1
cIAP2 - Cellular inhibitor of apoptosis 2
CReP - Constitutive repressor of eIF2α phosphorylation
eIF2 - Eukaryotic initiation factor 2
FADD - Fas-associated protein with death domain
GADD34 - Growth arrest and DNA damage-inducible protein
hOLA1 - Human obg like ATPase
IAPs - Inhibitors of apoptosis
PIC - Pre-initiation complex
RIPA - Radioimmunoprecipitation assay
TC - Ternary complex
TGS - ThrRS, GTPase, SpoT
TNFα - Tumor necrosis factor α
TRADD - Tumor necrosis factor receptor type 1-associated DEATH domain protein
TRAIL - TNF-related apoptosis-inducing ligand (TRAIL)
XIAP - X-linked inhibitor of apoptosis protein
Chapter 1: Introduction

1.1 Overview of eukaryotic gene expression
The deoxyribonucleic acid (DNA) encodes protein molecules that carry out all essential functions of life.1 The portion of a gene that encodes for a protein is called the coding region.2,3 Transcription is the process of copying the genetic information stored in the DNA to a ribonucleic acid (RNA) molecule by RNA polymerase enzymes.1 Once the primary transcripts are made they undergo post-transcriptional modifications such as 5’ capping, polyadenylation, and splicing to form mature mRNA.4 Capping is the addition of 7-methylguanosine (m7G) at the 5’ end of the mRNA.4 Polyadenylation involves the addition of around 250 adenine residues at the 3’ end.4 Splicing is the removal of the non-coding regions in the mRNA.4 Once the mature mRNA is made, it is transported to the cytoplasm. In the cytoplasm, mature mRNA is translated into proteins using the translation machinery.5 Translation is a fundamental cellular process which occurs in three different phases: initiation, elongation, and termination.5

1.1.1 Cap-dependent translation initiation
The initiation of translation involves several distinct steps leading to assembly of the 80S initiation complex.6 Notably, initiation is highly regulated and a rate-limiting step of translation.6 During initiation, the start codon is recognized and base-paired to initiator tRNA (Met-tRNAi) in the P site of 40S ribosomal subunit.5,7 The initiation process involves three major steps: formation of the 43S pre-initiation complex, formation of the 48S pre-initiation complex, and formation of the 80S ribosome.6 The first step of translation is the formation of ternary complex (TC; eIF2•GTP•Met-tRNAi) through the interaction of initiator tRNA with the eIF2 and guanosine triphosphate (GTP).8 eIF2 is a GDP/GTP
binding protein complex consisting of three subunits; regulatory subunit-α, tRNA/mRNA binding subunit-β, and guanine nucleotide binding subunit-γ. Formation of the 43S initiation complex involves interaction of the TC with the 40S ribosomal subunit and other initiation factors, namely 1, 1A, and 3 (Figure 1.1). The 43S initiation complex is stabilized by eIF1 and eIF3. eIF1A promotes assembly of 43S preinitiation complex and ribosomal scanning along with eIF1 to recognize the start codon. The 5’ m7G cap of cellular mRNA enhances the translation initiation through the interaction with cap-binding complex eIF4F. eIF4F is a heterotrimeric factor consisting of three initiation factors: RNA helicase eIF4A, cap-binding protein eIF4E, and molecular scaffolding protein eIF4G. The eIF4F complex binds to the 5’ cap structure along with eIF4B, whereas Poly(A)-binding protein (PABP) binds to the poly-A tail at the 3’ end of the mRNA. PABP has the ability to interact with eIF4G. This results in the circularization of mRNA. Circularization of mRNA enhances the recruitment of 43S pre-initiation complex (PIC) to the mRNA and formation of the 48S initiation complex. The RNA helicase eIF4A helps in unwinding 5’ mRNA secondary structures and facilitates ribosomal scanning to identify the initiation codon. Once the initiator tRNA is delivered and base-paired with the start codon, eIF5 induces the hydrolysis of GTP by eIF2. Subsequently, eIF5B facilitates the joining of the 40S and 60S ribosomal subunits. During this process eIF5B interacts with eIF1A on the C-terminus of eIF1A.

After the 80S initiation complex has formed, the translation proceeds to the elongation phase, where polypeptide chain formation occurs. During elongation, the elongation factor 1A (eEF1A) helps to recruit the next aminoacyl tRNA to the A site. Peptide bond formation between the adjacent amino acids is catalyzed by the peptidyl transferase activity
of the ribosome.\textsuperscript{15, 16} Elongation continues until the stop codon is encountered at the A site of the ribosome. Once the stop codon is recognized by eukaryotic releasing factor 1 (eRF1), eRF1 binds to eRF3 and GTP.\textsuperscript{15} Subsequently eRF3 induces the hydrolysis of GTP to promote peptide chain release from the ribosomal complex. Subsequently, ATP-binding cassette subfamily E member1 (ABCE1) promotes ribosomal dissociation and releases the factors by an energy-dependent mechanism.\textsuperscript{15} The released ribosomal subunits and eIFs are recycled and used in the next cycle of translation initiation.\textsuperscript{15}
Figure 1.1: Schematic representation of cap-dependent translation initiation: The 40S subunit along with TC and other factors form the 43S pre-initiation complex (PIC). 43S is believed to scan 5’ UTR of the mRNA to find a start codon in the right context. Following the Met-tRNAi delivery 48S pre-initiation complex is formed. Upon GTP hydrolysis and the release of eIFs, 60S ribosomal subunit unites with 40S subunit to form 80S ribosomal complex.
1.1.2 Cap-independent translation

The above described molecular mechanism of translation is also called the canonical or cap-dependent translation. However, during stress, cap-dependent translation is inhibited by cleavage or modulation of several initiation factors. During such stress conditions a subset of mRNAs continue to be translated via cap-independent translation initiation mechanisms. Internal ribosome entry site (IRES)-mediated translation initiation is an example of cap-independent translation initiation.

An IRES is a RNA element that has distinct secondary or tertiary structures and allows the translation of an mRNA through its interaction with few eIFs and trans-acting factors, without the involvement of the 5’ cap structure. IRES-mediated translation was first identified in the positive-stranded RNA poliovirus. Poliovirus infection leads to proteolytic cleavage of eIF4G, resulting in the loss of its interaction with eIF4E and eIF3, preventing the formation of the eIF4F complex, and thus inhibiting cap-dependent translation. Although cap-dependent translation of cellular mRNA is inhibited, poliovirus RNA translation was observed to be stimulated during these conditions. Poliovirus translation is conducted through a mechanism independent of eIF4F and the m⁷G cap structure. This cap-binding complex-independent mechanism of translation initiation was later termed as IRES-mediated translation. These viral IRESs were classified based on the RNA sequence and structural similarities and the factors required for translation initiation. For example, picornavirus IRESs classified into three types based on the different characteristics such as structure and the factors required for translation. Type I and type II requires eIF4G and eIF4A. Type III IRESs require eIF4G and only occurs in hepatitis A virus.
Cellular IRESs were identified in stress response genes that encode for pro- and anti-apoptotic proteins: B cell lymphoma extra-large (Bcl-xL), apoptotic protease-activating factor 1 (APAF1), B-cell lymphoma 2 (Bcl-2), Bcl-2 associated athanogene1 (Bag-1), cellular inhibitor of apoptosis protein 1 (cIAP1), X-linked inhibitor of apoptosis protein (XIAP).\textsuperscript{23-26} These pro-survival and death proteins are activated due to internal and external stress stimuli such as DNA damage, viral infection, hypoxia, nutrient deprivation and endoplasmic reticulum (ER) stress.\textsuperscript{24, 26} During stress conditions several signaling pathways are activated to regulate protein synthesis in order to recover cellular homeostasis.\textsuperscript{26-28}

1.1.3 Regulation of translation during the integrated stress response

During stress condition, translation is regulated by activation or inhibition of initiation factors such as eIF4E homologous protein (4E-HP), eIF4E-binding proteins (4E-BPs) and eIF2.\textsuperscript{29, 30} As mentioned below, these eIFs either regulate the recruitment of 40S subunit onto the mRNA or the recruitment of tRNA into the P site of 40S ribosomal subunit. Canonical translation initiation is extensively regulated by the 5’mRNA cap-binding protein eIF4E.\textsuperscript{31} The cap recognition by eIF4E is hindered due to binding of 4E-HP to the cap structure.\textsuperscript{32, 33} The 4E-HP protein could recognize cap structure, however, it lacks the ability to interact with eIF4G leading to the inhibition of protein synthesis.\textsuperscript{34} In addition, the eIF4G-eIF4E interaction in the eIF4F complex is inhibited by 4E-BPs.\textsuperscript{32, 33} 4E-BPs inhibit cap-dependent translation by competing with eIF4G for the shared binding site on eIF4E.\textsuperscript{33} The interaction of 4E-BP with eIF4E is regulated by phosphorylation of 4E-BP. Hypo-phosphorylation of 4E-BP leads to its stronger binding with eIF4E, whereas hyper-phosphorylation of 4E-BP leads to weakening of the interaction with eIF4E.\textsuperscript{33}
Phosphorylation of 4E-BP is regulated by a signaling pathway called mechanistic target of rapamycin (mTOR) pathway.\textsuperscript{35}

One of the key regulatory mechanisms during stress is phosphorylation of the α subunit of eIF2 complex.\textsuperscript{30, 36} eIF2 mainly exists in either: GTP- bound or GDP- bound states.\textsuperscript{37} During translation initiation, the inactive eIF2-GDP complex is recycled back to an active eIF2-GTP complex to regenerate the TC by a guanine nucleotide exchange factor called eIF2B.\textsuperscript{37} During stress conditions, the α-subunit of eIF2 is phosphorylated at serine 51 by one of the four eIF2α kinases (discussed below).\textsuperscript{5} Phosphorylation of eIF2α enhances its affinity for eIF2B, leading to the sequestration of eIF2 by eIF2B. Inhibition of active eIF2-GTP regeneration results in decreased TC formation, thus inhibiting overall translation and activating stress response pathway such as integrated stress response (ISR).\textsuperscript{36-38}

ISR is activated in response to various pathological and physiological conditions including both extrinsic and intrinsic stimuli. For example, nutrient deprivation, hypoxia, and ER stress can activate ISR.\textsuperscript{38-41} The common convergence of ISR activation is the phosphorylation of the eIF2α subunit.\textsuperscript{42} Increased phosphorylation of eIF2α leads to decreased cap-dependent translation while allowing the translation of selected mRNAs such as that encoding activating transcription factor 4 (ATF4) through 5’ upstream open reading frames (5’uORF).\textsuperscript{43-45} ATF4 translation is regulated through the involvement of two different uORFs: uORF1 and uORF2.\textsuperscript{46} The uORF1 element contains three amino acids residues in length, whereas uORF2 contains 59 amino acids residues in length that overlaps the ATF4 coding region in an out-of-frame manner.\textsuperscript{47} During normal conditions, ribosome translates uORF1 and resumes ribosomal scanning and re-initiation downstream of the uORF1 leading to the translation of uORF2 and bypassing start codon of ATF4.\textsuperscript{46} During
stress, eIF2α phosphorylation lowers the level of active eIF2-GTP complex which increases the time required for ribosomal scanning and re-initiation.\textsuperscript{47} This leads to a higher probability of re-initiation at the ATF4 coding region thus translating ATF4.\textsuperscript{47} ATF4 aids the cells to recover cellular homeostasis by reprogramming transcription of stress-related genes.\textsuperscript{38, 48, 49}

On the other hand, some of the cellular stress response transcripts are translated through IRES-mediated translation during stress.\textsuperscript{50} For example, XIAP is translated through IRES dependent mechanism during stress condition.\textsuperscript{50} XIAP is known to be encoded by two distinct mRNAs: shorter and longer 5’UTR.\textsuperscript{51} During normal conditions, XIAP is translated through the shorter 5’UTR.\textsuperscript{51} However, during stress the longer IRES-containing 5’ UTR helps in the translation of XIAP.\textsuperscript{51} The combination of cap-dependent translation and IRES-mediated translation helps in the maintenance of XIAP level in response to cellular stress and adaptation.\textsuperscript{51} eIF2α phosphorylation and dephosphorylation help to recover cellular homeostasis during stressed conditions.\textsuperscript{38} However, based on the severity and duration of the stress condition, eIF2α may or may not get dephosphorylated. Hence, ISR plays a dual role in cellular survival, cell death, hence in the ultimate cell fate.\textsuperscript{38}

1.2 Activation and termination of the ISR

The eIF2α can be phosphorylated by four distinct kinases: general control non-derepressible 2 (GCN2), heme-regulated eIF2α kinase (HRI), double-stranded RNA dependent protein kinase (PKR) and PKR-like ER kinase (PERK).\textsuperscript{52-54} These kinases are homologous in their catalytic domain and divergent in their regulatory domains.\textsuperscript{55} Each eIF2 kinase is activated by a distinct type of stress. For example, PKR is activated by the infection of double-stranded RNA viruses.\textsuperscript{56-58} The double-stranded RNA aids in the
dimerization of C-terminal domain of PKR and thus PKR gets activated. Activated PKR phosphorylates eIF2α which results into the inhibition of both viral and host protein synthesis. Notably, other types of stresses also activate PKR independent of dsRNA such as; oxidative stress, ER stress, and growth factor deprivation. The eIF2 kinase PKR is also activated through caspase activity in the early stages of apoptosis (apoptosis is discussed in section 1.3). Activation of caspase-8, -7 and -3 results in the cleavage of eIF2 kinase domain leads to activation of PKR. This results in the phosphorylation of eIF2α thus inhibiting translation. eIF2 kinase GCN2 is a highly conserved from yeast to human and is activated due to amino acid deprivation. During amino acid starvation, the levels of amino acetylated tRNA is decreased. GCN2 senses the increased pool of uncharged tRNA and gets activated. Also, UV light is reported to activate GCN2 through an unrevealed mechanism. Like PKR activation, GCN2 activation leads to the phosphorylation of eIF2α and attenuation of the global protein synthesis.

PERK is localized in the endoplasmic reticulum (ER) and activated due to several conditions including the accumulation of unfolded proteins in the ER, changes in the calcium homeostasis, and the oxidative stress. In normal conditions, the luminal domain of PERK is bound to glucose-regulated-protein 78 (GRP78). During ER stress PERK is activated by two different mechanisms. It is activated through the dissociation of ER luminal bound GRP78 due to the accumulation of unfolded proteins in the ER lumen. PERK is also activated by direct binding of an unfolded protein to its luminal domain.

HRI is mostly expressed in erythroid cells and is involved in erythrocyte differentiation. HRI kinase activity is regulated by heme through the formation of a disulfide bond between two monomers and preventing dimerization. HRI can be activated by many stresses.
independent of heme but with the support of heat shock proteins 70 and 90 (HSP70 and 90).\textsuperscript{67} Importantly, eIF2 kinase family members have interconnections; they can act together for similar stresses.\textsuperscript{68} For example, GCN2 and PERK can compensate for each other under a variety of stress conditions such as viral infection and ER stress.\textsuperscript{68, 69}

As mentioned earlier, besides inhibition of global protein synthesis, phosphorylation of eIF2\textalpha also induces the translation of transcriptional activator ATF4.\textsuperscript{70, 71} Increased translation of ATF4 induces the expression of the transcription factor C/EBP homologous protein (CHOP) and ATF3.\textsuperscript{72, 73} Together these protein factors activate ISR-CHOP signaling.\textsuperscript{74} CHOP promotes apoptosis through upregulation of BH3-only pro-apoptotic Bcl-2 family members.\textsuperscript{75} CHOP also regulates the expression of the pro-apoptotic protein death receptor 5 (DR5) protein and pro-apoptotic genes such as apoptotic protease activating protein 1 (APAF1).\textsuperscript{76} Moreover, ATF4 itself can promote apoptosis through direct activation of the pro-apoptotic-protein Bcl-2.\textsuperscript{77} Further, during ISR the inhibitors of apoptosis proteins cIAP1, cIAP2, Livin, Survivin, NAIP, and XIAP are downregulated upon chronic PERK signalling.\textsuperscript{78}

In addition to eIF2\textalpha kinases, phosphatases play a critical role in executing ISR.\textsuperscript{79} The eIF2\textalpha dephosphorylation is a key point for the termination of ISR to recover global protein synthesis and cellular homeostasis.\textsuperscript{2, 79} The eIF2\textalpha dephosphorylation is implemented by the phosphatase activity of the constitutive repressor of eIF2\textalpha phosphorylation (CReP), and growth arrest and DNA damage-inducible protein (GADD34).\textsuperscript{70, 80} CReP helps unstressed cells to maintain translational homeostasis by maintaining a low level of eIF2\textalpha phosphorylation through its constitutive function as eIF2\textalpha phosphatase.\textsuperscript{80} GADD34 expression is induced in later stages of ISR to dephosphorylate eIF2\textalpha.\textsuperscript{81} GADD34 helps in
the dephosphorylation of eIF2α by interacting with protein phosphatase 1 (PP1). The GADD34-PP1 complex acts as a negative feedback loop during ISR and helps the cells to recover homeostasis. Furthermore, dephosphorylation of eIF2α by GADD34 also helps the cells to allow the translation of the stress-responsive mRNAs such as HSP70. If the cellular homeostasis is failed by phosphatase activity of GADD34, that can facilitate cell death by inducing death inducing-proteins such as DR5 and Bcl-2 family pro-apoptotic proteins.

On the other hand, early/mild ISR can also promote cell survival through negative regulation of apoptosis. PERK-mediated ISR activation resulted in increased levels of cIAP1 and cIAP2 in both cancer and non-cancer cell lines and promote cellular survival. As mentioned earlier, the consequences of eIF2α phosphorylation differ based on the type of the stress condition such as chronic or acute and downstream effect of ATF4 translation and other factors.

### 1.3 Apoptosis or programmed cell death

The balance between cell survival and cell death pathways is important for maintaining cellular homeostasis. Cell death can be executed by programmed mechanisms such as apoptosis and necroptosis. Apoptosis is an energy requiring mechanism that occurs normally to maintain tissue/organ homeostasis during development and aging. Additionally, a wide variety of stress stimuli such as DNA damage and viral infection induce apoptosis. Apoptosis can be characterized by morphological changes such as cell shrinkage, nuclear condensation, membrane blebbing or biochemical features such as caspase (cysteine-aspartic proteases) cleavage and DNA fragmentation. Apoptosis is mediated by intracellular enzymes called caspases. Fourteen different caspases have been
found in humans so far, out of which caspases-2, 3, 6, 7, 8, 9, and 10 are known to play crucial roles in apoptosis. These enzymes are expressed as inactive proenzymes. Caspases consist of prodomain p20 large subunit and a p10 small subunit. Activation of these proenzymes requires a proteolytic cleavage between small and large subunits that results in heterodimer active caspases. Based on the length of their pro-domain, caspases are classified into two groups: initiator caspases (caspases 8, 9, 10) and executioner caspases (caspases 3, 6, 7). Initiator caspases contain a long pro-domain that comprises a protein-protein interacting domain, caspase recruiter domain (CARD) and death effector domain (DED). Upstream adaptor proteins such as fas-associated protein with death domain (FADD) and tumor necrosis factor receptor type 1-associated DEATH domain protein (TRADD) interact with caspases through CARD and DED domains. In contrast, effector caspases containing a short pro-domain are classically activated by upstream initiator caspases. The activation of executioner caspases leads to subsequent cleavage of multiple cellular proteins and execution of cell death. Apoptosis can be triggered by many stimuli and is executed through two major pathways namely the extrinsic pathway, and intrinsic (mitochondrial) pathway.

1.3.1 Mitochondrial or intrinsic pathway

The mitochondrion is an important cellular organelle that is critical for energy production and deciding cell fate. The intrinsic apoptotic pathway is activated through various intracellular signals such as DNA damage or nutritional deprivation leading to the mitochondrial outer membrane permeabilization (MOMP). In normal condition, the Bcl-2 family protein Bax is found in the cytosol as monomers or loosely attached to the outer membrane of the mitochondria. Another member of Bcl-2 family, Bak, is anchored into
the outer mitochondrial membrane in its monomeric form. During apoptosis, Bax is localized to the mitochondrial outer membrane. Bak and Bax oligomerize and form pores on the outer mitochondrial membrane. This leads to release of mitochondrial proteins such as cytochrome c (cyt c), endonuclease D (EndoG), HtrA2/Omi, apoptosis-inducing factor (AIF), and second mitochondria-derived activator of caspase/direct IAP binding protein (Smac/DIABLO) into the cytosol. Cytochrome c (cyt c) forms a complex in the cytoplasm with adaptor protein APAF1. Subsequently, the APAF1-cyt c complex forms a heptameric structure through oligomerization. Heptameric structure formation of APAF1-cyt c complex permits the interaction of procaspase-9 with APAF1. This interaction leads to the activation of Caspase-9. When caspase-9 is activated, it activates effector caspases (3 and 7) to execute cell death (Figure 1.3).

On the other hand, mitochondrial proteins Smac/DIABLO and HtrA2/Omi promote caspase activation by blocking IAP-mediated caspase inhibition.

1.3.2 Extrinsic pathway

The extrinsic pathway is initiated through the interaction of the death receptors present on the cell surface with their ligands. The extracellular receptor ligands, such as tumor necrosis factor α (TNFα), TNF-related apoptosis-inducing ligand (TRAIL)/Apo2L, and Fas ligand (FasL)/CD95L bind to their corresponding membrane receptors TNFR, DR4/DR5, and Fas/CD95 respectively. Death receptor contains a specific amino acid sequence termed death domain (DD). Death domain is crucial for the transduction of apoptotic stress stimuli from extracellular receptors. Once the receptors and the ligand bind together, a cytoplasmic adapter protein is recruited onto the receptor-ligand complex through DD. The pro-apoptotic ligand, receptor, pro-caspases and DD recruitment lead to the formation
of a new complex called death-induced signaling complex (DISC) and subsequent activation of initiator caspases.\textsuperscript{103, 104} Activation of initiator caspases-8 and -10 leads to the cleavage of executioner caspases such as caspases-3 and -7 (Figure 1.3).\textsuperscript{104} Notably, caspase-8 can also cleave BH3-only pro-apoptotic protein Bid to form truncated Bid (t-Bid) leading to MOMP and subsequent activation of the intrinsic apoptosis pathway.\textsuperscript{105}

As mentioned below, TRAIL and TNFα execute apoptosis through distinct components of the extrinsic apoptosis pathway. The major differences between TRAIL and TNFα induced apoptosis are the receptors and cytosolic adaptor proteins.\textsuperscript{106} TRAIL facilitates FADD recruitment onto the receptor upon binding to the appropriate receptors (DR4 or DR5) leading to the activation of caspase-8 and caspase-10.\textsuperscript{107} On the other hand, TNF-α induces TNFR signaling leading to both apoptotic and non-apoptotic responses through the formation of two different complexes: complex I and complex II.\textsuperscript{107} The complex I consists of TRADD, TRAF2, cIAPs, and receptor interacting protein-1 (RIP-1) to the activation of nuclear factor kappa-light-chain-enhancer of activated B cells (NF-κB) in order to promote cellular survival.\textsuperscript{106} When complex I dissociates from RIP I and associate with procaspase-8 and FADD results in the formation of complex II in the cytosol. Complex II formation leads to the activation of the apoptotic pathway.\textsuperscript{106, 107}
Figure 1.3: Schematic diagram of the apoptotic pathways. **Intrinsic pathway:** Intrinsic apoptosis is triggered by genotoxic stress. The major event is the recruitment of Bax and Bak on the outer mitochondrial membrane that causes the release of cytochrome c. Cytochrome C binds to APAF1 to form the apoptosome complex. The apoptosome activates caspase-9 that leads to the activation of caspase 3/7 and eventually executes apoptosis.

**Extrinsic pathway:** extrinsic apoptosis is triggered by external stimuli by binding of ligands (FasL, APO-2L, TRAIL, TNFα) to appropriate receptors (Fas, DR4, DR5, TNF-R1). Subsequent activation of procaspases-8 and -10 by recruiting adaptor protein FADD leads to the activation of caspase-8 or -10. This leads to the activation of executioner caspase-3, -6, -7 to mediate apoptosis. Also, there is a cross-talk between the intrinsic and extrinsic pathways. Caspase-8 can cleave Bid to form tBid leading to activation of the intrinsic apoptotic pathway.

### 1.3.3 Regulation of apoptosis

Apoptosis is regulated by anti- and pro-apoptotic proteins. Proteins corresponding to the negative regulation of apoptosis are called anti-apoptotic proteins. One of the main groups of anti-apoptotic proteins are the inhibitors of apoptosis proteins (IAPs), originally identified in the Baculovirus for their ability to regulate the enzymatic activity of caspases. Later, IAPs were identified in mammals as well as fruit flies. There are eight different human analogs from the IAP families: X-linked inhibitor of apoptosis
XIAP, Apollon/Bruce, cellular inhibitor of apoptosis protein1 (cIAP1), cellular inhibitor of apoptosis2 (cIAP2), Survivin, IAP-like protein 2 (ILP2), melanoma IAP (ML-IAP/Livin) and neuronal apoptosis inhibitory protein (NAIP). The important hallmark of IAP is that they all have at least one baculoviral IAP repeat (BIR) domain. With the exception of Survivin, IAPs have one or more functional domains either a really interesting new gene (RING) domain, or a caspase recruitment domain (CARD), or both. The RING domain possesses E3 ligase activity allowing the self-regulation of IAP by proteasomal degradation, whereas the CARD domain is involved in protein-protein interaction. Caspases are inhibited by BIR2 and BIR3 domains. The BIR2 domain specifically targets caspase-3 and caspase-7 whereas BIR3 domain targets caspase-9 and degraded by ubiquitylation.

XIAP is crucial among IAPs that suppress both extrinsic and intrinsic apoptotic pathways. It contains BIR2 and BIR3 domains to inhibit an upstream caspase (caspase-9) and downstream caspases (caspase-3 and 7). The BIR2 domain interacts with the N-terminal small subunit of caspase-3 and caspase-7 to prevent substrate binding. On the other hand, the BIR3 domain binds to the IAP binding motif of caspase-9. Also, XIAP can inhibit apoptosis via proteolytic degradation by E3 ligase activity of its RING domain. cIAP1 and cIAP2 are homologs that inhibit the extrinsic pathway through their interactions with TRAF1 and TRAF2. cIAP1 and cIAP2 contain three BIR domains, a CARD domain, and a RING domain. In addition, cIAP1 and cIAP2 inhibit apoptosis by activating the NF-κB pathway. Caspase inhibition by IAPs are antagonized by Smac/DIABLO family mediated poly-ubiquitylation. Although Survivin contains one
BIR domain, it does not inhibit caspases, whereas Livin contains one BIR domain that inhibits caspase-3 and 9.\textsuperscript{112, 113, 119} An additional group of anti-apoptotic proteins are the Bcl-2 family anti-apoptotic members (Bcl-2, Bcl-X, and Bcl-xL) that are associated with the outer mitochondrial membrane and prevent apoptosis.\textsuperscript{120} The Bcl-2 family has pro-apoptotic members as well, such as Bak, Bid and Bax that facilitate the release of cytochrome c into the cytosol and thus activate the intrinsic apoptotic pathway.\textsuperscript{120, 121} The anti-apoptotic protein of extrinsic apoptotic pathway is cellular-FADD-like IL-1β-converting enzyme-inhibitory protein (cFLIP).\textsuperscript{122} There are several splice variants of cFLIP. However, only three are expressed at the protein level: c-FLIP long (c-FLIP\textsubscript{L}), c-FLIP short (c-FLIP\textsubscript{S}) and a short variant c-FLIP\textsubscript{R}.\textsuperscript{122} All three isoforms of cFLIP can be recruited to the DISC complex through death effector domain (DED, a subclass of DD).\textsuperscript{122} cFLIP inhibit caspase-8 activation at the DISC by competing with caspase-8 to bind to FADD.\textsuperscript{123} Dysregulation of anti- and pro-apoptotic proteins have been reported in many type of cancers: pancreatic cancer, melanoma and lymphoma.\textsuperscript{124} This underlines the importance of studying detailed mechanisms of apoptotic regulators.\textsuperscript{112} 

### 1.4 Role of translational control in cancer

Dysregulation of gene expression at the translation level has been implicated in oncogenesis and cancer progression.\textsuperscript{125} As mentioned earlier, translation is mostly regulated at the initiation stage.\textsuperscript{126} Overexpression of the initiation factors such as eIF3, eIF4G, eIF5A is observed in many types of malignancies.\textsuperscript{25} Also, some initiation factors can act as both tumor inducer and as tumor suppressors.\textsuperscript{25} For example, the h subunit of eIF3-complex is overexpressed in breast cancer whereas the f subunit of the eIF3-complex
is observed to be decreased in melanoma and pancreatic cancer.\textsuperscript{127, 128} The crucial regulatory factors of cap-dependent translation, eIF4E, and eIF4G are overexpressed in breast cancer and lung carcinoma.\textsuperscript{129, 130} During the conditions when global protein synthesis is dysregulated by changes in the level of initiation factors, translation could rely on the cap-independent mechanism.\textsuperscript{6, 18} For example, during stress, genes that promote survival and tumor growth such as Bcl-2, VEGFA are translated via IRES-mediated mechanisms.\textsuperscript{25} Besides initiation factors other factors like \textit{trans}-acting factors, miRNAs and nucleotide-binding proteins also known to regulate protein synthesis.\textsuperscript{131, 132}

1.5 Human obg like ATPase1

Human obg like ATPase1 (hOLA1) is a nucleotide-binding protein that has numerous crucial roles in various cellular processes like translation, cell motility, cell division and cytoskeleton organization.\textsuperscript{133, 134} hOLA1 belongs to the phosphate-binding loop nucleotide triphosphatase (P-loop NTPase) group.\textsuperscript{133} The P-loop GTPase is a subclass of nucleotide-binding protein that facilitates the hydrolysis of GTP to regulate fundamental cellular processes such as cell division, signal transduction and translation.\textsuperscript{133} P-loop GTPases are further classified into two major groups based on structural features and conserved sequences: translation factor related class (TRAFAC) and signal recognition particle.\textsuperscript{133} P-loop GTPases contain a common core G-domain with five conserved motifs.\textsuperscript{133} These motifs are responsible for several functions such as GTP binding, GTP positioning, guanine nucleotide recognition, and magnesium (Mg\textsuperscript{2+}) coordination.\textsuperscript{133} These properties of the G domain provide conformational flexibility to the protein during nucleotide hydrolysis.\textsuperscript{133} Additionally, the central G domain of OLA1 is flanked on either side by a coiled-coil domain and a TGS (ThrRS, GTPase, SpoT) domain.\textsuperscript{133} P-loop GTPases are further
classified into four families: protein secretion factors, Era-related GTPases, Elongation factors (EF-G, EF-Tu, Lep A, and IF2) and Obg-related family (Obg and YchF).\textsuperscript{133} hOLAl is an ATP and GTP binding protein that belongs to obg1 family of P-loop GTPase.\textsuperscript{134} It was first identified and characterized in 2007.\textsuperscript{134} hOLAl is reported to have a higher affinity for ATP compared to GTP.\textsuperscript{134} Therefore, hOLAl was named as human Obg like ATPase1.\textsuperscript{134} hOLAl is abnormally expressed in different type of cancers, suggesting the importance of studying the role of hOLAl in stress conditions.\textsuperscript{135} Indeed, hOLAl is known to play numerous roles during stress conditions as discussed below.\textsuperscript{135-137} hOLAl is a negative regulator of antioxidant response.\textsuperscript{135} hOLAl prevents ubiquitylation-mediated degradation of HSP70.\textsuperscript{137} Also, hOLAl plays an important role in centrosome regulation through the interaction with breast cancer associated gene1 (BRCA1) and γ-tubulin in breast cancer cell lines.\textsuperscript{138, 139} In addition, hOLAl was also referred to as the DNA damage-regulated overexpressed in cancer 45 (DOC45) because it is implicated in DNA damage response.\textsuperscript{140} hOLAl is a protein synthesis, cell cycle, and ISR regulator.\textsuperscript{136} The next two sections describe how hOLAl regulates protein synthesis, ISR, and cell cycle.

1.5.1 hOLAl as a regulator of protein synthesis and ISR.

hOLAl is reported to inhibit protein synthesis and ISR by inhibiting TC formation.\textsuperscript{136} hOLAl is known to suppress protein synthesis by limiting the formation of TC.\textsuperscript{136} Importantly, hOLAl is known to interact with eIF2.\textsuperscript{136} In addition, Chen, H \textit{et al.} reported that hOLAl depletion leads to greater TC level and diminished ISR.\textsuperscript{136} Moreover, Chen, H \textit{et al.} also reported that hOLAl depletion leads to increased cellular survival and a diminished CHOP level.\textsuperscript{136} Further, \textit{in vivo} analysis revealed that hOLAl depletion leads to deficient ISR, tumor growth and metastasis.\textsuperscript{136} Chen, H \textit{et al.} suggested a mechanism
for the role of hOLA1 in regulating protein synthesis, they believed that hOLA1 facilitates the hydrolysis of eIF2-bound to GTP leading to the formation of an inactive eIF2-GDP complex, which is unable to bind to Met-tRNAi, thus inhibiting TC formation (Figure 1.3).

Figure 1.3: hOLA1 as regulator of protein synthesis, possible mechanism suggested by Chen, H et al. Chen, H et al. suggested a possible mechanism for the regulation of protein synthesis by hOLA1.\textsuperscript{136} hOLA1 is known to interact with eIF2 and thus possibly inhibiting TC formation, therefore Chen, H et al. suggested that hOLA1 inhibit TC formation by facilitating the hydrolysis of GTP bound to eIF2 into eIF2-GDP, thus inhibiting global protein synthesis.\textsuperscript{136}

1.5.2 hOLA1 as a cell cycle regulator

The cell cycle is a key cellular function that leads to the cell division.\textsuperscript{141} One of the main cause of cancer is dysregulation of the cell cycle.\textsuperscript{142, 143} Cyclins and cyclin-dependent kinases (CDKs) are essential enzymes for the transition of the cell cycle from one phase to the next phase in mammalian cells.\textsuperscript{142} The G1 to S phase transition requires cyclins D and E, the G2 to M transition requires the formation of the cyclin B-CDK1 complex by dephosphorylation of CDK1.\textsuperscript{142} This phase transition is inhibited by cyclin-dependent kinase inhibitors: p21, p27, and p53.\textsuperscript{142} hOLA1 is known to regulate the cell cycle by
suppressing CDK2 inhibitor p21. It was demonstrated that p21 is translationally regulated by hOLA1.\textsuperscript{143}
1.6 Aim of the project
As mentioned earlier hOLA1 suppresses protein synthesis by inhibiting TC formation. Moreover, hOLA1 is known to interact with eIF2 and depletion of hOLA1 leads to diminished ISR and enhanced cellular survival. Enhanced cellular survival could be the result of increased cell proliferation or decreased apoptosis. Depending on the type of the stress condition ISR plays an important role in deciding cell fate by regulating anti- and pro-apoptotic proteins. Therefore, I hypothesize that increased survival in hOLA1-depleted cells is due to the upregulation of anti-apoptotic proteins and inhibition of apoptosis. The main objectives of this project are as follows.

1. To elucidate the role of hOLA1 in apoptosis and ISR.

The role of hOLA1 in apoptosis was assessed using knockdown studies and Western blotting. Western blots were used to assess the levels of anti-apoptotic proteins, cleaved caspase, and PARP cleavage. The viability assay (alamarBlue™ assay) was used to measure cell viability. Propidium iodide (PI) staining and flow cytometry were used to quantify apoptosis.

2. To identify the proteins interacting with hOLA1.

As mentioned earlier, hOLA1 is known to regulate protein synthesis by inhibiting ternary complex formation. In addition, hOLA1 is reported to interact with eIF2. However, it is not clear as to how hOLA1 regulates protein synthesis. Therefore, I wanted to assess if there are additional proteins that regulate the interaction of hOLA1 with eIF2. To this end, I performed co-immunoprecipitation and mass spectrometry to identify additional protein(s) interacting with hOLA1.
Chapter 2: Materials and methods

2.1 Cell lines
The human breast cancer cell line: MDA-MB231 was obtained from the children’s hospital of eastern Ontario (CHEO). The human embryonic kidney cell line HEK-293T was purchased from American type culture collection (ATCC). Cells were cultured using Dulbecco's Modified Eagle's medium (DMEM) with 10% (v/v) Fetal Bovine Serum (FBS, Gibco), 50 U/ml Penicillin, 50 μg/ml Streptomycin, and 200 μM L-Glutamine. The cells were cultured in 10 cm Petri dishes under conditions of 5% CO2 and 37°C. Trypsin (Final concentration: 0.25% (wt/v) Trypsin, 1.45 mM NaCl, 0.5 mM EDTA in PBS) was used for detaching cells for sub-culturing.

2.2 Transfection
For hOLA1 knockdown, Human OLA1 cDNA (NM_013341.3) specific siRNA (SASI_Hs01_00244684) and the control siRNA (MISSION siRNA Universal Negative Control #1 SIC001) were obtained from Sigma-Aldrich. Lipofectamine® RNAiMAX transfection reagent (Thermofisher Scientific) was used to deliver the siRNA to the cells; the transfection mix was prepared according to the manufacturer’s instructions. 250 μl of Opti-MEM, 6 μl of Lipofectamine® RNAiMAX and 50 nM (final concentration) of siRNA were added and incubated for 20 minutes at 37°C in the incubator. The cells were stained with 0.4% (wt/v) Trypan Blue (Thermofisher, Catalog number: 15250061) and counted using hemocytometer, MDA-MB231 cells (1.5×10^5) and HEK-293T cells (1.0×10^5) were seeded in the 6-well plates. Following cell seeding, transfection mix was added, and the plate incubated at 37°C in an incubator with the 5% CO2 for 96 hours.
2.3 Western blotting

Cells were harvested in RIPA (Radio Immuno Precipitation Assay) lysis buffer containing 50 mM Tris-HCl pH 7.4, 1mM EDTA, 15mM NaCl, 1% (v/v) NP40, 0.5% (wt/v) deoxycholic acid, 0.05% (wt/v) SDS, protease inhibitors (Final concentration: 0.01 µg/µl Leupeptin, 0.005 µg/µl PMSF, 0.013 µg/µl Aprotinin, 0.001 µg/µl Pepstatin A) and phosphatase inhibitors (0.209 µg/µl NaF and 3.689 µg/µl NaV). The whole cell lysate was centrifuged using a Sorvall™ ST 8 Small Benchtop Centrifuge at 10,000g for 15 minutes to remove the cell debris. The supernatant was collected, and the protein concentration was quantified using Quick Star™ Bradford reagent as per manufacturer’s protocol (Bio-Rad, Catalog No:5000201). 10 µg of proteins were separated on 12% SDS-PAGE. The proteins were then transferred onto a nitrocellulose membrane. The membrane was blocked with 10% (wt/v) skim milk in PBST (137 mM NaCl, 2.7 mM KCl, 10 mM Na₂HPO₄, and 1.8 mM KH₂PO₄, 0.02% (v/v) Tween) for one hour at room temperature. Following blocking, the membrane was incubated with 10 mL of primary antibody (Diluted in 0.02% PBST with 1% skimmed milk as per manufacturer's instruction, Table 2.1) overnight at 4°C. The next day, blots were washed with PBST (0.02% (v/v) tween) two times and once with PBS (137 mM NaCl, 2.7 mM KCl, 10 mM Na₂HPO₄, and 1.8 mM KH₂PO₄). Subsequently, the blots were incubated with 10 mL of horseradish peroxidase (HRP) conjugated secondary antibody (1:3000 dilution in 1% skimmed milk in PBST) for one hour at room temperature. Subsequently, the bands were visualized using enhanced chemiluminescence (ECL) substrate (Thermofisher Scientific, catalog number: 32106) in Amersham imager 600.
<table>
<thead>
<tr>
<th>Antibody</th>
<th>Dilution</th>
<th>Catalog No</th>
<th>Source</th>
</tr>
</thead>
<tbody>
<tr>
<td>Anti-hOLAl antibody</td>
<td>1:1000</td>
<td>HPA035790</td>
<td>Sigma</td>
</tr>
<tr>
<td>Anti-GTPBP9 antibody</td>
<td>1:1000</td>
<td>ab51077</td>
<td>Abcam</td>
</tr>
<tr>
<td>Anti-cIAP1 antibody</td>
<td>1:1000</td>
<td>ab2399</td>
<td>Abcam</td>
</tr>
<tr>
<td>Anti-cIAP2 antibody</td>
<td>1:1000</td>
<td>ab23423</td>
<td>Abcam</td>
</tr>
<tr>
<td>Anti-XIAP antibody</td>
<td>1:1000</td>
<td>ab21278</td>
<td>Abcam</td>
</tr>
<tr>
<td>Anti-FLIP antibody</td>
<td>1:1000</td>
<td>ab8421</td>
<td>Abcam</td>
</tr>
<tr>
<td>Anti- Bcl-xL antibody</td>
<td>1:1000</td>
<td>ab31396</td>
<td>Abcam</td>
</tr>
<tr>
<td>Anti-ATF-4 antibody</td>
<td>1:1000</td>
<td>ab1371</td>
<td>Abcam</td>
</tr>
<tr>
<td>Anti-CHOP antibody</td>
<td>1:1000</td>
<td>D46F1</td>
<td>Cell signalling</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Technology (CST)</td>
</tr>
<tr>
<td>Anti-rabbit IgG, HRP-linked antibody</td>
<td>1:3000</td>
<td>7074S</td>
<td>CST</td>
</tr>
<tr>
<td>Anti-Caspase-8 antibody</td>
<td>1:1000</td>
<td>9496S</td>
<td></td>
</tr>
<tr>
<td>Anti-Caspase-7 antibody</td>
<td>1:1000</td>
<td>12827S</td>
<td></td>
</tr>
<tr>
<td>Anti-Caspase-9 antibody</td>
<td>1:1000</td>
<td>9502S</td>
<td>CST</td>
</tr>
<tr>
<td>Anti-Caspase-3 antibody</td>
<td>1:1000</td>
<td>9662S</td>
<td></td>
</tr>
<tr>
<td>Anti-Cleaved PARP antibody</td>
<td>1:1000</td>
<td>5625S</td>
<td></td>
</tr>
<tr>
<td>Anti-GAPDH antibody</td>
<td>1:10,000</td>
<td>ab9485</td>
<td>Abcam</td>
</tr>
</tbody>
</table>
2.4 alamarBlue™ assay
MDA-MB231 and HEK-293T cells were transfected with siRNA in a 96-well plate, at 24 hours post-transfection, apoptotic inducers were added: doxorubicin (10 nM), TRAIL (100 ng/mL), TNFα (100 ng/mL). Following 72 hours of apoptotic inducers treatment, 10 µl of alamarBlue™ dye (Resazurin sodium salt in water, final concentration-3 mg/mL, obtained from Sigma) was added. Subsequently, cells were incubated for 16 hours. At the end of 16 hours, the fluorescence absorbance at 600 nm was measured.

2.5 Propidium iodide (PI) staining and flow cytometry
MDA-MB231 cells (1.5×10^5) were seeded and transfected with 50 nM control siRNA and 50 nM si-hOLA1. After 92 hours of transfection, cells were treated with DMSO and 100 ng/mL TRAIL. Cells were trypsinized 96 hours post-transfection (after 4-hours of compound treatment), washed with PBS (137 mM, NaCl, 2.7 mM KCl, 10 mM Na₂HPO₄, and 1.8 mM KH₂PO₄). Cells were then re-suspended in 40 µl PBS. Subsequently, 400 µl of 70% (v/v) cold ethanol was added along the sides of the tube and mixed well. This step was repeated to make the total volume of 1 mL and kept at 4°C overnight. The next day, the cells were centrifuged using a Sorvall™ ST 8 Small Benchtop Centrifuge at 4,000g, 4°C for 5 minutes. The pellet was washed with PBS and centrifuged using a Sorvall™ ST 8 Small Benchtop at 4,000g, 4°C for 5 minutes, then the pellet was re-suspended in 30 µl PBS containing 2 µl of 10 mg/mL RNase. The re-suspended pellet was then incubated at 37°C for 4 hours. Following the incubation, the PI (PI staining solution obtained from Sigma, final concentration 1 µg/µl of PI with 2mM MgCl₂ in 1×PBS) solution was added and incubated at 37°C for 15-30 minutes. Subsequently, cells were submitted for flow cytometry analysis (flow cytometry facility, University of Alberta). Samples were acquired on an Attune NxT cytometer. Single cells (100,000) were acquired from each sample, based
on gating forward scatter (FSC-A vs SSC-A) versus side scatter (cells) and PI-A vs PI-W (single cells). All samples were analyzed using FlowJo v10 Cell Cycle Platform (Watson (Pragmatic) model).

2.6 Immunoprecipitation and mass spectrometry
MDA-MB231 cells were grown in a 10 cm dish for immunoprecipitation. Subsequently, cells were lysed when they reached 80% confluency using 1 mL RIPA lysis buffer (details are given in 2.1 section). The whole cell lysate was centrifuged using a Sorvall™ ST 8 Small Benchtop Centrifuge at 4°C, 10,000g for 15 minutes. The beads for the protein pulldown were obtained from ThermoFisher Scientific (Dynabeads™ protein G for immunoprecipitation, catalog number: 10003D). First, the beads were coated with hOLA1 antibody (Sigma, catalog number: HPA035790) by incubating 40 µl beads with 1 µg antibody overnight at 4°C in a Digital Tube Revolver (Thermo Scientific). Next, the cell lysate was incubated with anti-hOLA1 antibody-coated beads for 3 hours at 4°C in a rocker. The beads were washed with PBS, and the proteins were then eluted using 60 µl 2x Laemmli Sample Buffer (Bio-Rad, catalog number: #1610737). Next, a portion of the sample was assessed by Western blot technique. The membrane was probed with anti-hOLA1 antibody to verify the pull-down, and anti-eIF2α antibody (Cell Signalling Technology, catalog number: #9722). Subsequently, another portion of the sample was analyzed on SDS-PAGE and stained with Coomassie stain for mass spectrometry analysis. Protein bands were then excised from the gel and de-stained. Subsequently, the gel plugs were washed with 150 µl water and incubated in a thermoshaker at 1,050 rpm for 5 minutes. The gel plugs and water were centrifuged at max speed in a mini-centrifuge for 5 minutes and the solution was removed using pipette. Then, the gel plugs were washed with 150 µl
acetonitrile by incubating in a thermoshaker at 1050 rpm for 5 minutes. Subsequently, the gel plugs were incubated with 100 µl of 10 mM dichlorodiphenyltrichloroethane (DDT) at 56°C for 50 minutes. Next, the gel fragments were washed with acetonitrile as mentioned above. Then, the gel fragments were incubated with iodoacetamide for 20 minutes in a thermoshaker at 26°C and 1,050 rpm. Subsequently, the gel fragment was incubated with 150 µL of 100 mM NH₄HCO₃ and incubated at 26°C in a thermoshaker at 1,050 rpm for 5 minutes. Next, samples were centrifuged using a mini-centrifuge at maximum speed. Then, samples were incubated with acetonitrile as described above. The solutions were then removed by pipetting after a quick spin using micro-centrifuge. Subsequently, gel plugs were dried and digested with 15 µl trypsin (12.5 ng/µL Trypsin, 41.7 mM ABC, 4.17 mM CaCl₂) overnight. The next day, the solution was collected by a quick spin using micro-centrifuge and a fine pipette tip. The solution was transferred to a clean 1.5 mL microcentrifuge. Subsequently, gel fragments were washed with acetonitrile as mentioned above. The sample was dried and incubated with 5% formic acid for 15 minutes at 37°C and 1,050 rpm in a thermoshaker. Then, the sample was incubated with 50 µl acetonitrile at 37°C and 1,050 rpm in a thermoshaker. The gel plug was then spun down at maximum speed using a mini-centrifuge. The supernatant was collected and pooled with the supernatant from the digestion step. The combined supernatant was then evaporated using a speed vacuum (ThermoFisher Scientific, Savant™ SpeedVac™). Then, 30 µl of the sample was loaded for mass spectrometry analysis on the ThermoScientific™ Orbitrap Fusion™ Tribrid™ mass spectrometer at the mass spectrometry facility, University of Lethbridge.
2.7 Western blot quantification

The bands were quantified using an Amersham Imager 600 analysis software (GE healthcare). First, the number of lanes were chosen manually, and the bands were detected manually. The background was automatically subtracted by choosing the parameter ‘rubber band’. The same method was followed for the loading control GAPDH. The ratio of the protein of interest and GAPDH was calculated. The quantitative analysis was performed for all three biological replicates. The graph was plotted for si-Control and si-hOLA1 using Graph Pad Prism7 software.
Chapter 3: Results

3.1 hOLA1 depletion leads to upregulation of key anti-apoptotic proteins

As mentioned earlier, hOLA1 is known to suppress global protein synthesis by limiting ternary complex formation.\textsuperscript{136} In addition, hOLA1 depletion is reported to result in a diminished ISR level and increased cellular survival.\textsuperscript{136} ISR is known to play a key role in deciding cell fate through the effector proteins of ISR such as ATF4 and CHOP. These effector proteins are known to regulate several anti- and pro-apoptotic proteins.\textsuperscript{144, 145} Therefore, I hypothesized that the reason for increased cellular survival in hOLA1-depleted cells is due to decreased apoptosis caused by the effect of diminished ISR level on the levels of anti-apoptotic proteins. Therefore, I wanted to assess the effect of hOLA1 removal in the cellular levels of key IAPs and anti-apoptotic proteins.

Breast cancer is one of the most aggressive and heterogeneous diseases.\textsuperscript{146} hOLA1 is abnormally expressed in breast cancer.\textsuperscript{147} One of the well-established breast cancer cell lines is MDA-MB231.\textsuperscript{146, 148, 149} MDA-MB231 is a triple negative cell line, lacking progesterone receptor (PR) expression, human epidermal growth factor receptor 2 (HER2) and estrogen receptor (ER).\textsuperscript{149} Moreover, Chen, H \textit{et al.} have used MDA-MB231 cell line to investigate hOLA1 as a suppressor of protein synthesis and ISR regulator.\textsuperscript{136} Hence, I wanted to study the role of hOLA1 in apoptosis using the MDA-MB231 cell line. First, I optimized the si-hOLA1 concentration and duration time for the effective knockdown with a minimum off-target effects on gene expression. To this end, 1.5×10^5 MDA-MB231 cells were seeded in 6-well plates and transfected with two different concentrations of control siRNA and siRNA targeted against hOLA1 (25 nM, and 50 nM). 96 hours post-transfection, cells were lysed, and the cellular levels of proteins were analyzed by the
Western blot technique. The membrane was probed for hOLA1 and loading control glyceraldehyde 3-phosphate dehydrogenase (GAPDH). It is known that α-actin interacts with hOLA1, therefore GAPDH was used as a loading control. Following the incubation of the membrane with primary and secondary antibodies, the protein bands were visualized using an Amersham Imager 600 (GE healthcare). Subsequently, protein bands were quantified using the Amersham Imager 600 analysis software. Following the quantification, the relative expression was calculated by dividing the expression of hOLA1 by the expression of GAPDH in control siRNA and si-hOLA1 treated cells. The level of hOLA1 in control siRNA treated cells were considered as standard (100%) expression and the percent relative expression of hOLA1 was calculated. The numbers were plotted as bar graph using Prism Graph Pad software. According to the densitometry, cells treated with 25 nM of si-hOLA1 demonstrated 68% decrease in the level of hOLA1 compared to the control siRNA treated cells. However, when the cells were treated with 50 nM si-hOLA1, the protein expression level of hOLA1 was further decreased (almost 90%) (Figure 3.1). Therefore, I decided to use 50 nM si-hOLA1 for the rest of the study.
Figure 3.1: Optimization of the concentration of si-hOLA1 for the depletion of hOLA1. The expression of hOLA1 was depleted using 25 nM and 50 nM siRNA in MDA-MB231 cells. GAPDH was used as loading control. The relative expression of hOLA1 in control and si-hOLA1 transfected cells were quantified using Amersham 600 imager. According to the quantification, 25 nM si-hOLA1 demonstrated 68% decreased expression of hOLA1 compared to control siRNA treated cells. However, when 50nM si-hOLA1 was used, cells showed effective depletion (90%) of hOLA1 compared to 25 nM si-hOLA1 treated cells. siContr: control siRNA-treated cells, si-hOLA1: siRNA targeting hOLA1-treated cells.

Subsequently, I optimized the transfection duration time with the same cell line and cell number as described above; here I used 50 nM siRNA and different transfection time points (24h, 48h, 72h, and 96h). Cells were lysed, and the proteins were analyzed by the Western blotting technique. The relative band intensity of hOLA1 and GAPDH was quantified and analyzed as described above. According to the densitometry, 24 hours si-hOLA1 treated cells demonstrated 40% decrease in the expression of hOLA1 compared to control siRNA treated cells. However, when the transfection duration time was increased to 48, 72 and 96 hours the protein level of hOLA1 was decreased by 60%, 80%, and 90% respectively (Figure 3.2). The better transfection duration for the effective knockdown was 96 hours (Figure 3.2). Therefore, I decided to use 50 nM siRNA and 96 hours transfection duration time for this study.
Figure 3.2: Optimization of si-hOLA1 transfection duration time for the effective depletion. hOLA1 was depleted using 50nM si-hOLA1 in MDA-MB231 cell line. GAPDH was used as loading control. The bands were quantified using Amersham 600 imager. According to the quantification, depending on the transfection duration time, hOLA1 expression was decreased by 40%, 60%, and 90% for 24, 48, and 96 hours transfection duration time respectively. The better hOLA1 depletion was observed at 96 hours post-transfection. siConstr: control siRNA-treated cells, si-hOLA1: siRNA targeting hOLA1-treated cells.

As mentioned earlier, hOLA1 depletion leads to diminished ISR level and enhanced cell survival. ISR is known to regulate apoptosis. According to my hypothesis, increased cellular survival during hOLA1 depletion is due to decreased apoptosis caused by diminished ISR level. As mentioned in the introduction, CHOP is an effector protein of ISR and induced during ISR activation. Therefore, to verify that hOLA1 depletion leads to diminished ISR level and thus diminished CHOP level, I performed knockdown studies and Western blot analysis to assess the level of CHOP upon hOLA1 depletion. GAPDH
was used as the loading control. The level of CHOP is decreased in hOLA1-depleted cells compared to control cells (Figure 3.3). This suggests that hOLA1 depletion leads to decreased ISR level resulting in decreased level of CHOP in hOLA1-depleted cells (Figure 3.3).

![Image: Western blot analysis showing successful hOLA1 depletion, decreased level of CHOP, and GAPDH as a loading control.](image)

**Figure 3.3: hOLA1 depletion leads to diminished ISR.** The experiment was performed in MDA-MB231 cells using 50 nM siRNA and 96 hours post-transfection time. The cropped blot on the top represents successful hOLA1 depletion, middle blot implies the level of CHOP is decreased upon hOLA1 depletion. The third blot represents GAPDH, loading control to assess if equal lysates are loaded in all lanes. Decreased levels of CHOP in hOLA1-depleted cells suggests the diminished ISR levels upon hOLA1 depletion. siContr: control siRNA-treated cells, si-hOLA1: siRNA targeting hOLA1-treated cells.

As mentioned earlier, ISR regulates apoptosis through the regulation of anti- and pro-apoptotic proteins. Therefore, I wanted to assess the effect of diminished ISR caused by hOLA1 depletion on the levels of key IAPs and anti-apoptotic proteins. To this end, hOLA1 was depleted using siRNA targeting hOLA1, and the levels of IAPs were assessed by Western blot analysis. The experiment was performed in two different cell lines: HEK-293T and MDA-MB231. I have used two different cell lines to verify if the role of hOLA1 in apoptosis is cell line specific or not. To this end, MDA-MB231 cells (1.5×10⁵) and HEK-293T cells (1.0×10⁵) were seeded and transfected with control siRNA and si-hOLA1. Cells were lysed 96 hours post-transfection and analyzed by Western blotting. The
membrane was probed for key anti-apoptotic proteins cIAP1, cIAP2, XIAP, cFLIP, and Bcl-xL.

The partial silencing of hOLA1 was verified by Western blotting (Figure 3.4). It is known that ISR regulates cIAP1.\(^8\) Hence, first I assessed the level of cIAP1 upon hOLA1 depletion. cIAP1 is an inhibitor of the extrinsic apoptotic pathway. It inhibits apoptosis by interacting with extrinsic pathway signal transduction proteins TRAF1 and TRAF2 thus inhibiting the activation of caspase-8.\(^1\) In addition, cIAP1 inhibits apoptosis by activation of RIP1 ubiquitylation.\(^1\) Activation of RIP1 leads to RIPoptosome or complex II formation results in the activation of NF-κB pro-survival pathway.\(^1\) The experiment was performed in three biological replicates. The bands were quantified using Amersham 600 software. The relative band intensity was calculated by dividing the protein of interest’s expression by the expression level of the loading control (GAPDH). According to the densitometry, hOLA1-depleted cells demonstrated an increased level of cIAP1 compared to control cells in both cell lines. Error bars on the bar graph represent the standard error of the means. The reason for the variation between the biological replicates may be due to the saturated protein bands. This may show the higher variability among biological replicates and thus showing huge error bars. This suggests that there is a clear trend in the increase of cIAP1 levels by hOLA1 depletion. However, the statistical variability can be improved by avoiding saturation of protein bands. However, the level of cIAP1 in hOLA1-depleted cells is upregulated in all three biological replicates suggesting that hOLA1 depletion likely lead to upregulation of cIAP1 in MDA-MB231 and HEK-293T cell lines (Figure 3.4).
Figure 3.4: hOLA1 depletion leads to increased levels of cIAP1 in MDA-MB231 and HEK-293T cell lines. The left panel shows Western blot results from the MDA-MB231 cell line, and the right panel shows Western blot results from the HEK-293T cell line. hOLA1 was depleted in both cell lines and the membrane was probed for cIAP1 and GAPDH. The protein bands were quantified using Amersham 600 imager, and the numbers represent the ratio of cIAP1 to GAPDH. The relative expression was assessed, according to the quantification, hOLA1 depletion leads to increased levels of cIAP1 in both MDA-MB231 and HEK-293T cells. These Western blots are from three biological replicates. Error bars show the standard errors of the means (SEM). siContr: control siRNA-treated cells, si-hOLA1: siRNA targeting hOLA1-treated cells.

cIAP2 is a key anti-apoptotic protein and a homolog of cIAP1.\textsuperscript{108} cIAP2 is known to have functional redundancy with cIAP1.\textsuperscript{151} Hence, I wanted to examine the levels of cIAP2 in hOLA1-depleted cells. hOLA1 was depleted in MDA-MB231 and HEK-293T cell lines. Subsequently, cell lysate was analyzed by Western blotting. The membrane was probed for cIAP2 and GAPDH. As described in the methods section, protein bands were quantified using an Amersham 600 Imager and the ratio of cIAP2 and GAPDH was assessed. The relative expression of cIAP2 was plotted using Prism Graph Pad. Error bars on the bar graph represent the standard error of the means. As mentioned earlier, some of the proteins
bands being saturated which may have resulted in a higher variability between biological replicates and thus showing huge error bars. However, according to the quantification, all three biological replicates of MDA-MB231 and HEK-293T cells suggest increased levels of cIAP2 upon hOLA1 depletion (Figure 3.5). The data indicate that hOLA1 depletion seems to upregulate cIAP2. However, this is not statistically significant. Taken together, hOLA1 depletion likely lead to the upregulation of cIAP1 and cIAP2.

**Figure 3.5:** hOLA1 depletion leads to increased levels of cIAP2 in MDA-MB231 and HEK-293T cell lines. The left panel shows Western blot results from the MDA-MB231 cell line, and the right panel shows Western blot results from the HEK-293T cell line. hOLA1 was depleted in both cell lines and the membrane was probed for cIAP2 and GAPDH. The protein bands were quantified using Amersham 600, the numbers represent the ratio of cIAP2 to GAPDH. According to the quantification, hOLA1 depletion leads to increased levels of cIAP2 in MDA-MB231 and HEK-293T cell lines. These Western blots are from three biological replicates. Error bars show SEM. siContr: control siRNA-treated cells, si-hOLA1: siRNA targeting hOLA1-treated cells.

Following that, I evaluated the levels of an anti-apoptotic protein that inhibits the extrinsic apoptotic pathway, cFLIP_L. cFLIP inhibits apoptosis by competing with procaspase-8 to be
recruited onto the DISC. Removal of hOLA1 did not show any considerable effect on the level of cFLIP splice variant cFLIP_L in MDA-MB231 and HEK-293T cell lines (Figure 3.6). The middle blot of cFLIP_L in HEK-293T was previously probed for cIAP2. The blot was not stripped and, like a cIAP2 antibody, the hOLA1 antibody was raised in rabbit, therefore it shows two bands (Figure 3.6, the middle blot in the right panel). The upper band was excluded during quantification. Error bars on the bar graph represent the standard error of the means. HEK-293T bar graph shows a huge error bar for the level of cFLIP_L in hOLA1-depleted cells. This result may be due to the saturated cFLIP_L band in the hOLA1-depleted cells (Figure 3.6, middle blot in the right panel). According to the quantification, hOLA1 depletion did not show any considerable effects on the level of cFLIP_L.
**Figure 3.6: hOLA1 depletion does not have a considerable effect on the levels of cFLIP<sub>L</sub> in MDA-MB231 and HEK-293T cell lines.** The left panel shows Western blot results from the MDA-MB231 cell line, and the right panel shows Western blot results from the HEK-293T cell line. hOLA1 was depleted in both cell lines and the membrane was probed for cFLIP<sub>L</sub> and GAPDH. The protein bands were quantified using Amersham 600 software, and the numbers represent the relative expression of cFLIP<sub>L</sub> and GAPDH. According to the quantification, hOLA1 depletion does not have a considerable effect on the level of cFLIP<sub>L</sub> in both cell lines. These Western blots are from three biological replicates. Error bars show SEM. siContr: control siRNA-treated cells, si-hOLA1: siRNA targeting hOLA1-treated cells.

It is known that removal of hOLA1 leads to diminished CHOP.<sup>136</sup> CHOP triggers apoptosis by repressing Bcl-2 family anti-apoptotic proteins.<sup>153</sup> Therefore, I evaluated the levels of the Bcl-2 family anti-apoptotic protein: Bcl-xL. Bcl-xL is crucial anti-apoptotic protein, that prevents pore formation on the mitochondrial membrane thus inhibit intrinsic and extrinsic apoptotic pathways.<sup>96</sup> The level of Bcl-xL was assessed by Western blotting. HEK-293T always produced double bands for Bcl-xL. When the control siRNA treated cells are compared to si-hOLA1 treated cells, the level of Bcl-xL seems to be upregulated.
upon hOLA1 depletion in both cell lines (Figure 3.7). As mentioned earlier, some of the bands are saturated which may have resulted in a higher variability between biological replicates likely showing huge error bars.

Figure 3.7: hOLA1 depletion leads to increased levels of Bcl-xL in MDA-MB231 and HEK-293T cell lines. The left panel shows Western blot results from the MDA-MB231 cell line, and the right panel shows Western blot results from the HEK-293T cell line. hOLA1 was depleted in both cell lines and the membrane was probed for Bcl-xL and GAPDH. The protein bands were quantified using Amersham 600 software, and the numbers represent the relative expression of Bcl-xL and GAPDH. hOLA1 depletion leads to upregulation of Bcl-xL in HEK-293T and MDA-MB231 cell lines. These Western blots are from three biological replicates. Error bars show the SEM. siContr: control siRNA-treated cells, si-hOLA1: siRNA targeting hOLA1-treated cells.

Previously it was reported that ATF4 regulates XIAP during ISR. XIAP is known to inhibit both intrinsic and extrinsic apoptotic pathways by inhibiting caspase-9, caspase-3, and caspase-7. Therefore, I wanted to assess the level of XIAP upon hOLA1 depletion. To this end, I performed hOLA1 depletion and Western blot analysis, and the relative expression was assessed as mentioned in the previous section. According to the quantification, there are no detectable changes in the levels of XIAP between control cells.
and hOLA1-depleted cells in both cell lines (Figure 3.8). The reason for the variation between the biological replicates may due to some of the protein bands being saturated which may have resulted in more variation between biological replicates likely showing huge error bars. Taken together, the findings of this section suggest that hOLA1 depletion likely leads to upregulation of key anti-apoptotic proteins: cIAP1, cIAP2, and Bcl-xL, but not XIAP or cFLIPL.

**Figure 3.8:** hOLA1 depletion does not have a considerable effect on the levels of XIAP. The left panel shows Western blot results from the MDA-MB231 cell line, and the right panel shows Western blot results from the HEK-293T cell line. hOLA1 was depleted in both cell lines and the membrane was probed for XIAP and GAPDH. The protein bands were quantified using Amersham 600 software, the numbers represent the relative expression of XIAP and GAPDH. According to the quantification, hOLA1 depletion doesn’t have considerable changes on the level of XIAP in MDA-MB231 and HEK-293T cells. Error bars show SEM. siContr: control siRNA-treated cells, si-hOLA1: siRNA targeting against hOLA1-treated cells.
3.2 hOLA1 depletion leads to decreased caspase activation in the MDA-MB231 cell line.

The previous section of this chapter shows that hOLA1 depletion might lead to upregulation of key anti-apoptotic proteins: cIAP1, cIAP2, and Bcl-xL. The major role of an anti-apoptotic protein is to inhibit caspase activation through direct and or indirect interactions. Therefore, I wanted to examine the cellular levels of initiator and executioner caspases in the combination of si-hOLA1 and apoptotic inducers treatment. In order to induce caspase activation, MDA-MB231 cells were treated with various apoptotic agents such as doxorubicin, TRAIL, and TNFα. Doxorubicin induces apoptosis by damaging DNA through intercalation and torsional stress while TRAIL and TNF-α induce apoptosis by binding to the receptors on the cell membrane.\textsuperscript{154, 155} MDA-MB231 (1.5×10\textsuperscript{5}) cells were seeded in a 6-well plate, and hOLA1 was depleted using siRNA targeted against hOLA1. After 92 hours of siRNA treatment, cells were treated with 10 nM of doxorubicin, 100 ng/mL of TRAIL and 100 ng/mL of TNFα for 4 hours. Consequently, cells were lysed, and the proteins were analyzed by Western blotting. The membrane was probed for caspase-8, caspase-9, caspase-7, and caspase-3. In addition, the membrane was probed for cleaved PARP. PARP is a polymerase involved in DNA repair.\textsuperscript{156} This protein is cleaved by caspase enzymes, therefore I wanted to assess the protein level of cleaved PARP by Western blotting.

The previous section suggests that hOLA1 depletion leads to upregulation of cIAP1, cIAP2, and Bcl-xL. As mentioned in the introduction, extrinsic anti-apoptotic proteins: cIAP1, and cIAP2 inhibit initiator caspase of the extrinsic pathway: caspase-8. Moreover, intrinsic apoptotic pathway activation could result in the activation of the extrinsic pathway. Intrinsic anti-apoptotic protein upregulation could also result in the inhibition of extrinsic
initiator caspase: caspase-8. Therefore, I assessed the level of caspase-8 upon hOLA1 depletion and apoptotic inducers treatment. When the membrane was probed for caspase-8, the cleaved caspase-8 level is higher in the combination of TRAIL and control siRNA compared to TRAIL combined with si-hOLA1. TRAIL is an extrinsic apoptotic pathway inducer, decreased caspase-8 cleavage may possibly due to the upregulation of extrinsic anti-apoptotic proteins such as cIAP1 and cIAP2 in hOLA1-depleted cells (Figure 3.9). However, executioner caspases: caspase-7 or caspase-3 did not show any detectable levels of activation. Nevertheless, the level of full-length caspase-7 seems to be increased in the cells treated with apoptotic agents and siRNA targeted against hOLA1 (Figure 3.9). This may possibly due to transcriptional or translation upregulation of caspase-7 upon hOLA1 depletion.\textsuperscript{157} This may also possibly due to stabilization of caspase-7.\textsuperscript{158} However, TNF\textgreek{a} treated cells did not show detectable differences in the levels of cleaved caspase-8 (Figure 3.9). This may possibly due to the duration of TNF\textgreek{a} treatment may not enough to induce caspase activation.

The intrinsic apoptotic pathway is mainly initiated through intrinsic pro-apoptotic proteins that leads to the activation of intrinsic initiator caspases and executioner caspases.\textsuperscript{89} Previous section of this study suggests that hOLA1 depletion leads to increased levels of Bcl-xL. As mentioned earlier, Bcl-xL indirectly inhibits caspase-9 activation.\textsuperscript{108} Therefore, I wanted to assess the protein level of caspase-9 upon hOLA1 depletion and apoptotic inducers treatment as described above. When the membrane was probed for caspase-9, an initiator caspase of the intrinsic apoptotic pathway, very faint cleaved caspase-9 was detected in the control lane treated with doxorubicin, whereas no detectable band was observed in si-hOLA1 combined with doxorubicin (Figure 3.9). This may possibly due to
the effect of increased Bcl-xL levels upon hOLA1 depletion. Increased levels of Bcl-xL may possibly have inhibited the activation of caspase-9 upon hOLA1 depletion. Interestingly, increased level of cleaved caspase-9 was observed in control cells treated with TRAIL compared to the combination of si-hOLA1 with TRAIL. As mentioned earlier in the introduction, TRAIL induces the activation of caspase-8. Activation of caspase-8 may result in the cleavage of Bid into t-Bid thus activating the intrinsic apoptotic pathway. Therefore, the increased level of cleaved caspase-9 is perhaps due to the crosstalk between intrinsic and extrinsic apoptotic pathways (Figure 3.9).

hOLA1 depletion resulted in the upregulation of anti-apoptotic proteins and the activation of caspase-8 and caspase-9. Activation of caspases leads to the cleavage of PARP. PARP helps in the DNA repairing process during stress. Therefore, I wanted to assess the level of cleaved PARP in the combination of si-hOLA1 and apoptotic inducers. When the membrane was probed for cleaved PARP, the level of PARP activation was inhibited in the combination of sihOLA1-doxorubicin-treated cells compared to combination of siContr-doxorubicin-treated cells. Moreover, si-hOLA1-TRAIL-treated cells showed a clear inhibition of PARP activation compared to control si-RNA-TRAIL-treated cells. However, TNFα treatment does not show considerable PARP cleavage. This is perhaps due to the duration time of the TNFα treatment may not enough to induce apoptosis in the cells. Notably, TRAIL treatment shows more PARP cleavage compared to DNA damage inducer doxorubicin, this may possibly due to the effect of doxorubicin on PARP1 enzyme activity and expression. Doxorubicin is known to induce the suppression of PARP1 enzyme activity and the expression, therefore we may see decreased level of cleaved PARP in doxorubicin treated cells. Taken together, the data suggest that, hOLA1 depletion leads
to upregulation of anti-apoptotic proteins that may possibly result in the inhibition caspase activation.

**Figure 3.9: hOLA1 depletion leads to decreased caspase activation.** hOLA1 was depleted in MDA-MB231 cells using siRNA transfection. DMSO was used as a control, and the cells were treated with 10 nM, 100 ng/mL, 100 ng/mL doxorubicin, TRAIL and TNFα respectively. Upon hOLA1 depletion and TRAIL treatment, cells showed decreased caspase-8 activity. Also, the level of cleaved PARP is decreased in hOLA1-depleted cells in the presence of apoptotic inducers, especially with TRAIL. The bar graph on the right represents the relative expressions of hOLA1 in all the treatment.

3.3. **hOLA1 depletion does not have considerable effects on the inhibition of cell death**

Previous experiments of this study showed that hOLA1 depletion leads to upregulation of key anti-apoptotic proteins such as cIAP1, cIAP2, and Bcl-xL. Also, hOLA1 depletion demonstrated a decreased caspase-8 activation upon apoptotic agent treatment. In addition, hOLA1 depletion and apoptotic agent treated cells demonstrated a decreased PARP
cleavage compared to control siRNA and apoptotic agent treated cells suggesting that hOLA1 depletion may inhibit apoptosis. Hence, I wanted to further examine the viability of hOLA1-depleted cells in the presence and absence of apoptotic inducers. The alamarBlue™ reagent was used to assess cell viability. MDA-MB231 cells (1×10^5) and HEK-293T cells (6×10^3) were seeded in a 96-well plate. Cells were treated with siRNA targeted against hOLA1 (si-hOLA1) and control siRNA (si-Cont). After 24 hours of siRNA treatment, cells were treated with apoptotic inducers (doxorubicin 10 nM, TRAIL 100 ng/mL, and TNFα 100 ng/mL) for 72 hours. DMSO was used as a control because doxorubicin is dissolved in DMSO. Subsequently, the alamarBlue™ solution was added, and the fluorescence reading was measured using a fluorescent plate reader to measure cell viability.

In each biological replicate, all the treatments were normalized to control (DMSO). The average of three biological replicates was obtained. The differences between means were determined by the student t-test at P ≤ 0.05 using Graph Pad Prism software. According to the data from the MDA-MB231 cell line, there are no detectable differences in the viability between hOLA1-depleted cells and control cells in the absence of apoptotic agents (Figure 3.10). However, when the cells were treated with apoptotic agents such as TRAIL and TNFα, and si-hOLA1 shows a modest and statistically non-significant inhibition of cell death compared to control cells treated with the apoptotic agent. This modest difference may possibly be due to increased key anti-apoptotic proteins such as cIAP1 and cIAP2. When the cells were treated with intrinsic pathway apoptotic inducer doxorubicin, hOLA1-depleted cells exhibited a modest and statistically non-significant decrease in cell death.
This modestly decreased viability may possibly due to the upregulation of intrinsic anti-apoptotic protein Bcl-xL upon hOLA1 depletion.

Figure 3.10: hOLA1 depletion and apoptotic induction do not have considerable effects on the inhibition of cell death in MDA-MB231 cell line. hOLA1 was depleted in MDA-MB231 cells using siRNA transfection. The light grey bar shows control siRNA treated cell viability and the dark grey bar shows hOLA1-depleted cell viability. DMSO was used as a control. The cells were treated with 10 nM doxorubicin, 100 ng/mL TRAIL, and 100 ng/mL TNF-α. The graph represents means of three biological replicates. Upon hOLA1 depletion and apoptotic inducers treatment, cells show modest inhibition of cell death. An unpaired t-test was used to detect significant differences at P ≤ 0.05. The difference was found to be non-significant. Therefore, hOLA1 depletion and apoptotic induction do not have a considerable effect on the inhibition of cell death. The error bars represent the SEM.

The same experiment was performed in HEK-293T. HEK-293T cells showed similar trend like MDA-MB231. However, cells did not show a modest cell death inhibition upon hOLA1 depletion and apoptotic agent treatment like MDA-MB231. Notably, the alamarBlue™ assay may not be represent the correct physiological state of the cell. For example, cells undergoing apoptosis may not represented as dead in the alamarBlue™ assay. Therefore, the alamarBlue™ assay does not validate if the modest cell death
inhibition is due to decreased apoptosis. Hence, to validate that the increased cell survival upon hOLA1 depletion is due to the upregulation of anti-apoptotic proteins resulted in decreased apoptosis, I performed PI staining and flow cytometry.

Figure 3.11: hOLA1 depletion and apoptotic induction do not have considerable effects on cell death inhibition in HEK-293T cell line. hOLA1 was depleted in HEK-293T cells using siRNA transfection. The light grey bar represents the viability of control siRNA treated cells, and the dark grey bar represents the viability of hOLA1-depleted cells. DMSO was used as a control, and the cells were treated with 10 nM doxorubicin, 100 ng/mL TRAIL, and 100 ng/mL TNF-α. The graph represents means of three biological replicates. Cells do not demonstrate any considerable effects on the inhibition of cell death upon hOLA1 depletion and apoptotic agent treatment. An unpaired t-test was used to detect significant differences at P ≤ 0.05. Error bars represent the SEM.

3.4. Measurement of apoptosis in hOLA1-depleted cells

The PI staining experiment was performed using MDA-MB231 cells. MDA-MB231 (1.5×10^5) cells were seeded in a 6-well plate, and hOLA1 depletion was performed as described previously. At 92 hours of post-transfection, cells were treated with 100 ng/mL apoptotic inducer TRAIL. After 4 hours of treatment with TRAIL, cells were trypsinized and fixed in 70% ethanol overnight. Following that, cells were stained with PI staining and submitted for flow cytometry.
Apoptotic cells are characterized by DNA fragmentation and the loss of DNA content. PI is a fluorescent intercalating agent that can be used to label DNA and measure the cellular DNA content by flow cytometry analysis.\textsuperscript{161} Hence, PI staining and flow cytometry analysis was used to measure apoptosis upon hOLA1 depletion. Apoptotic cells are represented in the sub $G_0$ phase because cells containing fragmented DNA would be in the $G_0$ phase. Due to some technical errors made in gating, I am presenting the data from two different biological replicates (control cells without apoptotic inducers and cells treated with apoptotic inducers). Hence, I did not compare control cells (no treatment) and apoptotic agent-treated cells. Based on the control cells treated with nocodazole, each phase was gated by the flow cytometry facility. From the histogram, the percentage of cells sub-population in each phase was obtained (Figure 3.12, numbers in each phase of the histogram). The control siRNA treated cells were considered as 100% and si-hOLA1 treated cells were normalized to control cells in each phase. The percentage of cells in each phase was plotted using Prism Graph Pad. The percentage of cells sub-population in each phase was compared between control siRNA and si-hOLA1 treated cells. From the bar graph on the left (representative of histogram) the proportion of cell number in $G_0$ phase was decreased in hOLA1-depleted cells. A 20% decrease in apoptosis (number of cells in the sub $G_0$) was observed (Figure 3.12). Also, when cells treated with the combination of control siRNA and TRAIL were compared to cells treated with a combination of TRAIL and si-hOLA1, a 40% decrease in apoptosis was observed (Figure 3.12, the bar graph on the right). This data suggests that hOLA1 depletion leads to decreased apoptosis. Moreover, control siRNA treated cells (Figure 3.12, left-bar graph) demonstrated an increased proportion of the cells in $G_0/G_1$ phase, suggesting that hOLA1 depletion leads to cell cycle arrest in $G_0/G_1$ phase (Figure 3.12). This is possibly due to hOLA1’s role as a suppressor
of p21. Removal of hOLA1 leads to increased level of p21 thus causing a cell cycle arrest in the G0/G1 phase. However, a combination of si-hOLA1 and apoptotic agent treatment did not demonstrate a cell cycle arrest in G0/G1 phase. This may be possibly due to the effect of TRAIL treatment. hOLA1 depletion demonstrated a decreased proportion of the cell number S and G2/M phase. The same trend was observed for S and G2/M phase in hOLA1-depleted-TRAIL-treated cells. Together the findings of my study suggest that hOLA1 depletion leads to upregulation of anti-apoptotic proteins such as cIAP1, cIAP2, and Bcl-xL thus possibly decreases caspase activation and apoptosis.

Figure 3.12: hOLA1 depletion leads to decreased apoptosis. hOLA1 was depleted in MDA-MB231 cell line using si-hOLA1. From the data file, DNA content from each phase in the histogram was plotted and represented as a bar graph. DNA content represents the proportion of the number of cells in each phase. According to the bar-graph hOLA1-depleted cells showed a 20% decrease in the number of apoptotic cells compared to control cells. The bar graph on the right side shows cells treated with 100 ng/mL TRAIL. Upon hOLA1 depletion and TRAIL treatment, cells showed 40% decrease in the number of apoptotic cells (sub-G0 cells).
3.5 Co-immunoprecipitation and mass spectrometry to identify the protein(s) interacting with hOLA1

As mentioned earlier, hOLA1 interacts with eIF2 and inhibits global translation. However, it remains unclear as to how hOLA1 interacts with eIF2. The interaction of hOLA1 with eIF2 could be regulated by additional protein(s). To identify the protein(s) interacting with hOLA1, I performed co-immunoprecipitation of hOLA1 using MDA-MB231 cells followed by mass spectrometry. Cells were lysed when they reached 80% confluency using the RIPA lysis buffer. Subsequently, the lysate was incubated with anti-hOLA1 antibody-coated beads for 3 hours at 4˚C. A portion of IP eluate was subjected to SDS-PAGE separation and stained with Coomassie stain in preparation for the mass spectrometry analysis. Protein bands were then excised and submitted for mass spectrometry.

The pull-down of hOLA1 was validated by Western blotting using the remaining portion of IP eluate, then the membrane was probed for Anti-hOLA1 antibody (Figure 3.13, left panel). The same membrane was probed for eIF2α. However, as mentioned by Chen, H et al., I did not see the eIF2α interaction of hOLA1 with endogenous IP using anti-hOLA1 antibody. On the Coomassie-stained gel, I observed two bands, one at about 45kDa (Sample I), and the other at about 150kDa (Sample II). I anticipated the protein observed at about 45kDa could be hOLA1 (MW:42kDa). Hence I excised both bands and submitted them for mass spectrometry. The mass spectrometry resulted in many hits for both samples. I looked at the molecular weight of excised band corresponding to the given hits and higher coverage (coverage is the percentage of the protein sequence covered by all identified peptides in the sample). Next, I checked sequence similarity between the proteins similar to hOLA1’s
molecular weight with hOLA1’s sequence using basic local alignment search tool (BLAST) in National Center for Biotechnology Information (NCBI). As a result, the protein similar to hOLA1’s molecular weight had 90% coverage from mass spectrometry and 67% similarity with hOLA1. This strongly suggests that hOLA1 is pulled down. Next, I assessed the hits received for sample II, according to the molecular weight and the better coverage. Three hits demonstrate a similar molecular weight as sample II and better coverages such as Ubiquitin C, Rho-associated protein kinase, and Actin like filament (FLJ00119) (Table 3.2). The Ubiquitin C variant protein demonstrates higher coverage (40%) compared to the other two. It is known that hOLA1 protects HSP70 from ubiquitylation-mediated protein degradation.\textsuperscript{137} Hence, it is possible that hOLA1 may interact with Ubiquitin C variant and prevent ubiquitylation-mediated degradation. However, the interaction of hOLA1 with Ubiquitin C variant needs to be further validated by reciprocal IP and Western blotting. The Rho-associated protein kinase and Actin like filament (FLJ00119) demonstrate similar molecular weight and 2% or 5% coverage respectively. Rho-associated protein kinase and Actin like filament (FLJ00119) are known to be involved in cell adhesion.\textsuperscript{162} It is previously reported that hOLA1 negatively regulates cell adhesion.\textsuperscript{137} Taken together, the data of IP and mass spectrometry suggests that hOLA1 may not strongly interact with the α subunit of eIF2 as previously reported by Chen, H \textit{et al.}\textsuperscript{136} Also, hOLA1 may possibly interact with Ubiquitin C variant or Rho-associated protein kinase or Actin like filament and negatively regulate cell adhesion.
Figure 3.13: Co-immunoprecipitation to identify the protein(s) interacting with hOLA1. The left image shows the western blot analysis of hOLA1 immunoprecipitation. The first lane represents the whole cell lysate, second lane represents the control-IP and the third lane represents the hOLA1-IP. The protein band at 42kDa in third lane validates the pulldown of hOLA1. The right image represents the SDS-PAGE and Coomassie staining. The observed bands on the Coomassie-gel (sample I and II) were excised and analyzed by mass-spectrometry.

Table 3.1: Mass spectrometry data analysis of sample I (MW ~45kDa). Hits with higher coverage and similar molecular weight of hOLA1 were assessed for sequence similarities with hOLA1 to see if the pull-down is worked. There is a 64%-67% sequence similarity between hOLA1 and mass spectrometry hits with higher coverage and similar molecular weight validating the pulldown of hOLA1 has worked.

<table>
<thead>
<tr>
<th>Hits (coverage)</th>
<th>MW(kDa)</th>
<th>Sequence similarity with hOLA1</th>
</tr>
</thead>
<tbody>
<tr>
<td>Putative uncharacterized protein (91%)</td>
<td>40.4</td>
<td>67% Identity</td>
</tr>
<tr>
<td>Guanine nucleotide-binding protein G (87%)</td>
<td>40.7</td>
<td>64% Identity</td>
</tr>
</tbody>
</table>
Table 3.2: Possible interacting partner of hOLA1. Mass spectrometry data for sample II. Hits with higher coverage and similar MW of sample II from the mass spectrometry data is listed on the table. These proteins are previously reported to be involved in cell adhesion. hOLA1 negatively regulates cell adhesion. This suggests that hOLA1 was successfully pulled-down and it could interact with one of the proteins listed above.

<table>
<thead>
<tr>
<th>Hits(coverage)</th>
<th>MW(kDa)</th>
<th>Known roles of hOLA1 of the hits</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ubiquitin C variant (40%)</td>
<td>147.25</td>
<td>hOLA1 is known to prevent HSP70 from ubiquitylation-mediated degradation</td>
</tr>
<tr>
<td>Rho-associated protein kinase (2%)</td>
<td>158</td>
<td>hOLA1 is a negative regulator of cell adhesion</td>
</tr>
<tr>
<td>Actin like filament (FLJ00119) (5%)</td>
<td>158</td>
<td>hOLA1 is a negative regulator of cell adhesion.</td>
</tr>
</tbody>
</table>
Chapter 4: Discussion and future directions

4.1 Discussion

hOLAl is known to have numerous roles in tumor progression like cancer metastasis, and inhibition of centrosome regulation.\textsuperscript{135, 137, 138, 143} Moreover, hOLAl is reported to exhibit a regulatory effect on the global mRNA translation.\textsuperscript{136} hOLAl suppresses translation initiation by inhibiting ternary complex formation, and removal of hOLAl leads to enhanced cellular survival and diminished ISR.\textsuperscript{136} Decreased level of ISR leads to diminished ISR effector proteins: ATF4 and CHOP.\textsuperscript{163} Depending on the type of the stress condition, ATF4 and CHOP regulate pro-apoptotic genes at the transcription level (Bcl-2) and anti-apoptotic proteins.\textsuperscript{85, 164} Therefore, I hypothesized that increased cell survival upon hOLAl depletion is due to decreased apoptosis caused by the effect of diminished ISR on anti-apoptotic proteins. In this study, I evaluated the role of hOLAl in apoptosis by assessing the levels of anti-apoptotic proteins, caspase activation and measuring apoptosis. Diminished ISR in hOLAl-depleted cells was verified by assessing the levels of CHOP (Figure 3.3). Removal of hOLAl resulted in decreased CHOP levels (Fig. 3.3) and upregulation of key anti-apoptotic proteins; cIAP1, cIAP2, and Bcl-xL (Figures: 3.4, 3.5, 3.7 respectively). Previous studies have reported that ISR plays a crucial role in regulating cellular survival and cell death pathways depending on the type of stress condition.\textsuperscript{38} CHOP and ATF4 is known to have numerous roles in promoting and inhibiting apoptosis by regulating anti- and pro-apoptotic proteins as described below.\textsuperscript{165}

During chronic PERK signaling, ATF4-CHOP signaling is reported to show decreased levels of the IAPs: cIAP1, cIAP2, Livin, and Survivin.\textsuperscript{78} Furthermore, PERK signaling leads to decreased synthesis of XIAP through dysregulation of cap-dependent translation.\textsuperscript{78}
Also, increased levels of ATF4 during PERK mediated ISR promotes proteasomal degradation of XIAP through ubiquitylation resulting in decreased XIAP level.\(^7\) Moreover, CHOP is a direct target of Bcl-2 family proteins.\(^{166}\) CHOP-mediated apoptosis is known to be inhibited by anti-apoptotic protein: Bcl-xL and CHOP are known to promote the expression of pro-apoptotic proteins: DR5 and Bax.\(^{166,167}\) Moreover, CHOP is known to be degraded by cIAP1 and CHOP degrades cFLIP\(_L\) through ubiquitylation and proteasome degradation.\(^{144,145}\) Together, all these previous studies of ATF4-CHOP mediated regulation of anti- and pro-apoptotic proteins suggesting that decreased CHOP levels could be the reason for the upregulation of anti-apoptotic proteins upon hOLA1 depletion. Hence, I propose that upregulation of anti-apoptotic proteins: cIAP1, cIAP2, and Bcl-xL may possibly occur due to the diminished levels of CHOP and ATF4 caused by hOLA1 depletion.

On the other hand, hOLA1 is known to regulate the expression of heat shock protein 70 (HSP70).\(^{137}\) HSP70 is stress response protein that has the cellular defense mechanism against environmental stresses.\(^{137}\) HSP70 is known to inhibit apoptosis by interacting with APAF1.\(^{168}\) hOLA1 prevents ubiquitylation mediated degradation of HSP70 by replacing the E3 ubiquitin ligase: C-terminus of Hsc70-interacting protein (CHIP) at the carboxyl-terminus of HSP70.\(^{137}\) Therefore, hOLA1 may have a direct or indirect interaction with anti-apoptotic proteins leading to the inhibition of its function. Hence, hOLA1 may negatively regulate anti-apoptotic proteins independent of CHOP and ATF4. It is also possible that the upregulation of cIAP1 and cIAP2 upon hOLA1 depletion leads to the activation of the NF-κB pro-survival pathway, resulting in the enhanced cellular survival.\(^{169}\)
Notably, when I performed Western blot analysis in order to assess the levels of anti-apoptotic proteins upon hOLA1 depletion, some of the bands were too saturated. This may have been caused by issues related to Western blotting such as immunoprobing, membrane transfer, concentrated antibody or increased exposure time of the blot in the imaging system. Different type of optimization step may reduce the saturation of bands such as linearity test of the antibody with different concentration of the proteins and varying exposure time of the blots while imaging.\(^{170}\)

The major function of an anti-apoptotic protein is to inhibit apoptosis by directly or indirectly inhibiting caspase activation.\(^{108}\) Inhibitors of extrinsic apoptotic pathway cIAP1 and cIAP2 inhibit caspase-8 and caspase-10.\(^{108}\) TRAIL induces extrinsic apoptotic pathway by binding to the receptor on the cell membrane.\(^{155}\) In this study, activation of caspases was assessed by evaluating the cellular levels of cleaved caspases. Figure 3.9 demonstrates increased level of caspase-8 activation in control siRNA-TRAIL-treated cells compared to si-hOLA1-TRAIL-treated cells. This suggests that increased levels of extrinsic inhibitors of apoptosis proteins cIAP1 and cIAP2 may have resulted in decreased caspase-8 activation upon hOLA1 depletion and TRAIL treatment. Notably, cleaved caspase-9 was reduced in hOLA1-depleted-TRAIL-treated cells compared to control siRNA-TRAIL-treated cells, suggesting that this effect is possibly due to the crosstalk between the extrinsic and intrinsic apoptotic pathways. Caspase-8 may cleave Bid into truncated Bid (t-Bid) thus activating the intrinsic pathway of apoptosis.\(^{105}\) However, this needs to be verified by assessing the cellular levels of Bid and tBid by Western blotting. Notably, PARP is a DNA repairing enzyme, however, when the cells were treated with DNA damaging agent doxorubicin, the western blot image did not demonstrate considerable PARP cleavage, this could possibly
due to doxorubicin is known to induce the suppression of PARP1 enzyme activity and the expression thus showing decreased PARP level (Figure 3.9).  

Furthermore, viability assay to assess the effects of hOLA1 depletion revealed that depletion of hOLA1 demonstrates modest inhibition of cell death in the presence of apoptotic inducers. However, the western image did not show considerable inhibition of cell death. This could be due to variable reason: cells were treated with apoptotic agents for 72 hours. Even though hOLA1 depletion leads to the upregulation of anti-apoptotic proteins when cells were treated with apoptotic inducers like TRAIL they undergo lots of other stresses like oxidative stress and plasma membrane depolarization. Therefore, the upregulation of anti-apoptotic proteins may not enough to overcome the effect of the apoptotic inducers treatment. Moreover, TNFα induces cell survival pathway through cFLIP\textsubscript{L} activation. This is perhaps a reason for the failure of hOLA1-depleted cells to demonstrate considerable difference in the cell viability compared to control cells and caspase-8 activation (Figure 3.9, 3.10 and 3.11). Also, doxorubicin treated cells did not show considerable cell death compared to cells treated with DMSO (control cells) in viability assay. It may possibly due to the concentration of doxorubicin may not enough to induce cell death. As mentioned earlier, the alamarBlue™ assay may not represent cells undergoing apoptosis. Therefore, I performed PI staining and flow cytometry to show that hOLA1 depletion leads to inhibition of cell death. The results of PI staining and flow cytometry analysis demonstrate that hOLA1 depletion leads to inhibition of apoptosis.

The mechanism of how hOLA1 inhibiting TC and the interaction with eIF2 is still unclear. In order to identify the protein(s) interacting with hOLA1, I attempted to perform IP and mass spectrometry. To verify the pull-down and interaction of hOLA1 with
eIF2α as mentioned by Chen, H et al., I performed Western blotting with a portion of the IP sample. When I probed the membrane with anti-hOLA1 and anti-eIF2α antibody, I did not detect eIF2α as mentioned by Chen, H et al.\textsuperscript{136} Notably, Chen, H et al. did not validate the interaction of hOLA1 with eIF2α by endogenous IP of hOLA1.\textsuperscript{136} The reason for not detecting the interaction of hOLA1 with eIF2α by Western blotting may possibly due to weak interaction of eIF2α with hOLA1, or hOLA1 may not interacting with α subunit of eIF2, it could possibly interact with GTP binding subunit of eIF2: γ subunit. In future, the interaction of eIF2 and hOLA1 can be assessed by crosslinking IP and Western blotting. Mass spectrometry analysis resulted in three possible proteins for the interacting partner of hOLA1: Ubiquitin C, Rho-associated protein kinase and Actin like filament (FLJ00119) (Table 3.1). hOLA1 is known to regulate cell adhesion and prevent HSP70 ubiquitylation-mediated protein degradation.\textsuperscript{150} Rho-associated protein kinase and Actin like filament (FLJ00119) are known to involved in cell adhesion.\textsuperscript{162} Ubiquitin C is involved in ubiquitylation-mediated protein degradation.\textsuperscript{172} Therefore, it is possible that one of these proteins interacts with hOLA1 and regulate cell adhesion or protein degradation.\textsuperscript{162} Taken together, this study suggests that hOLA1 depletion leads to upregulation of anti-apoptotic proteins such as cIAP1, cIAP2, Bcl-xL and decreased caspase activation resulting in decreased apoptosis.
4.2 Future perspective

hOLA1’s roles as a protein synthesis suppressor and ISR regulator provide a different aspect of hOLA1’s role in cell survival. My study suggests that hOLA1 depletion inhibits apoptosis through upregulation of anti-apoptotic proteins such as; cIAP1, cIAP2, and Bcl-xL. However, further validation of the role of hOLA1 in anti-apoptotic protein regulation by overexpression of hOLA1. In order to effectively conclude that hOLA1 depletion inhibits apoptosis, it needs to be assessed as to if hOLA1 negatively regulate anti-apoptotic proteins independent of CHOP. Further, is this effect exerted due to the diminished CHOP level caused by hOLA1 depletion? Additional studies are required to validate the effect of hOLA1 removal on anti-apoptotic proteins in two conditions: CHOP overexpression and CHOP downregulation. Moreover, the effect of hOLA1 depletion in transcription and translation of anti-apoptotic proteins needs to be studied further. This can be studied by measuring the steady-state levels of mRNAs encoding anti-apoptotic proteins by qPCR. Polysome profiling study is required to validate if hOLA1 is translationally regulating anti-apoptotic proteins.

It would be interesting to assess the combination of hOLA1 depletion and Thapsigargin treatment (ISR inducer) and analyze at the levels of anti-apoptotic proteins. Following the treatment of the cells with this compound, assessing levels of CHOP and of anti-apoptotic proteins by Western blotting or qPCR will reveal if the effect is CHOP dependent or not. An important question arising from this study is how cIAP1 and cIAP2 are upregulated at the same time, it is known that cIAP1 and cIAP2 have functional redundancy. The survival pathway NF-κB is regulated by the expression of both cIAP1 and 2. Therefore, I
suggest that hOLA1 may have some role in modulating NF-κB pro-survival pathway. This can be assessed by the mRNA and protein levels of survival pathway protein(s).

Furthermore, co-IP and mass spectrometry to assess the additional protein(s) interacting with hOLA1 was inconclusive as it was done only once. If we want to confirm and study further the interaction between the proteins pulled down with hOLA1, we can perform co-IP with overexpression of hOLA1. Overexpression of hOLA1 can be achieved by introducing a vector containing hOLA1 (for example FLAG-tag protein) by transfection. Following overexpression of hOLA1, IP, Western blotting and mass spectrometry may reveal the interaction of the unknown protein. Furthermore, pull-down can be verified by Western blotting with the antibody against the pulled down protein (mass spectrometry hits) and hOLA1. Moreover, the interaction between hOLA1 and eIF2α can be validated by in vitro assays: for example, this interaction can be investigated by purifying each subunit of eIF2. These subunits can be incubated with hOLA1, and subsequent protein-pulldown will reveal the interacting partner of eIF2 with hOLA1.173

In future, this study can be further advanced using xenograft and syngeneic mouse models.174 It is important to study the effect of overexpression of hOLA1 and tumor progression as we propose that hOLA1 promotes apoptosis. Hence overexpression of hOLA1 may result in decreased cancer progression. It would be interesting to see the synergistic effect of hOLA1 overexpression and anti-cancer drug treatment in tumorigenesis. Moreover, expression levels of hOLA1 in different types of cancer and the reason for aberrant expression need to be elucidated as since it can serve as a tumor marker. In conclusion, hOLA1 may represent a potential therapeutic target for cancer.
References


40. Brostrom, C.O., Prostko, C.R., Kaufman, R.J. & Brostrom, M.A. Inhibition of translational initiation by activators of the glucose-regulated stress protein and heat


