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Probing the Role of Nascent Helicity in Protein Function: p27^{kip1} as a Regulator of the Cell Cycle

Abstract

p27^{kip1} binds to and regulates the activity of cyclin-dependent kinases (Cdks) which are the master timekeepers of the cell division cycle. Members of the p27 family of proteins, also including p21 and p57, are called cyclin-dependent kinase inhibitors (CKIs). The amino terminal domain of p27 inhibits Cdk activity and is referred to as the kinase inhibitory domain (KID). The KID is comprised of a cyclin-binding domain (D1) and a Cdk binding domain (D2) joined by a 22 residue linker domain (LH). Structural analysis of the KID in solution before binding its Cdk targets revealed that D1 and D2 are largely unstructured and that the linker domain exhibits nascent helical characteristics. The nascent helical structure of the linker domain is conserved amongst the CKI proteins and I hypothesize that it is an important determinant of their functional properties. To test this hypothesis, I probed the interactions of peptides corresponding to D1 and D2 (LH domain deletion mutant) with the Cdk2/cyclin A complex. Results from isothermal titration calorimetry (ITC) and kinase inhibition assays, show that the interaction of either D1 or D2 with the Cdk2/cyclin A complex is less favorable relative to that of the p27-KID and that the D2 peptide cannot completely inhibit Cdk2/cyclin A kinase activity. These results indicate that the LH domain that couples D1 and D2 is necessary for the function of p27 as a Cdk inhibitor.

Despite having a conserved structure, the linker domains of the Cip/Kip proteins do not have a conserved sequence. However, the sequence of this domain in each Cip/Kip protein in different species is well conserved suggesting that sequence may play a role in the function of these proteins. I explored this sequence-divergence/structure-conservation relationship of the linker domains of the Cip/Kip proteins by constructing chimeric p27-KID molecules in which the p27 linker domain was replaced with the corresponding segments of either p21 or p57. ITC and kinase inhibition assay results show that the chimeric molecules bind and inhibit Cdk2/cyclin A in a manner similar to that of p27-KID. Moreover, thermal denaturation studies show that the complexes formed by these proteins bound to Cdk2/cyclin A have comparable melting temperatures. Taken together, these results indicate that the different linker domains, despite their structural differences, play similar structural roles in p27 binding to Cdk2/cyclin A.

In a third study, I investigated the consequences of perturbing the conserved structure of the linker domain. I successfully designed and prepared p27 variants in which domain LH was either more or less helical with respect to the wild-type protein. The secondary structural properties of the mutants were characterized by circular dichroism spectropolarimetry (CD). Thermal denaturation experiments showed that the ternary complexes of the p27 variants bound to Cdk2/cyclin A were less stable compared to the ternary complex formed by wild-type p27 (p27-KID^{wt}). Thermodynamic analysis showed that there was a decrease in the enthalpy of association of the mutants with Cdk2/cyclin A relative to p27-KID^{wt}. The free energies of binding varied within a much narrower range. In addition, *in vitro* Cdk2 inhibition assays showed that the p27 variants exhibited disparate inhibitory potencies. Further, when we over-expressed the p27 variants in mouse fibroblasts, they were less effective in causing cell cycle arrest relative to wild-type p27. These results indicate that the conserved structure of the linker domain is important for p27 function. It seems that nature has selected the sequence of the p27 LH domain to be partially structured and that bolstering or eliminating this structure is deleterious to function; intrinsic structure is critical for function.

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**PROBING THE ROLE OF NASCENT HELICITY IN PROTEIN FUNCTION:
p27^{kip1} AS A REGULATOR OF THE CELL CYCLE**

A Dissertation
Presented for
The Graduate Studies Council
The University of Tennessee
Health Science Center

In Partial Fulfillment
Of the Requirements for the Degree
Doctor of Philosophy
From The University of Tennessee

By
Steve Biko Otieno
May 2007

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Dedication

This dissertation is dedicated to my father, the late Mr. Elly Otieno, and my mother, Mrs. Margaret Otieno for their unwavering support.

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I would like to acknowledge all those who supported and assisted me during my research. First, I would to thank Dr. Richard Kriwacki for his tutelage and for giving me an opportunity to work in his laboratory. My gratitude is also extended to all members of the Kriwacki laboratory, the Structural Biology Department at St. Jude Children's Research Hospital, and the Department of Molecular Sciences at The University of Tennessee Health Science Center. I would also like to thank my family for the support they have accorded me in the course of graduate school. Finally, I acknowledge the National Institutes of Health (NIH) and the American Lebanese and Syrian Associated Charities (ALSAC) for funding my research project.

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List of Abbreviations

ATP.....	Adenosine triphosphate
CaCl ₂	Calcium chloride
CAK	Cdk-activating kinase
cAMP	Cyclic adenosine monophosphate
CD.....	Circular dichroism spectropolarimetry
Cdk	Cyclin-dependent kinase
Cip	Cyclin-dependent kinase inhibitory protein
CKI.....	Cyclin-dependent kinase inhibitor
CML	Chronic myelogenous leukemia
CREB.....	cAMP response element binding protein
D1	Cyclin-binding domain of p27
D2	Cdk-binding domain of p27
ΔG.....	Gibbs free energy change
ΔH.....	Enthalpy change
DISOPRED ..	Disorder prediction server
DLS.....	Dynamic light scattering
DMEM.....	Dulbecco's modified eagle medium
DNA	Deoxyribonucleic acid
ΔS	Entropy change
DTT.....	Dithiothreitol
<i>E.coli</i>	<i>Escherichia coli</i>
EDTA	Ethylene diaminetetraacetic acid
EGTA.....	Ethylene glycol bis (2-amino-ethylether)-N, N, N', N'-tetraacetic acid
FBS.....	Fetal bovine serum
FTIR.....	Fourier transformed infra-red spectroscopy
GFP	Green fluorescent protein
GST	Glutathione S-transferase
H ₂ O	Water
HIV.....	Human immuno-deficiency virus
HPLC	High performance liquid chromatography
IC ₅₀	Concentration giving 50% inhibition
IDP	Intrinsically disordered protein
IPTG	Isopropyl β-D-1-thiogalactopyranoside
ITC.....	Isothermal titration calorimetry
IUP	Intrinsically unstructured protein
K	Degrees Kelvin
kcal	Kilocalorie
KCl.....	Potassium chloride
kDa	Kilodaltons
KH ₂ PO ₄	Potassium phosphate (monobasic)
KID.....	Kinase inhibitory domain
Kip.....	Kinase inhibitory protein

KPC	Kip1 ubiquitination promoting complex
LB	Luria-Bertani media
LH	Linker helix domain of Cip/Kip proteins
MgCl ₂	Magnesium chloride
μl	Microliter
ml	Milliliter
μM	Micromolar
mM	Millimolar
mRNA	Messenger ribonucleic acid
Na ₂ HPO ₄	Sodium phosphate (dibasic)
Na ₃ VO ₄	Sodium orthovanadate
NaCl	Sodium chloride
NDP	Natively disordered protein
NFATC1	Nuclear factor of activated T-cells 1
ng	Nanogram
(NH ₄) ₂ SO ₄	Ammonium sulfate
nm	Nanometer
nM	Nanomolar
NMR	Nuclear magnetic resonance
NUP	Natively unstructured protein
oePCR	Overlap extension polymerase chain reaction
PBS	Phosphate buffered saline
P-Cdk2	Cdk2 phosphorylated at threonine 160
PDB	Protein data bank
PI	Propidium iodide
PO ₄ -H1	Phosphorylated histone H1
PONDR	Predictor of natural disordered regions
Rb	Retinoblastoma protein
RNAse	Ribonuclease
ROA	Raman optical activity
S200	Superdex 200
SASA	Solvent accessible surface area
SAXS	Small angle X-ray scattering
SCF	Skp, cullin, F-box containing complex
SDS-PAGE ..	Sodium dodecyl sulfate polyacrylamide gel electrophoresis
TFA	Trifluoroacetic acid
TGF-β	Transforming growth factor β
T _m	Melting temperature
UV	Ultra-violet
¹³ Cα	¹³ C – alpha

Chapter 1: Introduction

Self-replication is a fundamental property of eukaryotic cells. A complex, coordinated succession of events ensures that the parent cell is divided into two identical daughter cells. The sequence of events that take place between one cell division and the next is referred to as the cell division cycle or the cell cycle.^{1,2} This series of events can be separated into four distinct phases:

- G1 phase – characterized by cell growth, synthesis of new organelles, increased protein synthesis, and an accelerated metabolic rate.
- S phase – characterized by synthesis of deoxyribonucleic acid (DNA).
- G2 phase – the cells prepare for mitosis and checkpoints ensure that any insults to the genome since the previous division are not propagated. The G1, S and G2 phases are collectively referred to as interphase.
- M phase – the genetic material is divided into two identical fractions (mitosis) and the cell divides into two daughter cells each containing one half of the genetic material (cytokinesis).

Proliferating cells may exit the cell cycle temporarily, or reversibly, and enter into a state called quiescence or G₀ phase. Cells typically enter the G₀ phase from the G1 phase. Figure 1.1 shows the various phases of the cell cycle.

The cell cycle is one of the most extensively regulated processes in eukaryotic organisms. A number of proteins including kinases, kinase activators and inhibitors, phosphatases, and transcription factors, just to mention but a few, are involved in the modulation of the cell division cycle.

Figure 1.1 Schematic diagram of the eukaryotic cell division cycle.

1.1 Cyclin-dependent kinases

Strict temporal control of cell cycle progression is required for faithful duplication of eukaryotic cells. This is achieved predominantly through the activity of a family of highly conserved serine/threonine kinases called cyclin-dependent kinases (Cdk's).³⁻¹¹ The Cdk's govern phase transitions during the cell cycle in a timely manner and are referred to as "master timekeepers" of the cell cycle.¹² Two events are required for complete activation of the Cdk's: (i) a regulatory subunit called a cyclin binds to the Cdk;¹³⁻¹⁹ and (ii) the Cdk is phosphorylated on a conserved threonine residue (T160 in Cdk2) by a family of proteins called Cdk-activating kinases (CAK's).²⁰⁻²⁷ The kinase active site is located in the Cdk subunit of the heterodimer. Upon activation, the Cdk's phosphorylate a number of substrates orchestrating an orderly progression of the cell cycle from one phase to the next. As an example, Cdk4/cyclin D, Cdk2/cyclin A and Cdk2/cyclin E phosphorylate the retinoblastoma (Rb) protein facilitating the transition of proliferating cells from the G1 to the S phase.²⁸⁻³¹ In the hypophosphorylated state, the Rb protein represses transcription by binding and negatively regulating the activity of transcription factors such as the E2F family of transcription factors, chromatin remodeling complexes, and histone deacetylases.³²⁻³⁴ Phosphorylation of the Rb protein by the Cdk complexes relieves its repressor activity and activates the transcription factors culminating in progression from G1 to S phase.

The protein level of the cyclins oscillates depending on the phase of the cell cycle ensuring that the Cdk's are activated at the appropriate time.³⁵ The

abundance of the cyclins in any cell cycle phase is regulated through programmed synthesis and degradation.¹⁷ A classic example is the regulation of protein levels of cyclin E and cyclin B. The synthesis of cyclin E is induced in late G1 and its level peaks at the G1 to S transition.³⁶ As previously mentioned, cyclin E activates Cdk2 and promotes G1 to S transition. However, as the S phase starts and progresses, cyclin E activity is no longer required and its protein levels are reduced by degradation.¹⁷ Cdk1/cyclin B is required for the G2/M transition and cyclin B synthesis is induced in late S phase.³⁷ The cyclin B accumulates in complexes with Cdk1 as the cells approach the G2 to M transition.³⁷ When the cells enter the M phase, cyclin B is rapidly degraded.³⁷ In addition to cyclins, other regulators of cell cycle are also temporally degraded to ensure proper progression of the cell cycle.^{35,38} Protein degradation is therefore an important regulatory mechanism in cell cycle progression.

The Cdk's were originally discovered in yeast where they were shown to be indispensable players in progression of the cell cycle. The *Saccharomyces cerevisiae* Cdk, Cdc2, and the *Saccharomyces pombe* Cdk, Cdc28, were the first to be identified.³⁹⁻⁴¹ The human orthologs were soon discovered based on their capacity to complement yeast mutants.⁴² In yeast, only a single Cdk regulates cell cycle events. In contrast, the cell cycle in higher eukaryotes is regulated by multiple Cdk's. To date, 11 Cdk's (Cdk 1-11) and 9 Cdk-related proteins have been identified.⁴ It is important to note that not all the Cdk's and Cdk-related proteins are involved in cell cycle progression (Table 1.1). Only Cdk's 1, 2, 4, and 6 have been directly implicated in cell cycle progression. On the other hand,

Table 1.1 Cyclins, cyclin-dependent kinases and their biological roles.

Cdk	Associated cyclin(s)	Function
Cdk1	Cyclins A, B, E	Regulation of G1-S, G2-M transitions
Cdk2	Cyclins A, E	Regulation of G1-S transition
Cdk3	Cyclins E, A, C	Cell cycle exit
Cdk4	Cyclins D1, D2, D3	Regulation of G1-S transition
Cdk5	Cyclins p35, p39	Neuronal regulation
Cdk6	Cyclins D1, D2, D3	Regulation of G1-S transition
Cdk7	Cyclin H	Cdk-activating kinase
Cdk8	Cyclin C	Regulation of transcription
Cdk9	Cyclins T1, T2, K	Regulation of transcription
Cdk10	Unknown	Regulation of transcription
Cdk11	Cyclins L1, L2	Regulation of transcription

Cdk7 acts as a CAK regulating the activity of other Cdk's^{43,44} and Cdk5 is predominantly expressed in neurons where it is involved in the regulation of neuronal functions including synaptic transmission and cytoskeletal remodeling.^{45,46} Other Cdks like Cdk8 and Cdk9 are mainly involved in regulation of transcription.⁴⁷⁻⁵⁰ The best characterized members of the Cdk family are those involved in cell cycle progression.

Although Cdk's have different regulatory roles in cells, their catalytic domains are highly conserved and have a similar three-dimensional structure. Past studies have shown that in the unbound Cdk's, the active site is not properly configured for catalysis.⁵¹ The binding of a cyclin subunit and CAK phosphorylation induces conformational changes which enable the Cdk active site to become catalytically competent.^{52,53} In addition to Cdk activation, the cyclin subunit also contains the binding site for a number Cdk substrates.⁵⁴ The binding site is composed of a conserved set of residues that form a hydrophobic patch. Key cell cycle regulatory proteins including Rb have been shown to bind to this substrate-binding motif.⁵⁴ Cdk activity is also regulated by inhibition.

Phosphorylation within the N-terminal region of Cdk's can lead to inactivation. This inactivating modification occurs on a single tyrosine residue in Cdk's 4 and 6 (Y17 and Y24 of Cdk4 and Cdk6, respectively), while Cdk1 and Cdk2 are phosphorylated on adjacent threonine and tyrosine residues (T14 and Y15 on Cdk1 and Cdk2).⁵⁵⁻⁵⁹ Previous studies have demonstrated that Cdk1 and Cdk2 are phosphorylated by Wee1 and Myt1 kinases.⁶⁰⁻⁶⁴ However, the kinase(s) that phosphorylate Cdk's 4 and 6 have not yet been identified. This

inactivating phosphorylation event is reversed by the activity of Cdc25 family of phosphatases.⁶⁵⁻⁶⁷ In addition to phosphorylation, Cdk activity is also inhibited by a family of proteins termed the cyclin-dependent kinase inhibitors.

1.2 Cyclin-dependent kinase inhibitors

The cyclin-dependent kinase inhibitors (CKI's) negatively regulate the activity of the Cdk's.^{68,69} In humans, CKI's are divided into two families: the Ink4 family⁷⁰⁻⁷³ that includes p15^{Ink4b}, p16^{Ink4a}, p18^{Ink4c}, and p19^{Ink4d}; and the Cip/Kip family⁷⁴⁻⁸⁸ that includes p21^{Waf1/Cip1/Sdi1/Cap20}, p27^{Kip1}, and p57^{Kip2}. The Ink4 family inhibits Cdk activity by binding to the catalytic subunit of Cdk's 4 and 6 abolishing their ability to form active complexes with D-type cyclins⁸⁹ (cyclin D1, cyclin D2 and cyclin D3; expressed in an organ and tissue specific manner³⁷). Figure 1.2 shows the structures of p19/Cdk6, p16/Cdk6 and Cdk2/cyclin A complexes. The PDB accession numbers for the p16/Cdk6, p19/Cdk6 and Cdk2/cyclin A are 1BI7, 1BI8, and 1FIN, respectively. The binding of p19 and p16 to Cdk6 distorts the cyclin binding site (compare with the cyclin binding site in the Cdk2/cyclin A complex). By blocking cyclin D from binding, p19 and p16 maintain Cdk6 in an inactive state. Cdks adopt the bilobed fold common to many protein kinases.⁹⁰ This fold is comprised of an N-terminal lobe that is largely comprised of β -sheets and a C-terminal lobe that is mainly comprised of α -helices. The binding of p19 and p16 to Cdk6 push the N and C terminal lobes away from each other resulting in deformation of the binding site for the D-type cyclins. In addition, the two inhibitors distort the ATP binding pocket of Cdk6 contributing to further inhibition.

Figure 1.2 Structures of the Cdk2/cyclin A (PDB; 1FIN), p19/Cdk6 (PDB; 1B18), and p16/Cdk6 (PDB; 1B17). The binding of p19 and p16 to the Cdk6 distorts the cyclin binding site preventing the formation of Cdk6/cyclin D complexes.

On the other hand, members of the Cip/Kip family of CKI's inhibit Cdk catalytic activity by binding to the heterodimeric complex over an extended surface spanning from the cyclin to the Cdk subunit. The Cip/Kip proteins inhibit a broad spectrum of Cdk complexes and have been shown to regulate the activity of Cdk's in complex with cyclins D, E and A.^{91,92} Figure 1.3 shows the versatility of the Cip/Kip proteins in Cdk regulation during the G1/S transition. Each Cip/Kip protein family member contains an amino terminal domain that has been shown to be necessary and sufficient for Cdk inhibition. This functionally conserved segment is known as the kinase inhibitory domain (KID). The sequences of the KID's of p21 (residues 11 - 94) and p57 (residues 23 - 106) with respect to p27 (residues 22 - 106) are 48% and 42% identical. The Cdk regulatory activity of the Cip/Kip proteins causes cell cycle arrest during the G1/S transition. This phase transition is controlled by Cdk1/cyclin A or E, Cdk2/cyclin A or E, Cdk4/cyclin D, and Cdk6/cyclin D. While the inhibition of Cdk1 and Cdk2 complexes by the Cip/Kip proteins is well defined, their effect on Cdk's 4 and 6 is less clear. Early studies on the interaction of the Cip/Kip proteins with Cdk4/cyclin D and Cdk6/cyclin D complexes reported that the Cip/Kip proteins promote assembly of Cdk4/cyclin D and Cdk6/cyclin D complexes, in essence acting as activators.^{93,94} However, reports from other groups have cast doubt on these observations.^{95,96} It should be noted that a mechanism has recently been reported that could possibly explain the apparently paradoxical functions of the Cip/Kip proteins.⁹⁷ The study showed that phosphorylation of p27 by the oncogenic tyrosine kinase, BCR-ABL, on a tyrosine residue (Tyr88) that is

Figure 1.3 The Cip/Kip proteins regulate the activity of a broad range of Cdk/cyclin complexes.

inserted into the ATP binding pocket during Cdk inhibition, ejects the tyrosine out of the active site, consequently relieving inhibition of kinase activity. The authors propose that this mechanism also takes place in the interaction of p27 and Cdk4/Cdk6 thereby explaining the seemingly paradoxical functions of p27.

The first member of the Cip/Kip family to be identified was p21. p21 inhibits cell proliferation in response to genomic insults and is up-regulated in irradiated cells and in cells exposed to DNA damaging drugs. The up-regulation of p21 is mediated by p53.^{98,99} In contrast to p21, p27 expression is induced by diverse differentiation and anti-mitogenic stimuli including, transforming growth factor- β (TGF- β),^{80,100} rapamycin,¹⁰¹ contact inhibition,⁸⁰ depletion of growth factors,¹⁰² and loss of cell anchorage.¹⁰³

The structure of p27 bound to Cdk2/cyclin A provides detailed insights into the molecular basis of Cdk inhibition by the Cip/Kip proteins.¹⁰⁴ The KID fragment of p27 was used to prepare the ternary complex. Figure 1.4 shows the structure of the p27-KID/Cdk2/cyclin A complex (PDB accession number, 1JSU). In the complex, p27-KID folds into an extended, non-globular structure consisting of (from N to C terminus): an extended segment, an amphipathic α -helix, an amphipathic β -hairpin, a β -strand and a 3_{10} helix. The extended segment (domain D1) binds to the cyclin A subunit of the kinase complex, the amphipathic α -helix (domain LH) links the cyclin binding segment to the Cdk binding segment (domain D2) which is comprised of the amphipathic β -hairpin, the β -strand, and the 3_{10} helix. The extended segment binds to the conserved substrate binding site on the cyclin A subunit. The amphipathic α -helix does not make a significant

Figure 1.4 The structure of the p27/Cdk2/cyclin A complex (PDB; 1JSU). p27 binds to the Cdk2/cyclin A complex over an extended surface spanning from the cyclin A to the Cdk2 subunit. Binding to cyclin A prevents substrate binding and the binding of the 3₁₀ helix of p27 into the ATP binding pocket impedes ATP binding.

number of contacts with the kinase complex. The extended structure of the p27-KID enables it to interact with the Cdk2/cyclin A complex over a large surface area (a total buried surface area of 5,752 Å²). Most of the hydrophobic residues on p27-KID pack against the Cdk2 and cyclin A subunits. From this structure, the binding of p27-KID inhibits the Cdk kinase activity in three ways. First, the binding of p27-KID to the substrate binding site on the cyclin obstructs the docking of Cdk2 substrates. Secondly, p27-KID binding deforms the catalytic cleft on the Cdk2. This is caused by interactions between p27-KID and an amino-terminal β -sheet of Cdk2. Finally, the 3₁₀ helix of p27-KID is inserted into the catalytic cleft preventing ATP binding.¹⁰⁴ This model of inhibition has been extended to p21 and p57 since the three proteins have very similar primary structures.

1.3 p27 kinase inhibitory protein 1 (p27^{kip1})

Human p27 is comprised of 198 amino acids and, in addition to cell cycle regulation, has been implicated in regulating differentiation,^{105,106} and more recently the control of cell motility.¹⁰⁷ p27 is up-regulated in serum-starved and contact inhibited cell cultures resulting in cell cycle exit (quiescence). Suppression of p27 expression using anti-sense mRNA abrogates cell cycle exit.¹⁰⁸ When quiescent cultures are supplemented with mitogens, p27 is eliminated from cells and the cells re-enter the cell cycle.¹⁰⁹ However, quiescent cells do not re-enter the cell cycle, even in the presence of mitogens, when the elimination of p27 is blocked by treatment with cyclic adenosine monophosphate

(cAMP) analogs, rapamycin, or hypoxia.^{101,110,111} These results demonstrate that p27 is a downstream effector of anti-mitogenic signals.¹¹² The *in vivo* effects of p27 deficiency have been investigated using knock-out mice. Mice lacking p27, though viable, exhibit hyperplasia in multiple organs and gigantism with respect to their wild-type counterparts.¹¹³⁻¹¹⁵ As a negative regulator of cell cycle progression, p27 is an oncosuppressive protein. Circumvention of p27 cell cycle control has been linked to a myriad of human malignancies. The loss of function in most tumor suppressor genes is usually a consequence of deletion of one or both alleles of the gene(s). However, the p27 gene is rarely deleted in human cancers. Mouse model studies have shown that p27 is a haploinsufficient tumor suppressor.¹¹⁶ Mice with a single allele of the p27 gene have an intermediate frequency of tumor formation relative to their wild-type and null counterparts.^{116,117}

1.4 p27 and cancer

One of the hallmarks of many types of cancer is the reduced level of p27 protein relative to normal cells. However, this reduction in p27 protein levels does not stem from p27 gene mutations.¹¹⁸ In fact, germline mutations in the p27 gene are very rare.¹¹⁹⁻¹²¹ Studies of breast carcinoma have revealed that decreased p27 protein levels promote tumor progression and are associated with poor prognosis.¹²²⁻¹²⁴ The same has also been observed in colorectal,¹²⁵ lung non-small cell,¹²⁶ gastrointestinal,¹²⁷ prostate,¹²⁸ oral cavity,¹²⁹ and ovarian carcinomas.¹³⁰ In most of these cancers, the p27 mRNA transcript is expressed

at a normal level indicating that the low level of p27 protein is caused by protein degradation. There are two main ways by which tumor cells evade cell cycle regulation by p27: (i) degradation of p27 at an accelerated rate (cause of reduced p27 protein level)¹³¹⁻¹³³ and (ii) sequestration of p27 in the cytoplasm preventing it from inhibiting the nuclear-localized Cdk's.¹³⁴ Like many other cell cycle regulatory proteins, degradation is one of the main ways by which cells regulate p27 levels. The degradation of p27 is controlled in two ways: cytoplasmic ubiquitination by the KPC (Kip1 ubiquitination promoting) complex¹³⁵ and nuclear ubiquitination by the SCF^{skp2} complex.^{136,137,138} The two complexes chemically ligate poly-ubiquitin chains to lysine residues on p27. Prior to being poly-ubiquitinated by the KPC and SCF^{skp2} complexes, p27 must be phosphorylated on Ser10 and Thr 187, respectively. After ubiquitination, p27 is degraded by the 26S proteasome.¹³⁹ To be recognized by the SCF complex, p27 has to be in complex with either Cdk2/cyclin A or Cdk2/cyclin E.^{140,141} In most malignancies, there are increased levels of the skp2 protein leading to accelerated degradation of p27.¹⁴²⁻¹⁴⁵ Table 1.2 shows different types of cancer and the alterations in the p27 protein. It has also been reported that reduced expression of p27 plays a role in cancer drug resistance. Resistance to docetaxel, one of the most effective chemotherapeutic agents used in breast cancer treatment, has been related to reduced expression of p27.¹⁴⁶ It is therefore not surprising that designing small molecule inhibitors that mimic p27 inhibitory activity has generated a lot of interest.

Table 1.2 Alteration of the p27 protein in human malignancies.

Malignancy	Alteration in p27 protein
Breast cancer	Accelerated degradation ¹⁴⁷ Cytoplasmic localization ¹⁴⁸
Prostate cancer	Accelerated degradation ¹⁴⁹ Cytoplasmic localization ¹⁵⁰
Ovarian cancer	Accelerated degradation ¹⁵¹ Cytoplasmic localization ¹⁵²
Colorectal cancer	Accelerated degradation ¹⁵³
Non-small cell lung cancer	Accelerated degradation ¹⁴⁵
Glioma	Accelerated degradation ¹⁵⁴
Lymphoma	Accelerated degradation ¹⁵⁵
Chronic myelogenous leukemia (CML)	Accelerated degradation ¹³³
Liver cancer	Accelerated degradation ¹⁵⁶ Cytoplasmic localization ¹⁵⁷
Oral squamous cell carcinoma	Accelerated degradation ¹⁴³

1.5 p27 is an intrinsically unstructured protein

The classical structure-function paradigm postulates that proteins must be folded into distinct three-dimensional conformations in order to function.

However, there is now sufficient evidence that supports the existence of functional proteins and protein domains that are intrinsically disordered under physiological conditions.¹⁵⁸⁻¹⁶⁵ As used herein, intrinsic disorder of a protein or protein domain means that under physiological conditions: (i) the isolated protein or protein domain lacks tertiary structure; (ii) there are no significant, long-range interactions between its residues; and (iii) it exists as an ensemble of many highly dynamic conformations. Intrinsic disorder in proteins is either classified as global (disorder of the entire protein), or local (disorder of a protein domain).

Proteins that exhibit native structural disorder belong to a fast growing family of proteins referred to either as intrinsically unstructured proteins (IUP's), intrinsically disordered proteins (IDP's), natively unstructured proteins (NUP's), or natively disordered proteins (NDP's). For the purposes of this dissertation, I will use the term intrinsically unstructured proteins (IUP's). It is now estimated that about one third of the human genome is comprised of sequences that encode proteins that have local or global intrinsic structural disorder. Further analyses have shown that the prevalence of IUP's in a particular species varies directly in proportion with its genomic complexity.¹⁶⁶ The IUP's are involved in key cellular processes including, but not limited to, DNA recognition (DNA binding domains of the GCN4 and NFATC1),¹⁶⁷⁻¹⁷⁰ transcriptional activation (kinase-inducible activation domain of CREB, activation domains of p53 and VP16),¹⁷¹⁻¹⁷⁵ RNA

recognition (RNA recognition segments of HIV-1 Rev and BIV Tat proteins),¹⁷⁶⁻¹⁷⁹ translation (eIF4E binding segment of eIF4G),¹⁸⁰ microtubule and actin dynamics (caldesmon, thymosin, stathmin/RB3),¹⁸¹⁻¹⁸³ and cell cycle regulation (p21, p27, p57). Table 1.3 shows a list of some IUP's and their biological functions. IUP's have unique properties that distinguish them from globular proteins.

Relative to globular proteins, the primary structures of IUP's are characterized by low sequence complexity and compositional bias.¹⁸⁴ They are enriched in amino acids that promote structural disorder such as Ser, Ala, Gly, Arg, Gln, Pro, Glu, and Lys, and relative to globular proteins, exhibit reduced levels of residues that promote structural order such as Trp, Cys, Phe, Ile, Asn, Val, Tyr, and Leu.^{184,164} Globular proteins typically have a comparatively larger number of order promoting hydrophobic amino acids that form their cores and facilitate folding. Bioinformatics programs that predict structural disorder based on the amino acid attributes of the IUP's with a high degree of accuracy have been developed. Examples include PONDR,¹⁸⁵ DISOPRED,¹⁸⁶ and GLOBPLOT.¹⁸⁷ The sequence properties of IUP's indicate that their intrinsic disorder is encoded in their primary structures as is the case for the secondary and tertiary structures of globular proteins. Due their intrinsic disorder, IUP's often exhibit extended conformations and have larger hydrodynamic radii relative to globular proteins of comparable mass.¹⁶³ Because of their extended structures, IUP's are sensitive to proteolytic cleavage.¹⁸⁸ In addition, IUP's are heat stable,⁸¹ have high net charge,¹⁶³ and low mean hydrophobicity.¹⁶³ The disordered structures of IUP's have been characterized using X-ray

Table 1.3 Biological functions of IUP's.

Protein	Function
Naked2 ¹⁸⁹	Vesicle transport
Bim, Bad, BMF ¹⁹⁰	Apoptosis
C-terminal domain of CtBP ¹⁹¹	Nuclear transcriptional co-repression, golgi membrane tubule fusion
C-terminal domain of the measles virus nucleoprotein ^{159,192}	Measles virus RNA synthesis
Thymosin- β ¹⁹³	Actin dynamics
Acidic activation domain of NF- κ B ¹⁹⁴	Transcriptional activation
Basic DNA-binding region of the leucine zipper protein GCN4 ¹⁶⁷	Transcriptional activation
Amino-terminal domain of p53 ¹⁷³	Onco-suppressing transcription factor
CITED2 ¹⁹⁵	Hypoxic response
High mobility group (HMG) domain, of LEF-1 ¹⁹⁶	Regulation of the T-cell receptor-a gene enhancer

crystallography, nuclear magnetic resonance (NMR), circular dichroism (CD) spectroscopy, raman optical activity spectroscopy (ROA), fourier-transformed infra-red spectroscopy (FTIR), analytical ultracentrifugation (AUC), dynamic light scattering (DLS), small angle X-ray scattering (SAXS), and fluorescence spectroscopy. Although IUP's have disordered structures prior to binding their targets, NMR and CD studies have demonstrated that some IUP's have segments that exhibit a natural propensity for nascent secondary structures.^{197,198,188,199} One key property of most IUP's is that they fold on binding their targets and become ordered.^{197,175,200} This folding-on-binding phenomenon was manifested in the interaction of p27 and Cdk2/cyclin A complex.¹⁹⁷

p27 is a globally unstructured IUP with segments that exhibit nascent secondary structure. Of the three Cip/Kip proteins, p27 is the best characterized structurally because it can be expressed at high levels in *E. coli*, can be purified to homogeneity, and is highly soluble. p27 exhibits many characteristics of IUP's. Consistent with the properties of IUP's, full-length p27 migrates at about 27 kDa during a sodium dodecyl sulfate polyacrylamide gel electrophoresis (the calculated molecular weight of p27 is 22 kDa) indicating that it is a highly extended polypeptide with a large hydrodynamic radius.⁸⁰ Moreover, analysis of p27 by limited proteolysis shows that free p27 is very susceptible to cleavage.²⁰¹ However, when p27 interacts with the Cdk2/cyclin A complex, it is refractory to proteolysis.²⁰¹ The secondary structures of p27, p21 and p57 determined by CD at low concentrations are strikingly similar.^{188,198,202} The three proteins are

largely unstructured with a low population of helical secondary structure. The CD spectra show that the native structural disorder of the Cip/Kip proteins is well conserved. It should be noted that the results on secondary structure discussed above are derived from studies of KID constructs and not the full-length proteins. Like the KID's, the full-length proteins are also largely unstructured. NMR analysis shows that full-length p27 and p27-KID have very similar secondary structures.¹⁹⁷ The conservation of the disordered structures of the Cip/Kip proteins argues strongly that structural disorder plays a significant role biological function. p27 folds dramatically on binding the Cdk2/cyclin A complex and attains structural order as shown in Figure 1.4, above. The plasticity of the structure of p27 and its template driven folding on binding enables p27 to interact with a broad range of Cdk complexes.²⁰³ The folding on binding mechanism affords high specificity but at a great entropic cost. It therefore follows that the thermodynamic contributions of the binding-on-folding process due to enthalpy and entropy have to be appropriately balanced so that binding occurs spontaneously.

1.6 Thermodynamics of the association of p27 and Cdk2/cyclin A

The KID's of the Cip/Kip proteins are comprised of three domains: (i) a cyclin-binding domain (hereafter referred to as D1); a linker domain (hereafter referred to as LH); and (iii) a Cdk binding domain (hereafter referred to as D2). D1 has a motif that binds to the substrate binding site on the cyclin subunit that is comprised of Leu32, Phe33, and Gly34. These three residues are conserved

amongst the Cip/Kip proteins. On D2, one key residue is Tyr88 which is inserted into the ATP binding pocket when p27 interacts with the Cdk2/cyclin A complex. Tyr88 is part of the 3_{10} helix of p27 that was discussed in Section 1.2. Figure 1.5 shows a sequence alignment of the KID's of members of the Cip/Kip family. The sequences D1 and D2 are well conserved amongst the three family members, but the sequence of domain LH is not conserved. Only 3 of the 22 residues in this domain are conserved. The number of residues in domain LH, 22, is the same for all the Cip/Kip proteins. As discussed above, the secondary structures of the Cip/Kip proteins are mainly unstructured with nascent helical properties. Nuclear magnetic resonance (NMR) ^{13}C -alpha ($^{13}\text{C}\alpha$) chemical shift analysis showed that D1 and D2 are largely unstructured and the nascent helical structure observed using CD is localized in the LH domain.¹⁹⁷ The nascent helical structure of the linker domain is conserved amongst all three members of the family.^{188,198,202} This sequence divergence/structure conservation relationship is atypical since the domains whose sequences are conserved, D1 and D2, are unstructured, and the domain whose sequence is poorly conserved, LH, exhibits conserved structural features.

Thermodynamically, the folding of p27 on binding the Cdk2/cyclin A is very unfavorable since it requires a large negative entropy change. Lacy, *et al.*, demonstrated that this large negative entropic cost is overcome by the quantity of heat (enthalpy associated with binding, ΔH) released during the extensive folding of p27 on binding to Cdk2/cyclin A.¹⁹⁷ This results in a negative Gibbs free energy change (ΔG) for binding and spontaneous binding. The interaction of p27

Figure 1.5 Sequence alignment of the KID domains of the Cip/Kip proteins.

and the Cdk2/cyclin A complex is therefore an enthalpically driven process. The heat capacity change associated with binding (ΔC_p) was found to be large and negative, and comparable to values obtained from protein folding events. The large negative heat capacity change is an indicator of the extent to which hydrophobic interactions contribute to the formation of the ternary complex.^{204,205} Lacy, *et al*, further demonstrated that the interaction of p27 with the Cdk2/cyclin A complex is more favorable than the interaction of p27 with either cyclin A or Cdk2 alone. The binding of p27 to the Cdk2/cyclin A complex is sequential, with p27 binding to the cyclin A subunit first, followed by folding of the linker domain, and finally the folding of D2 into a β -hairpin, a β -strand and a 3_{10} helix. The kinetics of association and dissociation of p27 and cyclin A are much faster relative to those of p27 and Cdk2. While the functions of D1 and D2 are well characterized, the role of domain LH is not well understood. Is it just a spacer between D1 and D2 or does it play a role in the binding process? If so, why does it have a conserved structure?

1.7 Project hypotheses and strategy

To understand the role of the linker domain of p27, I first investigated the interactions of D1 and D2 with P-Cdk2/cyclin A (a mutant with the linker domain deleted). Given the structural conservation of the linker domain of p27, I hypothesize that linker domain is necessary for the interaction of p27 and the P-Cdk2/cyclin A complex and for p27 to perform its biological function.

Although the sequence of domain LH is not conserved amongst members of the Cip/Kip protein family, it is very highly conserved for each protein in different species. Figure 1.6 shows an alignment of the sequences of p27 LH domains in different species. In mammals (from mouse to man – a period of over 100 million years in the evolutionary timeline), the sequence of the p27 linker domain is 100% conserved and has resisted selective pressure. This indicates that the sequence of each individual protein may be significant in determining its functional properties. Otherwise, there would have been no reason for these sequences to be selected for and to persist during evolution. However, the evolutionary stability of these sequences raises a question as to whether it is the sequence or the structure of the LH domain that is important in determining the biophysical and functional properties of these proteins. To answer this question, I designed p27-KID mutants in which the LH domain of p27 was replaced with the LH domains of either p21 (p27-KID^{p21helix}) or p57 (p27-KID^{p57helix}) and studied their interaction with the Cdk2/cyclin A complex. These mutants are hereafter referred to as the chimeric mutants. Since D1 and D2 are well conserved in the three proteins, the chimeric mutants should be suitable for studying the sequence-divergence/structure-conservation relationship of the linker domains of the three Cip/Kip proteins. I hypothesize that if the structure of the LH domain is the dominant determinant of the biophysical and functional properties of the Cip/Kip proteins on interacting with the Cdk2/cyclin A complex, then the properties of the p27-KID^{wt} relative to the chimeric mutants would be very similar (recall that amongst the Cip/Kip proteins, the structures are conserved but the

Frog	HDELRSSELKRQLKEIQASDCQR	32
Pig	HEELTRDLEKHCRDMEEASQRK	37
Cat	HEELTRDLEKHCRDMEEASQRK	37
Mink	HEELTRDLEKHRRDMEEASQRK	37
Human	HEELTRDLEKHCRDMEEASQRK	37
Hamster	HEELTRDLEKHCRDMEEASQRK	37
Rat	HEELTRDLEKHCRDMEEASQRK	37
Mouse	HEELTRDLEKHCRDMEEASQRK	37
Chicken	HEELNRDLKKHRKEMEEACQRK	37
Zebrafish	HEELKKDFQRQLRTMEDASAEA	37

Figure 1.6 Alignment of p27-LH domains from different species. The sequences of LH domains from mammals are circumscribed in the red box.

sequences are different). On the other hand, if the sequence of the LH domain is the dominant determinant of the properties of the interaction between the Cip/Kip proteins and P-Cdk2/cyclin A, then the properties of the interaction of the chimeric mutants with P-Cdk2/cyclin A relative to p27-KID^{wt} would be different. In addition to the sequence-structure relationship of the linker domains of the Cip/Kip proteins, I also studied the role of the three conserved residues in domain LH by mutating the residues to glycines (Glu39Gly/Leu41Gly/Arg43Gly, the p27-KID^{ΔC} mutant) and probed the interaction of this mutant with the P-Cdk2/cyclin A complex.

Many IUP's exhibit a small amount of nascent helical secondary structure in the absence of their binding partners. The polypeptide segments giving rise to these features typically fold into α -helices upon binding their biological targets. I propose that one of the roles of the nascent helical structures is to attenuate the entropic cost associated with the folding-on-binding process. I further hypothesize that increasing the helical content of the linker domain would decrease the entropic cost associated with the folding on binding and thereby decrease the Gibbs free energy of interaction of the p27-KID and a Cdk/cyclin complex. Conversely, decreasing the helical content would increase the entropic cost and make the reaction less favorable. One underlying assumption of these hypotheses is that all the mutants fold in exactly the same way as the wild-type protein. To test these hypotheses, p27-KID mutants which have LH domains that are either more (p27-KID^{+H}) or less (p27-KID^{-H}) helical relative to the wild-type p27-KID (p27-KID^{wt}) were designed and their thermodynamic and functional

properties on binding to the Cdk2/cyclin A complex probed. In addition, I also designed a mutant in which the LH domain of p27 was replaced with a flexible loop, 22 residues long, and performed similar studies.

Chapter 2: Materials and methods

2.1 Design of the more and less helical p27-KID linker domain mutants

My approach of designing p27-KID mutants with linker domains that were either more or less helical than the wild-type p27-KID was to substitute solvent exposed residues within the linker domain with residues that are known to either stabilize or destabilize helices. Mutating the solvent exposed residues minimizes disruption of contacts between p27 and the Cdk2/cyclin A complex. This reduces the possibility that the results obtained from the succeeding studies are a consequence of the disruption of contacts between p27 and Cdk2/cyclin A. The solvent accessible surface area (SASA) for each residue within the linker domain was analyzed using the GETAREA program.²⁰⁶ The input used for the program was the pdb file of the p27-KID/Cdk2/cyclin A complex (1JSU). Residues that had more than 50% of their total surface area exposed to solvent were selected as suitable candidates for mutagenesis. From the output, 10 residues had more than 50% of their surface areas exposed (Table 2.1). The 10 candidate residues were further analyzed using the Insight II program (Accelrys, San Diego, CA). The PDB file, 1JSU, was used as an input and all the atoms within a 6 Å radius of every atom of each candidate residue were analyzed. This was done to determine whether any of the candidate residues interacted with the residues of the binary complex. No close contacts were observed for the 10 candidate residues. After this analysis, all the 10 candidate residues were determined to be suitable for mutagenesis. Figure 2.1 shows the orientation of the side chains of

Table 2.1 Solvent exposure of the residues in the linker domain of p27.

Residue	Percentage of surface area exposed to solvent
His38	40.9
Glu39	84.1
Glu40	79.9
Leu41	1.9
Thr42	22.7
Arg43	56.7
Asp44	33
Leu45	0.4
Glu46	26.9
Lys47	76.2
His48	27.2
Cys49	13.2
Arg50	67.1
Asp51	55.4
Met52	20.2
Glu53	37.9
Glu54	81.6
Ala55	87.1
Ser56	14.6
Gln57	30
Arg58	81.3
Lys59	56.1

The residues selected for mutagenesis are in bold font.

Figure 2.1 The structure of p27/Cdk2/cyclin A (PDB; 1JSU) showing the residues determined to be suitable for mutagenesis. The residues are colored red for acidic residues, blue for basic residues, and green for hydrophobic residues. Cdk2 is colored orange and cyclin A cyan.

the selected residues in the structure of the p27/Cdk2/cyclin A structure (PDB; 1JSU).

The stability an α -helix is determined both by the helix forming propensity of the amino acids in its primary structure and the interactions between side chains.²⁰⁷ Two different strategies were used in the design of mutants with stabilized linker domains. In the first approach, alanine was used as a possible stabilizer of the helical structure of the linker domain. Two alanine mutants were designed. In the first alanine mutant, all the residues that were selected for mutagenesis were substituted with alanines. Alanine was chosen because previous studies have shown that alanines stabilize and have a high propensity to occur in α -helices.²⁰⁸⁻²¹⁰ This mutant, in which the 10 candidate residues were substituted with alanines, was designated p27-KID^{A10}. In the second alanine mutant, all the 22 residues in the linker domain were substituted with alanines. This mutant was designated p27-KID^{A22}. In the second approach, leucine was used as a potential stabilizer of the linker domain helix. Like alanine, leucine has also been shown to stabilize and have a high propensity to occur in α -helices.^{207,209} Hydrophobic interactions between leucine side chains on one face of an α -helix shield the backbone amides and carboxyl groups from hydrogen bonding with water, thus stabilizing the helical conformation.²¹¹ One disadvantage of leucine, however, is its hydrophobic nature. To counter the hydrophobicity of leucine, I designed a library of mutants in which the number of leucines replacing the candidate residues was balanced with charged, polar residues. The substituted residues in these mutants consisted of either 7

leucines and 3 polar residues, 6 leucines and 4 polar residues, 5 leucines and 5 polar residues, etc. I also ensured that the polar residues had a high helical propensity. The leucines were strategically positioned three and four residues apart in the helix. Past studies have shown that leucines positioned three and four residues apart stabilize α -helices.²¹² In addition, leucines positioned four residues apart is one of the most commonly observed pairwise side chain interactions in protein α -helices.²¹³ A total of 6 mutants were designed. The mutants were designated p27-KID^{SL1} (SL1; stabilized by leucine, construct 1), p27-KID^{SL2}, p27-KID^{SL3} (p27-KID^{+H}), p27-KID^{SL4}, p27-KID^{SL5}, and p27-KID^{SL6}. In the p27-KID^{SL6} mutant, all the candidate residues were substituted with leucines. This mutant was designed to test if charged residues that were determined to be unsuitable for mutagenesis could counter the hydrophobicity of the 10 leucines.

For the less helical mutant, the candidate residues were replaced with glutamate (Glu) residues. The premise behind choosing Glu residues is that electrostatic repulsion between the negatively charged Glu side-chains should destabilize the helix backbone.²¹⁴ In addition, the substituting residues close to the carboxy terminus of the helix with Glu residues would result in further destabilization of the helix due to repulsion of the negatively charged Glu side-chain and the negative carboxy-terminus helix dipole.²¹⁵ Another advantage of using glutamate is that the polar Glu side-chain should enhance the solubility of this mutant. This mutant was designated p27-KID^{-H}.

A mutant with a flexible loop comprised of mainly glycines, a known helix breaker, and other residues with a low propensity for being in α -helices was also

designed. The residues that comprised this mutant were chosen with an objective of minimizing helical secondary structure in the linker domain. The number of residues in the loop equaled 22, as in the p27 linker domain. This mutant was designated p27-KID^{loop}. Table 2.2 shows the linker domain sequences of all the mutants designed.

2.2 Preparation of mutagenesis cassettes

The p27-KID linker domain mutants were cloned by cassette mutagenesis. First, the deoxyribonucleic acid (DNA) sequence encoding residues 22-71 of p27 (including the cyclin binding domain, the linker domain, and the Cdk binding domain up to E71) was determined. The bases encoding the wild-type p27 linker domain residues were changed such that they encoded the linker domain sequences of the mutants. Oligonucleotides used in the preparation of the mutagenesis cassettes were then designed. For each mutant, a forward and reverse oligonucleotide, each 104 bases long, with 40 overlapping complementary bases and end primers for each oligonucleotide, were designed. The sequences of the oligonucleotides used for the preparation of the mutagenesis cassettes are shown on Table A.1. End primers with the sequences 5' GTAACGCATATGGAGCACCCCAAGCCCTCGGCC 3' and 5' CCCC GCCTCGAGGGGTTTGTGATTCTGAAAATCGA 3' were designed for the p27-KID^{p21^{helix}} and p27-KID^{p57^{helix}} mutants. The sequences for the end primers designed for the other mutants were: 5' GCCTCCCATATGGAGCAC-CCCAAG 3' and 5' CTTGGGCTCGAGGGGTTTGTGATTC 3'. Included in the

Table 2.2 Linker domain sequences for the p27-KID variants designed to be more and less helical.

Variant	Sequence of linker domain
p27-KID ^{wt}	H <u>EE</u> L <u>T</u> <u>R</u> D <u>L</u> E <u>K</u> H <u>C</u> <u>R</u> D <u>M</u> E <u>E</u> A <u>S</u> <u>Q</u> <u>R</u> <u>K</u> ^a
p27-KID ^{A10}	H <u>A</u> <u>A</u> L <u>T</u> A <u>D</u> L <u>E</u> A <u>H</u> C <u>A</u> M <u>E</u> A <u>A</u> S <u>Q</u> <u>A</u> A ^b
p27-KID ^{A22}	<u>AAAAAAAAAAAAAAAAAAAAAAAAA</u> ^b
p27-KID ^{SL1}	H <u>L</u> <u>E</u> L <u>T</u> L <u>D</u> L <u>E</u> L <u>H</u> C <u>L</u> L <u>M</u> E <u>L</u> L <u>S</u> Q <u>R</u> K ^b
p27-KID ^{SL2}	H <u>E</u> E <u>L</u> T <u>L</u> D <u>L</u> E <u>L</u> H <u>C</u> L <u>L</u> M <u>E</u> L <u>L</u> S <u>Q</u> R <u>K</u> ^b
p27-KID ^{SL3} (p27-KID ^{+H})	H <u>E</u> E <u>L</u> T <u>K</u> D <u>L</u> E <u>L</u> H <u>C</u> L <u>L</u> M <u>E</u> L <u>L</u> S <u>Q</u> R <u>K</u> ^b
p27-KID ^{SL4}	H <u>E</u> E <u>L</u> T <u>K</u> D <u>L</u> E <u>L</u> H <u>C</u> K <u>L</u> M <u>E</u> E <u>L</u> S <u>Q</u> K ^b
p27-KID ^{SL5}	H <u>L</u> <u>E</u> L <u>T</u> L <u>D</u> L <u>E</u> E <u>H</u> C <u>L</u> K <u>M</u> E <u>L</u> E <u>S</u> Q <u>L</u> K ^b
p27-KID ^{SL6}	H <u>L</u> L <u>L</u> T <u>L</u> D <u>L</u> E <u>L</u> H <u>C</u> L <u>L</u> M <u>E</u> L <u>A</u> S <u>Q</u> L <u>L</u> ^b
p27-KID ^{-H}	H <u>E</u> E <u>L</u> T <u>E</u> D <u>L</u> E <u>E</u> H <u>C</u> E <u>E</u> M <u>E</u> E <u>E</u> S <u>Q</u> E <u>E</u> ^b
p27-KID ^{loop}	<u>T</u> N <u>P</u> G <u>N</u> G <u>G</u> H <u>G</u> G <u>T</u> G <u>V</u> G <u>G</u> N <u>G</u> G <u>N</u> G <u>H</u> G ^b

^a The residues determined to be suitable for mutagenesis are underlined and are in bold font in the linker domain sequence of the wild-type p27-KID (p27-KID^{wt}).

^b The mutated residues are underlined.

oligonucleotide design, were a 5' NdeI restriction site and a 3' XhoI restriction site for all the mutants. The mutagenesis cassettes were prepared by overlap extension polymerase chain reaction (oePCR). The overlap extension polymerase chain reaction is illustrated in Figure A.1. First, the single stranded non-overlapped ends of the long oligonucleotides are filled in by a proofreading polymerase (Expand high fidelity polymerase, Roche, Mannheim, Germany) to yield a double stranded DNA cassette. However, this double stranded DNA cassette product is in low quantities since only a small amount (10 ng) is added to the reaction. In the succeeding cycles, the end primers, which are present in large quantities in the reaction (125 ng), amplify the double stranded DNA cassettes. The details of the oePCR reaction that I performed including concentrations of reactants, and thermocycler settings are reported in Table A.2 and Table A.3. Figure 2.2 shows the product from the oePCR reaction for the preparation of the mutagenesis cassette for the p27-KID^H mutant. Similar product quantities were obtained for the other mutants.

2.3 Preparation of cassettes for cloning Cip/Kip linker domain peptides

Overlap extension PCR was also used in the preparation of oligonucleotide cassettes that were used to prepare expression plasmids for Cip/Kip linker domain peptides. For the preparation of each peptide cassette, two long oligonucleotides (98 bases each; 37 overlapping complementary bases) were designed and end primers were designed. Table A.4 shows the sequences of the oligonucleotides and end primers that were designed for each linker

Figure 2.2 Agarose gel analysis of the oePCR for the p27-KID^H mutant. The descriptions of the lanes are as follows: (a) PCR markers, (b) No primer control, (c) No enzyme control, (d) No template control, and (e) p27-KID^H oePCR. The expected size of the p27-KID^H oePCR product is 168 bp.

domain peptide. The cassette design for each peptide included (in the 5' to 3' direction with respect to the coding strand): (i) a 5' SacII restriction enzyme site, (ii) bases encoding a 6-histidine (His) tag, a thrombin cleavage site, the linker domain sequence, and a stop codon, and (iii) a 3' EcoRI restriction site. The Cip/Kip linker domain sequences designed included a Trp residue after the 22nd residue of the linker domain to facilitate protein concentration determination since these peptides otherwise lack chromophores that absorb light near 280 nm. It should be noted that there is a conserved Trp at the end of the Cip/Kip linker domain sequences (corresponding to Trp60 of p27). The cloning cassette for each peptide was produced in high yield using this procedure. The oePCR product for the p27 linker domain peptide is shown on Figure A.2. The concentrations of the reagents and the thermocycler settings were the same as in Section 2.2 above.

2.4 Preparation of *Escherichia coli* expression plasmids

The pET28a (+) plasmid (Novagen, San Diego, CA) was used as the starting vector to construct plasmids for p27 protein expression in *Escherichia coli* (*E.coli*). The multi-cloning site of the pET28 vector is shown on Figure 2.3. Three sets of plasmids were prepared.

The first set of plasmids prepared was used for cloning the Cip/Kip linker domain peptides. The starting plasmid was the pET28 vector. A cassette encoding a glutathione S-transferase tag with 5' NcoI site and 3' EcoRI restriction sites was cloned between the NcoI and EcoRI restriction sites. It should be

noted that there is a SacII site that was engineered into the cassette between the end of the GST sequence and the EcoRI site. The cassettes for the Cip/Kip linker domain peptides prepared in Section 2.3 were digested with SacII and NdeI and cloned into the vector. As mentioned in Section 2.3 above, the Cip/Kip linker domain peptides have a His tag at their 5' ends after the SacII restriction site. The plasmids formed after ligation of the Cip/Kip linker domain cassettes into the pET28-GST-His vector therefore have both GST and His tags. This vector is hereafter referred to as the pET28-GST-His vector. Cip/Kip linker domain peptides with both GST and His tags have two advantages. First, it provides more purification options. In addition, the use of a GST-His tag makes the fusion protein large enough to be visualized using sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE). Without the large GST tag, the small peptides (about 3.3 kDa each) would easily diffuse out of standard concentration polyacrylamide gels. Figure 2.4 shows a diagram of the pET28-GST-His vector used to express the p27 linker domain peptide.

The second set of plasmids prepared was used to express the p27-KID mutants. A pET28 plasmid with p27-KID cloned between the NdeI and EcoRI sites had previously been prepared in the laboratory.¹⁹⁷ The sequence of the p27-KID coding region is shown on Figure A.3. The XhoI site that follows the EcoRI site in the pET28 vector was mutated to prevent restriction by the XhoI enzyme. Within the p27 Cdk binding domain (D2) sequence, the bases encoding Leu70 and Glu71 were mutated from CTAGAG to CTGGAG (the XhoI site). The cassettes of the p27-KID mutants prepared in Section 2.2 were then digested

Figure 2.4 The pET28-GST-His p27-LH vector construct.

with NdeI and XhoI restriction enzymes and ligated into the pET28-p27-KID vector with the engineered XhoI site. Some of the cassettes for the p27-KID mutants were ligated into the pET28-GST-His vector. A cassette coding for p27-KID was amplified from the pET28p27-KID vector by PCR using the following primers: 5' GCGGCCTGGTGCCGCGCGGCAGCCAT 3' and 5' CTTGTCGACG-GAGCTCGAATTCGGA 3'. The cassette and the pET28-GST-His-p27-LH plasmid were both digested using NdeI and EcoRI enzymes followed by ligating the p27-KID cassette into the digested pET28-GST-His vector. This plasmid was termed pET28-GST-Hisp27-KID. The cassettes for the p27-KID mutants were ligated into the p27-KID coding region of the pET28-GST-Hisp27-KID by cassette mutagenesis in the same way that they were cloned into the pET28p27-KID plasmid. Table A.5 shows the p27-KID mutants and the plasmids that they were cloned into. Figure 2.5 shows a schematic diagram of the cassette mutagenesis design.

The last set of plasmids prepared was used for ligating mutants into a pET28vector with the full-length p27 (flp27) cloned between the NdeI and EcoRI sites. The sequence of the flp27 coding region of this vector is shown in Figure A.4. The DNA sequence encoding flp27 has a PstI restriction site (CTGCAG) within the bases coding for Ala28, Cys29, and Arg30 (GCCTGCAGG). These residues are located in the cyclin binding domain (D1) of p27. In addition, bases coding for the residues Pro92 and Arg93 constitute a SacII restriction enzyme recognition site (CCGCGG). These residues lie within the Cdk binding domain of p27. The pET28a (+) plasmid does not have PstI or SacII restriction sites. The

Cassette preparation

Preparation of the pET28-GST-His vector

Figure 2.5 Schematic diagram of the cassette mutagenesis design. LH* is a mutated cassette of a p27-KID linker domain variant. The same approach was used to clone the mutant cassettes into the pET28p27-KID vector.

following primers were used to amplify cassettes from the plasmids in which the p27-KID linker domain mutants had been cloned: 5' GCCCTCGGCCTGCAGGA-ACCTCTTCGGCCCGG 3' and 5' CAGGCACCTTTGGGGGGCCGCGGGGGTC 3'. The pET28p27-KID^{SL3} (pET28p27-KID^{+H}), pET28-GST-Hisp27-KID^{-H}, and the pETGST-Hisp27-KID^{loop} plasmids used as templates. The cassettes and the pET28flp27 vector were digested with PstI and SacII enzymes followed by cloning the cassettes into the digested vector.

2.5 Preparation of retroviral shuttle vectors

The MSCV-I-GFP vector was the starting plasmid. A vector map of the MSCV-I-GFP vector is shown on Figure A.5. Cassettes were amplified from the pET28flp27^{wt}, pET28flp27^{SL3} (pET28flp27^{+H}), pET28flp27^{-H}, and the pET28flp27^{loop} plasmids prepared in Section 2.4 above using the following primers: 5' GGACCCGAATTCATGTCAAACGTGCGAGTGTCTAACGGGAGCC-CTAG 3' and 5' CAGACCGAATTCTTATTACGCATAATCCGGCACATCATACG-GATACGTTTGACGTCTTCTGAGGCCAGGCTTCTTGGGC 3'. The primers were designed such that the amplified cassettes had EcoRI sites at both the 5' and 3' ends. The cassettes were digested with EcoRI and ligated into the MSCV-I-GFP vector. The plasmids were purified using an endonuclease free plasmid kit (Qiagen, Valencia, CA) and the sequences of the plasmid constructs were confirmed by DNA sequencing. The sequences of the MSCV-I-GFPflp27^{wt}, MSCV-I-GFPflp27^{SL3} (MSCV-I-GFPflp27^{+H}), MSCV-I-GFPflp27^{-H}, and the MSCV-I-GFPflp27^{loop} are shown on Table A.6.

2.6 Expression and purification of Cip/Kip linker domain peptides

The linker domain peptides were expressed as fusion proteins with amino-terminal glutathione S-transferase and histidine tags (GST-His). The fusion proteins were expressed in *E. coli* BL21 (DE3) cells (Novagen, San Diego, CA) at 37 °C in LB media and induced with 1 mM Isopropyl β -D-1-thiogalactopyranoside (IPTG) when the culture reached mid-log phase (absorbance at 600 nm (A_{600}) $\approx$ 0.8). Cells were allowed to grow for 2 hours following induction. Cells were harvested by centrifugation at 5,000 x *g* for 10 minutes at 4 °C using a Sorvall Evolution RC centrifuge (Mandel Scientific, Ontario, Canada). The harvested cells were then resuspended in a nickel chelating column loading buffer (Ni-A buffer; 20 mM Tris pH 8.0, 500 mM NaCl, 5 mM imidazole). Every gram of cell pellet was resuspended in 10 ml of Ni-A buffer. The resuspended cells were lysed by sonication using a Branson sonifier 450 (Branson, Danbury, CT). The cell lysate was cleared of cell debris by centrifugation at 20,000 x *g* for 30 minutes at 4 °C followed by filtration using a 0.45 μ m Millex syringe filter (Millipore, Bedford, MA). Unless stated otherwise, this procedure was used to prepare the soluble extract after protein over-expression in *E. coli*. The fusion proteins were then purified from the soluble cell extract using Ni²⁺-affinity chromatography using a Hi-Trap nickel chelating column (GE Healthcare, Piscataway, NJ). With the exception of the resin used in the high performance liquid chromatography columns, all chromatography resins used for protein purification were obtained from GE Healthcare, Piscataway, NJ. The soluble extract was loaded onto the nickel chelating column followed by a two

step washing procedure. In the first step, the column was washed with five column volumes of Ni-A buffer. This was followed by a second wash step in which the column was washed with five column volumes of a buffer containing 90% Ni-A buffer and 10% of nickel column elution buffer (Ni-B; 20 mM Tris pH 8.0, 500 mM NaCl, 500 mM imidazole). The fusion protein was eluted from the column by washing with a buffer containing 30% Ni-A and 70% Ni-B. The eluted protein fraction was dialyzed against GST-A buffer (140 mM NaCl, 2.7 mM KCl, 10 mM Na₂HPO₄, 1.8 mM KH₂PO₄, 5 mM dithiothreitol, pH 7.3) overnight at 4 °C. This was followed by glutathione-affinity chromatography. The protein was loaded onto a glutathione affinity column, washed with five column volumes of GST-A buffer and eluted from the column by washing with five column volumes of GST elution buffer (GST-B buffer; 50 mM Tris pH 8.0, 10 mM reduced glutathione, 5 mM dithiothreitol). The fraction eluted from the glutathione affinity column was dialyzed against the high performance liquid chromatography sample buffer (Buffer D; 10 mM Tris pH 7.2, 5 mM dithiothreitol). This was followed by thrombin cleavage at 4°C. Thrombin (No vagen, San Diego, CA) was added to the cleavage reaction at a concentration of 0.5 units per milligram of fusion protein. The cleaved protein was then purified using high performance liquid chromatography (HPLC). The cleaved protein was mixed in a 1:1 ratio (volume/volume) with the HPLC loading buffer (HPLC-A; 0.1% trifluoroacetic acid in water) and loaded onto a C18 column (Vydac, Hesperia, CA). The protein was eluted from the column by increasing the concentration of the HPLC elution buffer (HPLC-B; 0.1% trifluoroacetic acid in acetonitrile) at a rate of 1% per

minute up to a concentration of 80% HPLC-B. The HPLC silica-based column used here requires use of buffers with salt concentrations lower than 150 mM and pH lower than 7.5. Higher salt and pH conditions are detrimental to the performance and longevity of the column. The fractions were then lyophilized and analyzed by mass spectrometry. Figure 2.6 shows the purification of the p27 linker domain peptide. Similar results were achieved for the p21 and p57 linker domain peptides. The mass spectrometry analyses of the linker domain peptides are shown on Figure 2.7 and Figure 2.8.

2.7 Expression and purification of p27-KID variants

The p27-KID linker domain mutants listed in Table A.5 were expressed in *E.coli* as described in Section 2.6. Like the linker domain peptides, the bacterial cells that contained the expressed protein were resuspended in Ni-A buffer. After cell lysis and clearing, some of the mutants were found to be insoluble. Table A.6 shows all of the p27-KID mutants expressed, their solubilities, and the corresponding expression plasmid(s). Only the mutants with soluble extracts were purified. The soluble extract was loaded onto a Hi-Trap nickel column, followed by washing using five column volumes of Ni-A buffer. Protein was eluted from the column by washing with 100% Ni-B buffer. The eluted protein was dialyzed overnight against thrombin cleavage buffer (Buffer T; 20 mM Tris pH 8.0, 50 mM NaCl, 5 mM dithiothreitol) at 4 °C. The cleaved protein was boiled for 20 minutes and centrifuged at 30, 000 g for 30 minutes at 4 °C. The pellet mainly contains heat-denatured *E. coli* proteins and the supernatant

Figure 2.6 Purification of the p27 linker domain peptide (p27-LH).

Figure 2.7 Mass spectrometry analyses of p27 and p21 linker domain peptides.

Figure 2.8 Mass spectrometry analysis of p57 linker domain peptide.

contains the over-expressed mutant p27 protein. Figure 2.9 shows the purification of the p27-KID^{-H} mutant. Similar results were achieved in the expression of the other p27-KID linker domain mutants. After purification, the secondary structures of the more and less helical p27-KID mutants were determined by circular dichroism spectropolarimetry. For the more helical mutant the p27-KID^{SL3} was found to exhibit the desired secondary structural properties and was chosen for further study. Hereafter, this mutant is referred to as the p27-KID^{+H}. The p27-KID^{-H} and p27-KID^{loop} mutants were also found to have the desired structural properties. The p27-KID linker domain mutants that I studied in detail include: p27-KID^{+H}, p27-KID^{-H}, p27-KID^{loop}, p27-KID^{p21helix}, p27-KID^{p57helix}, and p27-KID^{ΔC}. These mutants were characterized by mass spectrometry (Figure 2.10, Figure 2.11, and Figure 2.12) and sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGE). Figure 2.13 shows the SDS-PAGE gels of the purified proteins.

The D1 peptide (residues 23-57) was prepared by solid phase peptide synthesis (Hartwell Center, St. Jude Children's Research Hospital, Memphis, TN). A pET28a plasmid encoding a peptide corresponding to the Cdk binding domain of p27 (D2; residues 58-105) was used to express a His-tagged D2 fusion protein. The D2 peptide was purified using the same protocol as used for the KID variants described above (the nickel buffers used in the purification of D2 were phosphate based, D2-Ni-A and D2-Ni-B, as specified in Section A.1). After purification, the D2 peptide was analyzed by mass spectrometry (Figure 2.14).

Prior to isothermal titration calorimetry (ITC) experiments, preparation of

1. Nickel affinity purification

2. Thrombin cleavage

3. Boiling

Figure 2.9 Purification of the p27-KID^H mutant.

Figure 2.10 Mass spectrometry analyses of p27-KID^{wt} and p27-KID^{p21helix}.

Figure 2.11 Mass spectrometry analyses of p27-KID^{p57helix} and p27-KID^{ΔC}.

Figure 2.12 Mass spectrometry analyses of purified p27-KID^{+H}, p27-KID^H, and p27-KID^{loop}.

Figure 2.13 SDS-PAGE analysis of purified p27-KID mutants. Note that the lanes in the upper panel were digitally captured from multiple, individual SDS-PAGE gels. However, each of these gels was electrophoresed for a similar period of time and, thus, this composite image accurately represents the relative mobility of the molecular weight markers and the p27-KID proteins.

Figure 2.14 Mass spectrometry analysis of the D2 peptide.

ternary complexes with Cdk2/cyclin A for thermal denaturation experiments or kinase assays, p27-KID^{wt} and its variants were treated with 100 mM dithiothreitol (DTT) at room temperature for 30 minutes followed by gel filtration chromatography purification using an Superdex S-75 column equilibrated with buffer GF (20 mM HEPES pH 7.5, 300 mM NaCl, 5 mM DTT). Unless stated otherwise, all gel filtration chromatography purifications were performed in columns equilibrated with buffer GF. The D2 peptide was also treated in the same manner as the p27-KID mutants before ITC experiments and kinase assays.

2.8 Expression and purification of cyclin A and phosphorylated Cdk2

A pET28 plasmid encoding human cyclin A (residues 173-432) had previously been prepared in the lab.¹⁹⁷ Cyclin A was expressed in *E.coli* at 37 °C in LB media grown to early log phase ($A_{600} \approx 0.4$). The temperature was reduced to 20 °C and protein expression was induced with 1 mM IPTG when the cell density reached an $A_{600} \approx 0.8$. The cells were then cultured for 20 hours and harvested by centrifugation. The protein was purified by nickel affinity chromatography (as described in Section 2.7; 50% Ni-B was used for elution) followed by ammonium sulfate ((NH₄)₂SO₄) precipitation. Maximal precipitation of the cyclin A protein during the (NH₄)₂SO₄ precipitation were achieved by adding finely ground (NH₄)₂SO₄ (0.56 g of (NH₄)₂SO₄ for every ml of protein solution; 85 % saturation) to protein solutions with a total protein concentration greater than 3 mg/ml. The precipitated protein was resuspended in TcycA –buffer

(20 mM Tris pH 8.0, 1.0 M NaCl, 2.5 mM CaCl₂) and cleaved using thrombin overnight at 4 °C. The cleaved cyclin A protein was further purified by anion exchange chromatography using Q-sepharose resin and stored at -20 °C after a second (NH₄)₂SO₄ precipitation step. Before use in experiments, the stored protein was purified by gel filtration chromatography using a superdex S-75 column.

A bicistronic plasmid vector encoding full length human Cdk2 and civ1 (a yeast CAK)²¹⁶ was kindly provided by Dr. Jane Endicott, Oxford University, Oxford, UK. Both the Cdk2 and civ1 proteins are fused to GST tags at their N-termini. civ1 is a Cdk-activating kinase that phosphorylates Cdk2 on threonine-160.²¹⁷ This vector was used to express P-Cdk2, full length human Cdk2 phosphorylated at threonine-160. P-Cdk2 expression was performed using the same procedure as discussed earlier for truncated human cyclin A. The protein was first purified by glutathione-affinity chromatography followed by dialysis into Pre-Scission cleavage buffer (50 mM Tris pH 7.0, 150 mM NaCl, 1 mM EDTA, 1 mM DTT). The purified protein was cleaved overnight using Pre-Scission protease (GE Healthcare, Piscataway, NJ) enzyme at 4°C. Only the GST-P-Cdk2 fusion protein contains the Pre-Scission cleavage site and *ipso facto* only the P-Cdk2 protein is cleaved from the GST fusion protein. Cleaved P-Cdk2 was purified using a second glutathione-affinity chromatography step. GST-civ1, the cleaved GST tag from GST-P-Cdk2 fusion protein, uncleaved GST-P-Cdk2, and pre-scission enzyme (Pre-Scission is GST tagged) were retained on the column while cleaved P-Cdk2 flowed through the column.

P-Cdk2 was then concentrated by ammonium sulfate precipitation (in the same way as cyclin A above) and stored at -20 °C. Before use in experiments, the stored P-Cdk2 protein was purified by gel filtration chromatography superdex S-75 column.

To prepare the P-Cdk2/cyclin A binary complex, the ammonium sulfate precipitated P-Cdk2 and cyclin A stored at -20 °C were first separately dissolved in buffer GF and the concentration of each protein was determined. The two proteins were then mixed in a 1:1 mole ratio (purified cyclin A to purified P-Cdk2) and incubated at 4 °C for 30 minutes. The binary complex was then purified by gel filtration chromatography using a Superdex 75 (S-75) column. The p27-KID/P-Cdk2/cyclin A ternary complexes used in the thermal denaturation experiments were made by mixing p27-KID^{wt} or mutants in a 1.1:1.0 mole ratio with the P-Cdk2/cyclin A complex, incubating the mixture for 30 minutes at 4 °C and purifying the complex using gel filtration chromatography using a Superdex 200 (S-200) column equilibrated in the thermal denaturation buffer (1 mM phosphate pH 7.0, 25 mM NaCl, 1 mM DTT).

2.9 Preparation of retrovirus

Retroviruses integrate their genetic material into the genomes of their hosts and use the transcription and translation machinery of the host to express viral proteins.²¹⁸ The viral genetic material is replicated by the host cell and new viruses are packaged. The packaged viruses exit the cell and invade the neighboring cells. The viral genome has four DNA modules that are

indispensable in invasion and replication: (i) a packaging signal (Ψ) that enables packaging of the viral genome into viral particles; (ii) the *env* gene that encodes the viral coat proteins used in the attachment of a virus to a host; (iii) the *pol* gene that encodes the viral polymerase; and (iv) the *gag* gene that encodes proteins that form the viral core structure.^{219,220} Viral particles are only packaged when the Ψ packaging signal is in *cis* with the *gag*, *pol* and *env* genes.²²¹

Retroviral expression systems were designed to take advantage of the modularity of the viral genome.²²²⁻²²⁴ First, a gene of interest (in this case *flp27^{wt}* or its variants) in *cis* with the retroviral packaging signal is ligated into shuttle vector. The *gag*, *pol*, and *env* genes were ligated into one plasmid or two separate plasmids called helper plasmid(s). The helper plasmid(s) and the shuttle vector are then transfected into a packaging cell line.²²⁵ The packaging cell lines have very high transfection efficiencies and are engineered to optimally and stably express the viral *gag*, *pol*, and *env* genes thereby ensuring production of high viral titres. Further, the packaging cell lines are genetically modified to prevent production of replication-competent viruses.^{226,227} In the packaging cell line, the *gag*, *pol* and *env* genes cloned into the helper plasmid(s) facilitate the packaging of the gene of interest into viral particles in *trans*. Only the gene of interest is packaged since it is in *cis* with the Ψ . The viral particles with the gene of interest are then used to transduce a non-packaging cell line. Since there are no *gag*, *pol* or *env* genes in the non-packaging cell line, the viruses with the gene of interest can only transduce the cells but cannot replicate. For my studies, the MSCV-I-GFP shuttle vectors in which the *flp27^{wt}* or its variants were cloned

(described in Section 2.5) and a helper plasmid were used to prepare retroviruses.

First, the the viruses were packaged in the 293T packaging cell line. 100 mm x 10 mm plates of 293T cells were cultured at 37°C in Dulbecco's Modified Eagle Medium (DMEM) supplemented with 10% fetal bovine serum (FBS), 2% of 200 mM glutamine (vol:vol), and 1% of 10,000 µg/µl penicillin/streptomycin (vol:vol). Hereafter, the DMEM supplements are referred to as additives. 3×10^6 cells were cultured on each plate. After 24 hours, the cell culture media (with additives) was substituted with additive free media (DMEM only) and incubated for 3 hours. During this time, transfection cocktails were prepared.

The cocktails contained a single helper DNA (containing the viral gag-pol-env genes), shuttle vector DNA plasmid, fugene 6 transfection reagent (Roche, Basel, Switzerland) and additive free DMEM media. The helper DNA was obtained from Dr. Martine Roussel (St. Jude Children's Research Hospital, Memphis, TN). In every cocktail, the ratio of the volume of the fugene reagent to the amount of the shuttle vector (by mass) was varied to identify the ratio that yielded the highest viral titre. The volume of the fugene reagent was kept constant at 15 µl and the amount of the shuttle vector added was varied to give different fugene to shuttle vector ratios. Cocktails with volume of fugene to mass of shuttle plasmid DNA ratios of 1:20, 1:15, 1:12, and 1:10 were prepared for each construct (each construct had 4 cocktails). A constant 6 µg of helper DNA was added to each cocktail. Serum free media was added to bring the total volume of each cocktail to 700 µl. Before addition of fugene to the cocktails, it

was first warmed to room temperature according to manufacturer's instructions. For each cocktail, the reagents were added in the following order: (i) serum free media, (ii) helper DNA, (iii) vector DNA, and (iv) fugene. The cocktails were then mixed. After 3 hours, the transfection cocktails were added to the incubated cells.

After 20 hours the additive free media was replaced with 10 ml of fresh media supplemented with additives. In the following steps, all the media was supplemented with additives unless stated otherwise. After 6 hours, the media was aspirated from the plates and 5 ml of media was added to each plate. After 18 hours, 5 ml of the media containing virus was harvested and 5 ml of fresh media was added to each plate. 8 hours later, 5 ml of media containing virus was again harvested and a fresh 7 ml of fresh media added to each plate. Virus-containing media (7 ml) was harvested a final time after an additional 20 hours. The harvested virus was filtered using a 0.22 μm steriflip filter (Millipore, Billerica, MA) and stored at 4 $^{\circ}\text{C}$.

2.10 Viral transduction and cell cycle arrest analysis

3×10^5 NIH 3T3 cells were cultured on 100 x 10 mm plates in DMEM media with additives (additives specified in section 2.9 above). After 20 hours, the media was aspirated from each plate. 5 ml of cold virus (virus kept in ice) supplemented with 30 μl of 1 mg/ml of polybrene (6 μl of polybrene for every ml of virus) was added to each plate and incubated at 37 $^{\circ}\text{C}$. After 3 hours, 10 ml of DMEM media with additives was added to each plate. After 48 hours, the cells

were washed, re-suspended in additive-free DMEM media, and the viral titre determined using GFP fluorescence analysis (Flow Cytometry Facility, St. Jude Children's Research Hospital, Memphis, TN). Before GFP analysis, viable cells were separated from non-viable cells by propidium iodide (PI) staining. For PI staining, the cells were first mixed until a single suspension of cells was realized. The cells were then counted and washed once with phosphate buffered saline (PBS). The supernatant was decanted and the cells were resuspended in PI solution (0.05 mg/ml propidium iodide) at 1×10^6 cells/ml. This was followed by treatment with RNase for 30 minutes at room temperature. Prior to and after the RNase step, the cells were kept in ice. The cells were then filtered using a 40 μ m-diameter mesh (Small Parts Inc., Miami Lakes, FL) to eliminate large multicellular clumps that might clog the flow cytometer. The filtered cells were analyzed by flow cytometry.

For cell cycle analysis, viral solutions with high viral titres, as determined by GFP analysis, were used to transduce NIH 3T3 cells as described above. The cells were washed, and analyzed for viral transduction using GFP analysis. The top 10% GFP positive cells were stained by PI and analyzed for DNA content. Each cell cycle arrest analysis was performed in triplicate for each construct.

2.11 Protein concentration determination

The concentrations of the prepared proteins were determined by UV absorbance at 280 nm in a denaturing buffer containing 20 mM phosphate pH 6.5, and 6.0 M guanidinium hydrochloride.^{228,229} The molar extinction coefficients

were determined using the ProteinParameters tool (ExPASy server; <http://us.expasy.org/tools/protparam.html>) as: 15 220 M⁻¹ cm⁻¹ for p27-KID^{wt}, p27-KID^{p57helix}, p27-KID^{ΔC}, p27-KID^{+H}, p27-KID^{-H} and p27-KID^{loop}; 15 470 M⁻¹ cm⁻¹ for D2 and p27-KID^{p21helix}; and 35 560 M⁻¹ cm⁻¹, 31 860 M⁻¹ cm⁻¹ and 33 710 M⁻¹ cm⁻¹ for P-Cdk2, cyclin A, and the P-Cdk2/cyclin A complex respectively. The extinction co-efficient for the p27-KID^{wt}, p27-KID^{p57helix}, p27-KID^{ΔC}, p27-KID^{+H}, p27-KID^{-H} and p27-KID^{loop} ternary complexes was determined as 24 465 M⁻¹ cm⁻¹. An extinction coefficient value of 24 590 M⁻¹ cm⁻¹ was used for the p27-KID^{p21helix} ternary complex. The p27, p21, and p57 linker domain peptides all had an extinction coefficient of 5690 M⁻¹ cm⁻¹.

2.12 Secondary structure analysis by circular dichroism (CD) spectropolarimetry

The secondary structures of the p27-KID and its mutants were analyzed using an AVIV model 62A DS CD spectropolarimeter. All the samples were analyzed in a buffer containing 10 mM phosphate pH 7.0 and 5 mM DTT. The experiments were performed at 25 °C. At least 3 CD traces were averaged for each sample with a 5 second signal averaging time for each data point. To determine the helical contents of the p27-KID mutants relative to the wild-type, the CD spectra of all the proteins to be compared were obtained under exactly the same conditions. For these experiments, the protein concentration in each sample was 20 μM. The CD spectra were collected in a 1 mm path length CD cuvette to maximize the signal to noise ratio.

2.13 Thermal denaturation experiments

The thermal denaturation of the ternary complexes formed by the p27-KID^{wt} and its mutants on binding the P-Cdk2/cyclin A complex was monitored in the temperature range of 15 °C to 95 °C using an AVIV model 62A DS circular dichroism spectropolarimeter equipped with a Peltier thermoelectric temperature control device. The samples, each containing 400 nM of protein in 1 mM sodium phosphate buffer, pH 7.0, 25 mM NaCl and 1 mM DTT, were analyzed in a 1 cm path length quartz cuvette. For these studies, denaturation was monitored by measuring ellipticity at 222 nm at 1 °C intervals at a heating rate of 5 °C min⁻¹ with constant stirring. Three experiments were performed with ternary complexes comprised of P-Cdk2/cyclin A and p27-KID^{wt}, p27-KID^{+H}, p27-KID^{-H}, p27-KID^{loop}, or p27-KID^{p57helix}, and two experiments were performed with ternary complexes containing p27-KID^{p21helix} or p27-KID^{ΔC}. The denaturation curves were analyzed as previously described by Bowman, *et al.*,²¹⁶ as follows: The fraction of unfolded protein (f_D) at any given temperature was calculated from the corresponding experimental ellipticity value θ using Equation 2.1:

$$f_D = [\theta - (\theta_N + m_N T)] / [(\theta_D + m_D T) - (\theta_N + m_N T)] \quad (\text{Eq. 2.1})$$

where T is the temperature, θ_N and θ_D are the ellipticity values corresponding to the native (N) and denatured (D) states extrapolated to $T = 0$, and m_N and m_D are the slopes of the baselines preceding and following the transition region obtained

by linear regression analysis. The free energy of unfolding, ΔG_{D-N} , was calculated using Equation 2.2:

$$\Delta G_{D-N} = -RT \ln K = -RT \ln \{ f_D / (1 - f_D) \} \quad (\text{Eq. 2.2})$$

to calculate T_m , ΔG_{D-N} values over the transition region of the denaturation curve were plotted against temperature and fit to a straight line by linear regression. T_m corresponds to the intercept of the x-axis where ΔG_{D-N} is equal to zero (this is the temperature at which $f_D = f_N$). The melting temperature for each sample was determined at least two times. The experimental error in the determination of the melting temperature was calculated by determining the standard deviation from the mean T_m for each sample.

2.14 Determination of thermodynamic parameters by ITC

The thermodynamic parameters associated with p27-KID^{wt} and the mutant proteins binding to P-Cdk2/cyclin A were determined using isothermal titration calorimetry (ITC). The experiments were performed using a VP-ITC calorimeter (Microcal, Northampton, MA). A major advantage of ITC is that the enthalpy of association (ΔH), binding stoichiometry (n), and binding affinity (K_a ; within certain limits) of the two interacting components can be determined in a single experiment. It is important to note that ITC cannot accurately determine the binding affinity associated with very tight interactions (low nM binding affinity) because such systems have a very high c value – c is defined by Wiseman, *et*

*al.*²³⁰ However, accurate estimates can be obtained. The Gibbs free energy of association (ΔG) is calculated using the equation $\Delta G = -RT \ln(1/K_a)$, where R is the gas constant ($1.987 \text{ cal mol}^{-1} \text{ K}^{-1}$) and T is the temperature (degrees Kelvin) at which the experiment was performed. Given that the above limitation applies, the values of the Gibbs free energy of association of p27-KID^{wt}, p27-KID^{+H}, p27-KID^{-H}, p27-KID^{p57helix}, p27-KID^{p21helix} and p27-KID ^{ΔC} binding to P-Cdk2/cyclin A are taken as estimates of the upper limit of the K_d and those of p27-KID^{loop}, D1, and D2 are considered to be accurate.

The experiments were performed at 25 °C in buffer containing 20 mM HEPES pH 7.5, 300 mM NaCl and 5 mM DTT. For these experiments, the syringe volume was 250 μL and the cell volume was 1.3 ml. In each experiment, the P-Cdk2/cyclin A complex was in the cell and p27-KID^{wt} or its variants were in the syringe (titrant). A standard injection protocol was used for each titrant. 2 μL of the titrant was first injected followed by 39 injections, 6 μL each, with 300 seconds between the injections. The following were the protein concentrations used in the ITC studies: 75 μM p27-KID^{wt} + 7.4 μM P-Cdk2/cyclin A; 73 μM D1 + 5.2 μM P-Cdk2/cyclin A; 72.3 μM D2 + 5.61 μM P-Cdk2/cyclin A; 56 μM p27-KID^{+H} + 4.2 μM P-Cdk2/cyclin A; 42 μM p27-KID^{-H} + 4.4 μM P-Cdk2/cyclin A; and 36.4 μM p27-KID^{loop} + 3.1 μM P-Cdk2/cyclin A; 56.87 μM p27-KID^{p21helix} + 6.22 μM P-Cdk2/cyclin A; 53.57 μM p27-KID^{p57 helix} + 4.6 μM P-Cdk2/cyclin A; and 35 μM p27-KID ^{ΔC} + 2.88 μM P-Cdk2/cyclin A. The samples were degassed before the experiments were performed and the instrument was regularly calibrated as specified by the manufacturer. The binding data were fit to a 1:1

binding model using the Origin software (Origin Lab Corporation, Northampton, MA). This analysis yielded values of the enthalpy of binding (ΔH), the equilibrium association constant (K_a), and the Gibbs free energy of binding (ΔG). The product of entropy and temperature ($-T\Delta S$) was calculated using Equation 2.3:

$$-T\Delta S = \Delta G - \Delta H \quad (\text{Eq. 2.3})$$

2.15 Kinase inhibition assays

A solution of 80 pM P-Cdk2/cyclin A binary complex containing 2.5 μM histone H1 (Upstate, Charlottesville, VA) and varying amounts of p27-KID^{wt} or its mutants was incubated at 4 °C for 1 hour. 6 μCi of $\gamma^{32}\text{P}$ -ATP (PerkinElmer, Inc., Wellesley, MA) was added to each reaction and incubated at 30 °C for 35 minutes. The concentration of the $\gamma^{32}\text{P}$ -ATP was 1 mCi/ml (1 part 10 mCi/ml of $\gamma^{32}\text{P}$ -ATP to 9 parts 111 μM cold ATP) and the total reaction volume was 15 μL . All the samples were in a kinase assay buffer containing 20 mM HEPES pH, 7.3, 25 mM sodium b-glycerolphosphate, 15 mM MgCl_2 , 16 mM EGTA, 0.5 mM Na_3VO_4 and 10 mM DTT. The kinase reactions were quenched by adding SDS-PAGE gel loading buffer. The samples were then analyzed by SDS-PAGE and the bands quantified using a phosphoimager (Molecular Dynamics, Sunnyvale, CA). The inhibition data was fit to a dose-response model using Graphpad Prism software (Graphpad Software, Inc., San Diego, CA). Each curve was fit from a triplicate set of experiments. The error was reported as the 95% confidence interval of the IC_{50} from a curve fit of a triplicate data set.

Chapter 3: The linker domains of proteins in the Cip/Kip family play similar structural roles

The structure of the linker domain of p21, p27, and p57 is conserved, and this domain serves to couple the D1 and D2 domains of Cip/Kip proteins. In the first study of this chapter, I have investigated the effects of deleting the linker domain on the function of p27 as a cyclin-dependent kinase inhibitor. The thermodynamics of association of peptides corresponding to the D1 and D2 domains of the KID with the P-Cdk2/cyclin A complex relative to the p27-KID^{wt} were analyzed. The functional consequences of deleting the linker domain were determined by studying inhibition of the P-Cdk2/cyclin A catalytic activity by D2 peptide.

Although the nascent helical structure of the linker domain is conserved amongst members of the Cip/Kip family, its sequence is not. However, the sequence of the linker domain for each Cip/Kip protein is highly conserved in different species suggesting that the sequences of these proteins may be important for their function. In the second study in this chapter, I investigated whether the linker domains of the Cip/Kip proteins, despite having different sequences, play similar structural roles. Chimeric Cip/Kip proteins were used for these studies. The chimeric proteins were prepared by replacing the linker domain of the p27-KID with either the p21 linker domain (p27-KID^{p21helix}) or the p57 linker domain (p27-KID^{p57helix}). The interaction of these chimeric proteins with the P-Cdk2/cyclin A kinase complex was then probed. This chimeric mutant

model is suitable for these studies for two reasons. First, the individual linker domains of p21, p27 and p57 have an inherent propensity to form α -helices indicating that this domain can fold independent of the rest of the KID polypeptide. Secondly, the residues neighboring the linker domains in the native proteins are well conserved. This minimizes the variation in stabilization of the linker helix by the neighboring residues. By replacing only the linker domain and keeping D1 and D2 constant, the consequences of linker domain swapping on the interaction with the P-Cdk2/cyclin A complex can be studied.

Only 3 out of the 22 residues in the linker domains of p21, p27, and p57 are conserved. The role of these conserved residues in the interaction of the Cip/Kip proteins with Cdk complexes is unknown. In the final study of this chapter, I have investigated the role of these conserved residues by preparing a p27-KID variant in which the three conserved residues in the linker domain are mutated to glycines (Glu39Gly/Leu41Gly/Arg43Gly, the p27-KID^{ΔC} mutant) and probing its interaction with the P-Cdk2/cyclin A complex.

In all the studies, CD was used to study the secondary structures of the mutants. The relative stabilities of the ternary complexes formed by the variants on binding to P-Cdk2/cyclin A were determined by monitoring the ellipticity of each complex at 222 nm as a function of temperature. ITC was used to determine the thermodynamics of association of the mutants with the P-Cdk2/cyclin A complex. Kinase inhibition assays were performed to provide insights into the consequences of the mutations on the kinase inhibition activity of the Cip/Kip proteins.

3.1 The nascent helical character of the linker domain is preserved in the chimeric mutants and the p27-KID^{ΔC} mutant

The secondary structures of the p21, p27, and p57 linker domain peptides, and the p27-KID^{wt}, p27-KID^{p21helix}, p27-KID^{p57helix}, p27-KID^{ΔC}, were determined by CD (Figure 3.1 and Figure 3.2). The linker domain peptides are mainly unstructured (ellipticity at about 200 nm), with some helical secondary structure (ellipticity at 222 nm). These results are consistent with secondary structure prediction for the linker domains of p27, p21, and p57.¹⁹⁷ The p21 helix peptide shows a significantly higher content of random coil than either p27 or p57 linker helix peptides. This is not surprising since its sequence has a glycine (Gly40) which would effectively decrease its helical content and increase its overall random coil composition relative to the p27 and p57 helix peptides. The presence of nascent α -helical secondary structure in all the linker domain peptides indicates that their sequences have an intrinsic tendency to form α -helices and can fold independent of the rest of the KID sequence. The stabilization of the linker domains in the context of the KID should be very similar since the residues neighboring the linker domains in the p21, p57, and p27 proteins are well conserved.

The chimeric mutants have secondary structures very similar to that of the p27-KID^{wt}. They are largely unstructured with nascent helical characteristics. The mean residue molar ellipticity values of the chimeric mutants at 222 nm ($[\Theta_{222}]$) are comparable to that of the p27-KID^{wt} indicating that they have comparable helical contents. The results of the secondary structural analysis of

Figure 3.1 CD spectra of the p21, p27, and p57 linker domain peptides.

Figure 3.2 CD spectra of p27-KID^{wt}, p27-KID^{p21helix}, p27-KID^{p57helix}, and p27-KID^{ΔC}.

the chimeric mutants suggest that the linker domains of p21 and p57, despite having different sequences and being bordered by D1 and D2 domains of p27, adopt structures similar to their previously reported secondary structures in the context of p21-KID and p57-KID, respectively.^{188,198} Further, this shows that our chimeric mutant model does not cause major changes in the secondary structure and hence is suitable for probing the structure conservation/sequence divergence relationship of the linker domains of p27, p21, and p57.

The p27-KID^{ΔC} mutant also exhibits a largely disordered secondary structure with nascent helical characteristics (Figure 3.2). The $[\Theta_{222}]$ of this mutant is comparable to that of the wild-type p27-KID indicating that the two p27-KID variants have comparable helical contents. These results suggest that the mutation of the conserved residues in the linker domain to glycines does not cause a major structural change. This structural similarity suggests that any disparities in the interaction between p27-KID^{ΔC} and the P-Cdk2/cyclin A complex relative to the p27-KID^{wt} are not a consequence of perturbation of the structure of the linker domain.

3.2 Binding thermodynamics determined by ITC

The thermodynamics of the association of the p27-KID^{wt} and its variants with the P-Cdk2/cyclin A complex were characterized by ITC. First, the thermodynamic parameters of the binding of either the D1 or the D2 peptide to the binary complex were compared to those of p27-KID^{wt}. The binding of either D1 or D2 to P-Cdk2/cyclin A is much weaker than that of p27-KID^{wt}. The

dissociation constants for the p27-KID^{wt}, D1, and D2 were determined to be 3.34 ± 1.64 nM, 42.20 ± 5.16 nM, and 82.96 ± 17.89 nM, respectively (Table 3.1).

These results show that the coupling of D1 and D2 via the linker domain greatly increases the affinity of p27 for the kinase complex. The ΔG 's of association of p27-KID^{wt}, D1, and D2 with the P-Cdk2/cyclin A complex are -10.06 ± 0.07 kcal mol⁻¹ and -9.66 ± 0.12 kcal mol⁻¹, and -11.61 ± 0.30 kcal mol⁻¹, respectively. The sum of the ΔH values for the interactions of D1 and D2 with P-Cdk2/cyclin A is -47.69 kcal mol⁻¹ relative to the -43.77 kcal mol⁻¹ of heat released when p27-KID^{wt} binds to the same complex. This suggests that the P-Cdk2/cyclin A bound conformation(s) of either the D1 or D2, or both peptides, is slightly different from its/their conformation(s) in the context of p27-KID.

The chimeric mutants bound to the P-Cdk2/cyclin A complex with affinities very similar to that of p27-KID^{wt} (Table 3.1). Figure 3.3 shows the binding isotherms for the interactions between p27-KID^{wt} or its variants with the P-Cdk2/cyclin A complex. The dissociation constants for p27-KID^{p21helix} and p27-KID^{p57helix} were 2.83 ± 1.37 nM, and 0.49 ± 0.14 nM respectively. Similar to that for p27-KID^{wt}, the binding of each chimeric mutant to the P-Cdk2/cyclin A complex was associated with a large entropic penalty. $-T\Delta S$ values of $+32.16 \pm 1.22$ kcal mol⁻¹, $+16.24 \pm 1.29$ kcal mol⁻¹, and $+18.44 \pm 0.35$ kcal mol⁻¹ were obtained for the p27-KID^{wt}, p27-KID^{p21helix}, and p27-KID^{p57helix} respectively. However, the disparities $-T\Delta S$ values suggest that the chimeric mutants do not fold to the same extent as the p27-KID^{wt}. Although there are differences in the folding of the chimeric mutants relative the p27-KID^{wt} on binding the

Table 3.1. Thermodynamic parameters determined by ITC at 25 °C.

Interaction	K_d (nM)	ΔG^a (kcal mol ⁻¹)	ΔH (kcal mol ⁻¹)	$-T\Delta S^b$ (kcal mol ⁻¹)
D1 + P-Cdk2/cyclin A	42.20 ± 5.16	- 10.06 ± 0.07	- 16.14 ± 0.20	+ 6.08 ± 0.27
D2 + P-Cdk2/cyclin A	82.96 ± 17.89	- 9.66 ± 0.12	- 31.55 ± 1.66	+ 21.89 ± 1.53
p27-KID ^{wt} + P-Cdk2/cyclin A	3.34 ± 1.64	- 11.61 ± 0.30 ^c	- 43.77 ± 1.38	+ 32.16 ± 1.22
p27-KID ^{p21helix} + P-Cdk2/cyclin A	2.83 ± 1.37	- 11.70 ± 0.27 ^c	- 27.94 ± 1.26	+ 16.24 ± 1.29
p27-KID ^{p57helix} + P-Cdk2/cyclin A	0.49 ± 0.14	- 12.71 ± 0.17 ^c	- 31.15 ± 0.51	+ 18.44 ± 0.35
p27-KID ^{ΔC} + P-Cdk2/cyclin A	1.97 ± 0.65	- 11.89 ± 0.21 ^c	-39.65 ± 0.13	+ 27.76 ± 0.17

In all the experiments above, the stoichiometry of binding was determined as one (1), with experimental error within 10%. The errors reported above are the standard deviations from the mean of three independent experiments.

^a Calculated with the equation $\Delta G = -RT \ln(1/K_d)$.

^b Calculated with the equation $\Delta G = \Delta H - T\Delta S$.

^c The c values for these p27-KID variants are very high and the ΔG values reported here are taken as estimates of the upper limits of K_d . Ideal c values for accurate determination of K_d have a range of 1 to 500. The c values for the binding of p27-KID^{wt}, p27-KID^{p21helix}, p27-KID^{p57helix}, and the p27-KID^{ΔC} to P-Cdk2/cyclin A complex are 2212, 2195, 9387, and 1461, respectively.

Figure 3.3 ITC binding isotherms for the p27-KID variants.

P-Cdk2/cyclin A complex, data from the binding of D1 and D2 with the same complex suggest that the D1 and D2 domains of these mutants bind to the kinase complex. The chimeric mutants have much higher binding affinities for the P-Cdk2/cyclin A complex than either D1 or D2 (Table 3.1). This stems from the linkage by the linker domain. In addition, the association of the the chimeric mutants with P-Cdk2/cyclin A is more favorable than that of D1 or D2. Further, the Gibbs free energy of association for the chimeric mutants with P-Cdk2/cyclin A is more favorable than ΔG values previously reported for the interactions of p27-KID^{wt} with either Cdk2 or cyclin A.¹⁹⁷ The differences in the folding of the p27-KID^{wt} and the chimeric mutants are further seen in the enthalpy values obtained for their interaction with the P-Cdk2/cyclin A complex. Enthalpy values of $-43.77 \pm 1.38 \text{ kcal mol}^{-1}$, $-27.94 \pm 1.26 \text{ kcal mol}^{-1}$, and $-31.15 \pm 0.51 \text{ kcal mol}^{-1}$ were obtained for the p27-KID^{wt}, p27-KID^{p21helix}, and p27-KID^{p57helix} respectively. The differences in ΔH are most likely caused by the interactions of different sets of residues in the linker domains of p27-KID^{wt}, p27-KID^{p21helix}, and p27-KID^{p57helix} (recall that the sequences linker domains of the Cip/Kip proteins are not conserved). The enthalpy values show that the interactions of the chimeric mutants with the P-Cdk2/cyclin A complex have fewer favorable contacts with respect to the p27-KID^{wt}. Nevertheless, the enthalpy generated by each chimeric mutant compensates for the entropic cost associated with the folding on binding the P-Cdk2/cyclin A complex making the free energies of the interactions negative and favorable – entropy-enthalpy compensation. Entropy-enthalpy compensation is a phenomenon in which the enthalpy change in a system during

a chemical process (in which the experimental conditions are varied) is partially or completely compensated for by a corresponding change in entropy such that the net free energy change is very small. As an example, the decrease in the absolute magnitude of ΔH in the interactions of p27-KID^{p21helix} and p27-KID^{p57helix} with P-Cdk2/cyclin A relative to p27-KID^{wt} caused by the differences in the sequences of the linker domains (variation in experimental conditions) is compensated for by a proportionate decrease in the entropic cost of binding; the residues that make fewer favorable contacts (release less heat) have more degrees of freedom (increase in entropy). Entropy-enthalpy compensation therefore results in a narrow range of variation of ΔG between the native system and the system with varied experimental conditions. The ΔG values of the interactions of p27-KID^{wt}, p27-KID^{p21helix}, and p27-KID^{p57helix} with P-Cdk2/cyclin A vary within a narrow range indicating that, despite the enthalpic differences, the linker domains have very similar ΔG contributions to the interaction of the Cip/Kip proteins with the P-Cdk2/cyclin A complex. The free energies of association for the p27-KID^{wt}, p27-KID^{p21helix}, and p27-KID^{p57helix} were $-11.61 \pm 0.30 \text{ kcal mol}^{-1}$, $-11.70 \pm 0.27 \text{ kcal mol}^{-1}$, and $-12.71 \pm 0.17 \text{ kcal mol}^{-1}$ respectively.

The interaction of the p27-KID ^{ΔC} mutant with the P-Cdk2/cyclin A complex relative to the p27-KID^{wt} shows the thermodynamic contributions of the conserved residues within the linker domain. This mutant binds very tightly to the kinase complex with a K_d of $1.97 \pm 0.65 \text{ kcal mol}^{-1}$ (Figure 3.3, Table 3.1). Similar to the chimeric mutants, it also folds extensively on binding the

P-Cdk2/cyclin A complex and has a $-T\Delta S$ value of $+27.76 \pm 0.17 \text{ kcal mol}^{-1}$. This extensive folding on binding, as is expected, is accompanied by the release of a large amount of heat. The enthalpy of association of the p27-KID^{ΔC} mutant with P-Cdk2/cyclin A complex is $-39.65 \pm 0.13 \text{ kcal mol}^{-1}$. The enthalpic contribution of the conserved residues is about $4.12 \text{ kcal mol}^{-1}$ (about 9.4% of the total heat released). The net free energy obtained for the interaction of the p27-KID^{ΔC} with the P-Cdk2/cyclin A complex is $-11.89 \pm 0.21 \text{ kcal mol}^{-1}$. The ΔG of the interaction of the p27-KID^{ΔC} mutant to the P-Cdk2/cyclin A complex is therefore the same as that for the p27-KID^{wt} within the error range. This shows that even though there is a slight decrease in enthalpy of interaction of the p27-KID with the P-Cdk2/cyclin A complex in the absence of the conserved residues within the linker domain, this loss is completely compensated for by a proportionately reduced entropic cost – entropy-enthalpy compensation.

3.3 The p27-KID^{wt} ternary complex is more stable than the corresponding complexes of its variants

Previous studies have shown that the binding of the p27-KID to P-Cdk2/cyclin A stabilizes that binary kinase complex and increases its melting temperature (T_m).²¹⁶ While the P-Cdk2/cyclin A complex has a T_m of $54.7 \pm 0.6 \text{ }^\circ\text{C}$, the p27-KID/ P-Cdk2/cyclin A ternary complex has a T_m of $82.3 \pm 0.5 \text{ }^\circ\text{C}$, an increase of $27.6 \text{ }^\circ\text{C}$ in the melting temperature. I investigated the consequences of linker domain swapping (chimeric mutants), and the role of the

conserved residues within the linker domain on the stabilization of the binary complex by determining the T_m 's of their ternary complexes on binding the P-Cdk2/cyclin A complex. The melting temperature (T_m) for each protein complex was determined by using standard thermal denaturation curve-fitting equations²³¹ to analyze the temperature dependence of $[\Theta_{222}]$ using CD. The thermal denaturation curves for the ternary complexes of p27-KID^{wt}, p27-KID^{p21helix}, p27-KID^{p57helix}, and p27-KID ^{Δ C} are shown in Figure 3.4. It is significant to note that the thermal denaturation process for the complexes in this study is irreversible due to protein aggregation at temperatures above their T_m values. Our results show that the ternary complex containing p27-KID^{wt} ($T_m = 82.3 \pm 0.5$ °C) is slightly more stable than those containing p27-KID^{p21helix} or p27-KID^{p57helix} which have T_m values of 80.7 ± 0.2 °C and 80.1 ± 0.1 °C (Table 3.2), respectively. The p27-KID ^{Δ C} is the least stable of the three mutants relative to the p27-KID^{wt} and has a T_m of 74.1 ± 0.6 °C. The relative instability of the p27-KID ^{Δ C} is probably caused largely in part by the mutation of Leu41 which in the crystal structure of the p27-KID^{wt}/Cdk2/cyclin A complex¹⁰⁴ is buried in the p27-KID^{wt}/cyclin A binding interface. By mutating this residue to a glycine, I have decreased the buried surface area at this position and consequently diminished the extent of favorable inter-molecular interactions. The other conserved residues, Glu39 and Arg43, are solvent exposed and most likely do not contribute significantly to the interaction of the linker domain with the cyclin A. This result indicates that the conserved residues of the linker helix contribute significantly to the stability of the ternary complex (by about 8°C).

Figure 3.4 Thermal denaturation curves of the p27-KID mutants.

Table 3.2 Melting temperatures of ternary complexes formed by either p27-KID^{wt} or its variants bound to P-Cdk2/cyclin A complex.

Complex	T_m (°C)
p27-KID ^{wt} /P-Cdk2/cyclin A ^a	82.3 ± 0.5
p27-KID ^{p21helix} /P-Cdk2/cyclin A ^b	80.7 ± 0.2
p27-KID ^{p57helix} /P-Cdk2/cyclin A ^a	80.1 ± 0.1
p27-KID ^{ΔC} /P-Cdk2/cyclin A ^b	74.1 ± 0.6

^a The error reported is the standard deviation from the mean of three independent experiments.

^b The error reported is the standard deviation from the mean of two independent experiments.

3.4 Relative inhibition potency determined by *in vitro* kinase assays

Kinase inhibition assays were performed to probe the P-Cdk2/cyclin A kinase inhibition potency of the p27-KID mutants with respect to the p27-KID^{wt}. In these studies, the P-Cdk2/cyclin A kinase concentration was 80 pM to avoid depletion of the inhibitors which would be the case at much higher kinase concentrations. All the proteins used in these studies were exchanged into the kinase assay buffer (buffer KA) by gel filtration chromatography. This was necessary because these assays are very sensitive to differences in solution conditions. It was therefore very important to prepare all the samples under exactly the same buffer conditions in order to obtain comparable results for the various p27-KID variants.

Studies of the inhibition potency of D2, Figure 3.5, Table 3.3, shows that it is a poor inhibitor ($IC_{50} = 125$ nM) and cannot completely inhibit kinase activity. even at very high D2 inhibitor concentrations, as high as 17.7 μ M, P-Cdk2/cyclin A retains about 25% of full kinase activity. This indicates that the isolated D2 peptide does not fold into the same inhibitory conformation as does D2 in the context of the full KID.

On the other hand, the chimeric mutants are potent kinase inhibitors with IC_{50} values in the low nM range. Figure 3.6 shows the gels and kinase inhibition curves of the chimeric mutants. The IC_{50} values for the p27-KID^{wt}, p27-KID^{p21helix}, and p27-KID^{p57helix} were 0.22 nM, 1.73 nM, and 3.45 nM respectively (Table 3.3). These results clearly show that the D2 domains of the chimeric mutants are folded into conformations with a capacity of inhibiting the kinase

Figure 3.5 Inhibition of P-Cdk2/cyclin A kinase activity by the D2 peptide. (a) SDS-PAGE gels showing the inhibition of the P-Cdk2/cyclin A kinase by the p27-KID^{wt} and D2. PO₄-H1 is phosphorylated histone H1. (b) Inhibition curves for the p27-KID^{wt} and D2 peptide.

Table 3.3 Inhibition potencies of the p27-KID variants.

Variant	IC₅₀ (nM)	95% confidence intervals of IC₅₀ (nM)^a
p27-KID ^{wt}	0.22	0.17 – 0.28
p27-KID ^{p21helix}	1.73	1.09 – 2.73
p27-KID ^{p57helix}	3.45	2.32 – 5.13
p27-KID ^{ΔC}	2.56	1.70 – 3.87
D2	125 ^b	45.94 – 340.80

^a Error is reported as the 95% confidence interval of the IC₅₀ from a curve fit of a triplicate data set.

^b This variant only partially inhibits kinase activity. 25% residual kinase activity observed at inhibitor concentrations as high as 17.7 μM.

(a)

(b)

Figure 3.6 Inhibition of P-Cdk2/cyclin A kinase activity by the p27-KID variants. (a) SDS-PAGE gels showing the inhibition of the P-Cdk2/cyclin A kinase by the p27-KID mutants. PO_4 -H1 is phosphorylated histone H1 (b) Inhibition curves of the p27-KID mutants.

complex. Even though the chimeric mutants are potent kinase inhibitors, their potencies are slightly decreased relative to the p27-KID^{wt}. The p27-KID^{p21helix} mutant is about 8 fold less potent than the p27-KID^{wt}, while the p27-KID^{p57helix} is about 16 fold less potent relative to the wild-type p27-KID.

The p27-KID^{ΔC} mutant is also a potent inhibitor of the P-Cdk2/cyclin A kinase activity. This mutant has an IC₅₀ value of 2.56 nM. Like the chimeric mutants, the p27-KID^{ΔC} mutant is less potent than the p27-KID^{wt} (about 12 times less potent). This suggests that the conserved residues contribute marginally to the kinase inhibitory function of p27.

3.5 Summary

In this chapter, it has been demonstrated that the binding the D1 and D2 peptides to the P-Cdk2/cyclin A complex is less favorable relative to that of p27-KID^{wt}. It has also been shown that the D2 peptide cannot completely inhibit the P-Cdk2/cyclin A kinase activity.

In addition, I have shown that the linker domains of the Cip/Kip proteins play similar structural roles, the differences in their sequences notwithstanding. The free energies of association of chimeric mutants (created by substituting the linker domain of p27 with p21 and p57 linker domains) with the P-Cdk2/cyclin A complex are very comparable to the free energy of association of wild-type p27-KID with the same complex. The melting temperatures of the ternary complexes formed by the chimeric mutants on binding the P-Cdk2/cyclin A complex are very similar to that of the ternary of the p27-KID^{wt}. The chimeric

mutants are potent inhibitors of the kinase activity of P-Cdk2/cyclin A, with IC_{50} values in the low nM range.

Data from the third study shows that three conserved residues within the linker domains of the Cip/Kip proteins do not contribute significantly to the Gibbs free energy of association in the binding of p27-KID to the P-Cdk2/cyclin A. However, the conserved residues contribute to the thermal stability of the ternary complex formed when p27-KID binds the P-Cdk2/cyclin A complex. Further, these residues contribute, albeit marginally, to the inhibition of the P-Cdk2/cyclin A kinase activity.

Chapter 4: Consequences of perturbation of the structure of the linker domain of p27

The nascent helical structure of the linker domain is conserved amongst members of the Cip/Kip family of cyclin-dependent kinase inhibitors; however, its role in function is not known. To investigate the role of the nascent structure of the linker domain, mutants with higher or lower helical contents than the wild-type p27-KID in the linker domain were prepared and the consequences of the structure perturbations on the interaction of p27-KID with P-Cdk2/cyclin A were monitored. First, the solvent exposed residues within the linker domain were substituted with residues that either stabilize or destabilize helices. The secondary structures of the mutants were determined by CD. The consequences of the mutations on the thermodynamics of association of p27 and the P-Cdk2/cyclin A were analyzed by ITC. CD thermal denaturation studies were used to determine the thermal stability of the ternary complexes formed by p27 and its variants bound to P-Cdk2/cyclin A. The effects of the mutations on p27 function were determined by Cdk2 inhibition assays and cell cycle arrest analysis.

4.1 Secondary structure analysis of the more and less helical p27-KID mutants

The mutants selected for these studies were part of a larger library of mutants as listed in Table 2.2. Figure 4.1 shows the CD spectra of p27-KID^{wt},

Figure 4.1 CD spectra of the p27-KID linker domain mutants.

p27-KID^{+H}, p27-KID^{-H}, and p27-KID^{loop} and the differences in their helical contents. p27-KID^{loop} exhibits random coil structure, suggesting strongly that the helical contribution at 222 nm observed in the p27-KID^{wt} is localized in the linker domain and not in D1 or D2. This is consistent with previous studies.¹⁹⁷

4.2 The free energy of the interaction of p27-KID with P-Cdk2/cyclin A does not decrease when the helical content in the linker domain of p27 is increased

The thermodynamic binding parameters associated with the interaction of the p27-KID linker domain mutants and the P-Cdk2/cyclin A complex were determined by ITC. Figure 4.2 shows the ITC binding isotherms for these mutants. The interactions of the linker helix mutants with the P-Cdk2/cyclin A complex, involve dramatic folding-on-binding and are accompanied by large entropic costs, but not to the same extent as the p27-KID^{wt}. The observed $-T\Delta S$ values for the p27-KID^{wt}, p27-KID^{+H}, p27-KID^{-H}, and p27-KID^{loop} are $+32.2 \pm 1.2$ Kcal mol⁻¹, $+22.3 \pm 2.5$ Kcal mol⁻¹, $+19.0 \pm 1.0$ Kcal mol⁻¹, and $+20.0 \pm 1.4$ Kcal mol⁻¹ respectively (Table 4.1). This suggests that p27-KID^{+H}, p27-KID^{-H}, and p27-KID^{loop} do not fold to the same extent upon binding to P-Cdk2/cyclin A as p27-KID^{wt}. The folding of the mutants on binding the P-Cdk2/cyclin A complex results in a large number of favorable contacts. All the mutants release a large amount of heat when they bind the P-Cdk2/cyclin A complex. Nevertheless, the enthalpies of interaction for the mutants are smaller in magnitude relative to that for p27-KID^{wt}. The enthalpies of interaction for the p27-KID^{wt}, p27-KID^{+H},

Figure 4.2 ITC binding isotherms for the interaction of the p27-KID mutants with the P-Cdk2/cyclin A complex. The transition regions for p27-KID^{wt} and p27-KID^{loop} are expanded.

Table 4.1 Thermodynamic binding parameters of the p27-KID mutants.

Interaction	K_d (nM)	ΔG^a (kcal mol ⁻¹)	ΔH (kcal mol ⁻¹)	$-T\Delta S^b$ (kcal mol ⁻¹)
p27-KID ^{wt} + P-Cdk2/cyclin A	3.34 ± 1.64	- 11.61 ± 0.30 ^c	- 43.77 ± 1.38	+ 32.16 ± 1.22
p27-KID ^{+H} + P-Cdk2/cyclin A	23.41 ± 6.09	- 10.42 ± 0.15	- 32.72 ± 2.57	+ 22.3 ± 2.46
p27-KID ^{-H} + P-Cdk2/cyclin A	3.66 ± 0.91	- 11.52 ± 0.14 ^c	- 30.48 ± 0.82	+ 18.96 ± 0.95
p27-KID ^{loop} + P-Cdk2/cyclin A	138.3 ± 43.05	- 9.37 ± 0.18	-29.12 ± 1.19	+ 19.7 ± 1.36

In all the experiments above, the stoichiometry of binding was determined as one (1), with experimental error within 10%. The errors reported above are the standard deviations from the mean of three independent experiments.

^a Calculated with the equation $\Delta G = -RT \ln(1/K_d)$.

^b Calculated with the equation $\Delta G = \Delta H - T\Delta S$.

^c The c values for these p27-KID variants are very high and the ΔG values reported here are taken as estimates of the upper limits of K_d . Ideal c values for accurate determination of K_d , have a range of 1 to 500. The c values for the binding of p27-KID^{wt} and p27-KID^{-H} to P-Cdk2/cyclin A complex are 2212 and 1698, respectively.

p27-KID^{-H}, and p27-KID^{loop} are $-43.8 \pm 1.4 \text{ kcal mol}^{-1}$, $-32.7 \pm 2.6 \text{ kcal mol}^{-1}$, $-30.5 \pm 0.8 \text{ kcal mol}^{-1}$, and $-29.1 \pm 1.2 \text{ kcal mol}^{-1}$ in the order given. These enthalpies of interaction, for the wild-type and mutants, overcome the respective unfavorable interaction entropies resulting in overall favorable Gibbs free energies of interaction. ΔG values of $-11.6 \pm 0.3 \text{ kcal mol}^{-1}$, $-10.4 \pm 0.2 \text{ kcal mol}^{-1}$, $-11.5 \pm 0.1 \text{ kcal mol}^{-1}$, and $-9.4 \pm 0.2 \text{ kcal mol}^{-1}$ were obtained for the p27-KID^{wt}, p27-KID^{+H}, p27-KID^{-H}, and p27-KID^{loop}, respectively. The $\Delta\Delta G$ value for the p27-KID^{+H} with respect to the p27-KID^{wt} is $+1.2 \text{ kcal mol}^{-1}$ demonstrating that increasing the helical content of the linker domain does not make the interaction of p27-KID^{wt} with P-Cdk2/cyclin A more favorable; in fact, it actually makes it slightly less favorable.

4.3 The ternary complexes formed by the linker helix mutants on binding P-Cdk2/cyclin A are thermodynamically less stable relative to that of p27-KID^{wt}

The P-Cdk2/cyclin A complex has a modest thermal stability with a melting temperature (T_m) of $54.7 \pm 0.6 \text{ }^\circ\text{C}$.²¹⁶ When p27-KID^{wt} binds to the P-Cdk2/cyclin A complex, the thermal stability of the ternary complex is dramatically increased to $82.3 \pm 0.5 \text{ }^\circ\text{C}$. Figure 4.3 shows the thermal denaturation curves of the ternary complexes formed by the interaction of the p27-KID linker helix mutants and the P-Cdk2/cyclin A complex. The T_m for values for complexes containing p27-KID^{+H}, p27-KID^{-H}, and p27-KID^{loop} were $80.3 \pm 0.2 \text{ }^\circ\text{C}$, $78.5 \pm 0.6 \text{ }^\circ\text{C}$, and $73.0 \pm 0.3 \text{ }^\circ\text{C}$ respectively (Table 4.2). The results indicate that perturbation of

Figure 4.3 Thermal denaturation curves for the p27-KID variants.

Table 4.2 Melting temperatures of the ternary complexes formed by the p27-KID mutants on binding the P-Cdk2/cyclin A complex.

Complex	T_m (°C)
p27-KID ^{wt} /P-Cdk2/cyclin A	82.3 ± 0.5
p27-KID ^{+H} /P-Cdk2/cyclin A	80.3 ± 0.2
p27-KID ^{-H} /P-Cdk2/cyclin A	78.5 ± 0.6
p27-KID ^{loop} /P-Cdk2/cyclin A	73.0 ± 0.3

the helical content of the linker domain has a net effect of decreasing the thermal stability of the ternary complexes. Reducing the helical content (p27-KID^{-H}) or completely eliminating the nascent helical structure (p27-KID^{loop}) has a greater impact on the thermal stability than increasing the helical content (p27-KID^{+H}). The decrease in the melting temperature of the ternary complex containing the p27-KID^{+H} mutant relative to that of p27-KID^{wt} indicates that restricting the dynamics of the linker domain by stabilizing its helical structure has a destabilizing effect on the p27-KID^{wt} ternary complex. On the other hand, the T_m 's obtained for the p27-KID^{-H} and p27-KID^{loop} mutants suggest that excessive flexibility of the linker domain is also destabilizing.

4.4 The linker helix mutants are less potent inhibitors of P-Cdk2/cyclin A kinase activity relative to p27-KID^{wt}

In the last chapter, it was shown that p27-KID is a potent inhibitor of the kinase activity of the P-Cdk2/cyclin A complex and that a Cdk binding domain peptide (D2) cannot completely inhibit kinase activity. The D2 peptide partially inhibits P-Cdk2/cyclin A kinase activity with an IC_{50} of about 125 nM. However, the D2 peptide fails to completely inhibit P-Cdk2/cyclin A with 25% residual kinase activity observed even at inhibitor peptide concentrations as high as 17 μ M. These data suggest that the Cdk binding domain requires the other domains of the KID (the cyclin binding domain and the linker domain) for effective inhibition. The structure of the p27-KID/Cdk2/cyclin A complex shows that the the LH domain of p27 is an α -helix in this ternary complex.¹⁰⁴ However, the role

of the helical propensity of the LH domain of unbound p27 in binding and inhibition is not known. I performed kinase inhibition assays to uncover the contribution of the structure of the linker domain to Cdk inhibition.

All of the linker helix mutants inhibit kinase activity in the low nanomolar range. Nevertheless, none of the mutants is more potent than the wild-type KID. Figure 4.4 shows the raw inhibition data and normalized inhibition curves for the p27-KID mutants. The values for the p27-KID^{wt}, p27-KID^{+H}, p27-KID^{-H}, and p27-KID^{loop} were 293.8 pM, 5.1 nM, 353.0 pM, and 16.0 nM (Table 4.3), respectively. The results obtained for the p27-KID^{loop} mutant are very interesting because they show that a KID molecule formed by coupling the cyclin and Cdk binding domains with an unstructured flexible linker can inhibit the P-Cdk2/cyclin A complex with an IC₅₀ as low as 16 nM. Increasing the helical content of the linker domain decreases inhibition potency relative to the p27-KID^{wt}. The p27-KID^{+H} variant is about 17 times less potent than the wild-type KID in inhibiting P-Cdk2/cyclin A kinase activity. However, reducing the helical content (p27-KID^{-H}) does not diminish the inhibition potency of the KID to the same extent. The p27-KID^{-H} is almost as potent as the wild-type KID. Since the linker domain does not have extensive interactions with either cyclin A or P-Cdk2, its effects on phosphorylation are most likely a consequence of the role it plays in the folding of the Cdk binding domain.

It should be noted that the disparity between the IC₅₀ value, (16 nM), and K_d value, (138 nM), for the p27-KID^{loop} mutant is likely due to kinetic effects. In Figure 4.2, the expanded regions compare the raw data for the ITC binding

(a)

(b)

Figure 4.4 P-Cdk2/cyclin A kinase inhibition by the p27-KID mutants. (a) Gels showing kinase inhibition the p27-KID mutants. (b) Inhibition curves of the p27-KID variants.

Table 4.3 Inhibition potencies of the p27-KID mutants.

Variant	IC₅₀ (nM)	95% confidence intervals of IC₅₀ (nM)^a
p27-KID ^{wt}	293.8 pM	0.17 – 0.28
p27-KID ^{+H}	5.06 nM	3.3 – 5.1
p27-KID ^{-H}	353 pM	0.26 – 0.33
p27-KID ^{loop}	15.95 nM	9.9 – 25.7

^a Error is reported as the 95% confidence interval of the IC₅₀ from a curve fit of a triplicate data set.

isotherms for p27-KID^{wt} and p27-KID^{loop}. From the figure it is clear that unlike the p27-KID^{wt} which equilibrates to a completely flat baseline between injections, the injection peaks of the p27-KID^{loop} mutant do not fully equilibrate to a flat baseline in the time between injections. The equilibration time between injections is 5 minutes. On the other hand, during the kinase assays, P-Cdk2/cyclin A was mixed with p27-KID^{loop} and incubated for 1 hour prior to addition of ATP. It is conceivable that in an hour the binding reaction reaches equilibrium explaining the difference between K_d and IC_{50} . Such a long equilibration time would not be practical in an ITC experiment because it would require a total experiment time of at least 40 hours. In addition, the P-Cdk2/cyclin A complex would precipitate if kept at 25 °C for such an extended period of time making such an experiment unfeasible.

4.5 Perturbation of the structure of the linker domain is deleterious to p27 function in cells

Previous studies have shown that up-regulation of p27 causes cell cycle arrest during the G1/S cell cycle transition.¹¹¹ The effects of increasing or decreasing the helical content of the linker domain on cell cycle arrest were probed by over-expressing the p27 mutants in mouse fibroblast cells immortalized using the *NIH 3T3* protocol.²³² The linker LH domain mutants were expressed in the context of the full-length p27 protein. The level of protein expression was evaluated by measuring the GFP fluorescence of *NIH 3T3* cells using fluorescence activated cell sorting (FACS) analysis (Figure A.7). The DNA

content of cells expressing the greatest amount of GFP (top 10%) was measured for cell cycle analysis. Relative to a vector only negative control, the flp27^{wt}, caused cell cycle arrest and reduced the percentage of cells in S-phase from 28.4% to 14% (Table 4.4, Figure A.8). The mutant with the stabilized helix, flp27^{+H}, also caused cell cycle arrest. However, its cell cycle arrest capacity relative to flp27^{wt} is diminished. 18.2% of the cells progressed to S-phase when flp27^{+H} mutant was over-expressed. The greatest effects on cell cycle arrest were observed when the structure of the linker domain was either destabilized (flp27^{-H}) or completely eliminated (flp27^{loop}). When either flp27^{-H} or flp27^{loop} was over-expressed, 28.5% and 30.5% of the cells respectively progressed to the S-phase from the G1 phase. Compared to the vector only negative control (28.4%), the flp27^{-H} and the flp27^{loop} mutants do not appear to cause cell cycle arrest.

4.6 Summary

In this chapter, the thermodynamic and functional effects of perturbations of the structure of the linker domain of p27 have been reported. There was a decrease in the enthalpy of association of p27 with the P-Cdk2/cyclin A complex when the structure of the linker domain was perturbed. However, this reduction in enthalpy was compensated for by a proportional reduction in the entropic cost associated with the folding of the p27-KID mutants on binding the P-Cdk2/cyclin A complex. This compensation caused the Gibbs free energies of association of all the p27-KID variants to vary within a very narrow range. The ternary

Table 4.4 Cell cycle analysis of cells.

Cell cycle phase	No plasmid	Empty vector (GFP only)	p27 ^{wt}	p27 ^{+H}	p27 ^{-H}	p27 ^{loop}
G0/G1	55.3 ± 2.6	22.2 ± 2.1	57.9 ± 0.4	48.5 ± 3.0	38.7 ± 3.2	39.9 ± 2.3
S	34.4 ± 2.1	28.4 ± 2.4	14 ± 1.0	18.2 ± 1.7	28.1 ± 1.1	30.5 ± 1.2
G2/M	10.3 ± 1.3	49.5 ± 2.6	28.2 ± 0.6	33.3 ± 1.4	33.1 ± 3.0	29.6 ± 1.5

The values represent the percentage of cells at the various phases of the cell cycle. Triplicate experiments were done for each control and p27-KID variant.

complexes formed when the mutants bind to the P-Cdk2/cyclin A complex were thermally less stable than that formed with wild-type p27-KID. All the mutants were less potent inhibitors of the P-Cdk2/cyclin A complex relative to the p27-KID^{wt}. While enhancing the helical structure of the linker domain has a slight negative impact on the ability of p27 to cause cell cycle arrest, destabilizing or replacing this structure with a loop is deleterious to p27 function.

Chapter 5: Discussion

5.1 The linker domain of p27 is necessary for function

The studies of the interactions of D1 and D2 with P-Cdk2/cyclin A support my hypothesis that the linker domain of p27 is necessary for function. The crystal structure of the p27/Cdk2/cyclin A complex shows that the linker domain makes few contacts with the Cdk2/cyclin A complex. It follows that the sum of the enthalpy of association of D1 with the P-Cdk2/cyclin A complex and the enthalpy of association of D2 with the same complex should either be equal to or slightly greater than (more positive) the interaction of p27 with the P-Cdk2 cyclin A complex. However, the total observed enthalpy for the D1 and D2 interactions ($-47.69 \text{ kcal mol}^{-1}$) with the binary complex was less than (more negative) that for p27-KID^{wt} ($-43.77 \text{ kcal mol}^{-1}$). This suggests that even though D1 and D2 bind to the P-Cdk2/cyclin A complex, one or both domains does (do) not make exactly the same contacts with P-Cdk2/cyclin A as they do in the context of p27-KID^{wt}. A previous study reported by our laboratory suggests that it is unlikely that D1 misfolds.¹⁹⁷ First, the binding of D1 to the cyclin does not involve large conformational changes which would require overcoming many local free energy barriers. Secondly, the enthalpy of the interaction D1 to the P-Cdk2/cyclin A complex ($-16.1 \pm 0.2 \text{ kcal mol}^{-1}$) is very similar to that of the p27-KID^{wt} on binding cyclin A ($-17.2 \pm 0.3 \text{ kcal mol}^{-1}$).¹⁹⁷ This suggests that favorable contacts made in both cases are very similar, if not identical.

On the other hand, the binding of D2 to the Cdk2 subunit of the kinase complex involves a number of significant conformational changes and dramatic folding of p27-KID and remodeling within Cdk2. This makes it likely that the D2 peptide does not make the same contacts with Cdk2 as does the D2 domain in the context of p27-KID^{wt}. The results of the kinase inhibition assays confirm this (Figure 3.5). ITC shows that D2 binds to the kinase complex with reduced affinity ($K_d = 82.96 \pm 17.89$ nM). However, D2 is unable to completely inhibit Cdk2 kinase activity. Differences in the dynamics of bound D2 peptide relative to that of bound D2 in the context of the p27-KID may account for the differences in Cdk2 kinase inhibition observed for the D2 peptide and p27-KID^{wt}. I propose that the D2 peptide bound to Cdk2 is more dynamic than Cdk2-bound D2 in the context of the p27-KID^{wt}. The more dynamic nature of the Cdk2-bound D2 peptide allows some ATP molecules to access to the ATP-binding pocket thereby enabling the 25% residual kinase activity observed during the kinase inhibition assays. I further propose that the coupling of D2 to D1 by the linker domain in p27-KID^{wt} restricts the motion of D2 on binding Cdk2. Because of the motional restriction, the Cdk-binding domain of p27-KID^{wt} completely occludes the ATP binding pocket thereby blocking ATP access and *ipso facto* completely inhibiting Cdk2 kinase activity.

In addition to facilitating kinase inhibition, the coupling of D1 and D2 by the linker domain makes the interaction of p27-KID^{wt} with P-Cdk2/cyclin A more favorable than the interactions of either D1 or D2 peptides with the same complex. The ΔG of the binding of p27-KID^{wt} to P-Cdk2/cyclin A complex (-11.61

$\pm 0.30 \text{ kcal mol}^{-1}$) is more favorable relative to that of either D1 ($-10.06 \pm 0.07 \text{ kcal mol}^{-1}$), or D2 ($-9.66 \pm 0.12 \text{ kcal mol}^{-1}$). This raises a question as to how the coupling of D1 and D2 by the linker domain makes the binding of p27-KID^{wt} to the P-Cdk2/cyclin A more favorable than the binding of D1 and D2 peptides to the same complex.

For the binding of p27-KID^{wt} to P-Cdk2/cyclin A to be more favorable than the binding of either D1 or D2, the coupling of D1 and D2 with the linker domain has to either cause net increase the amount of heat released (net increase in the enthalpy of binding) or increase the overall disorder of the system (net increase in entropy). From the structure of the p27-KID/Cdk2/cyclin A it is evident that the linker domain makes very few contacts with either Cdk2 or cyclin A. It is therefore unlikely that the linker domain decreases the free energy of the binding of p27-KID^{wt} to P-Cdk2/cyclin A (relative to the binding of D1 and D2 peptides to P-Cdk2/cyclin A) by causing a net increase in the enthalpy of binding.

The other explanation is that the coupling of D1 and D2 with the linker domain decreases the free energy of binding of p27-KID^{wt} to P-Cdk2/cyclin A, relative to the binding of D1 and D2 to the same complex, by causing a net increase in the entropy of binding. The net change in the entropy of binding is determined by two components: (i) the entropy of water resulting from the burial of hydrophobic side-chains (favorable entropy); and (ii) the conformational entropy that results from loss of translational and rotational degrees during protein folding (unfavorable entropy) and the motion of the protein after binding (favorable entropy).^{233,234} Since there is no significant burial of hydrophobic

residues within the linker domain when p27-KID^{wt} binds to Cdk2/cyclin A, the contribution of the entropy of water toward a net entropy increase should be minimal. This suggests that the net increase in entropy contribution of the linker domain toward the binding of p27-KID^{wt} to P-Cdk2/cyclin A is caused by a net positive change in conformational entropy. This positive change in entropy indicates that the contribution of the dynamics (positive entropy change) of the linker domain of p27 when bound to P-Cdk2/cyclin A is greater than the entropic contribution of the folding of the linker domain (negative entropy change). This is consistent with recent results from our laboratory (unpublished results). NMR analysis of p27-KID in complex with Cdk2/cyclin A has showed that the linker domain is dynamic even when p27 is bound to Cdk2/cyclin A. The same has also been observed for the linker domain of p21-KID in a p21-KID/Cdk2/cyclin A complex (unpublished results). I propose that the dynamic nature of the linker domain of the Cip/Kip proteins contributes significantly to the free energy of their association with Cdk complexes.

5.2 The linker domains of the Cip/Kip proteins play similar structural roles despite their sequence differences

The sequences of the linker domains of the Cip/Kip proteins are very variable but have a conserved helical structure. An analysis of the linker domain of each Cip/Kip protein in different species shows very high sequence conservation. It is curious why nature has selected for disparate sequences in the linker domains of the Cip/Kip proteins. It is reasonable to argue that if just a

helical structure was required to couple D1 and D2, then any sequence that could adopt a helical structure could have been selected for; but, this is clearly not the case. This may indicate that these sequences may play well defined roles in the interactions of the Cip/Kip proteins with Cdk's. This poses a question as to whether the different sequences of the linker domains of the Cip/Kip proteins confer each protein different thermodynamic and functional properties when they bind to their Cdk targets.

ITC studies show that the free energies of association of the chimeric mutants on binding the P-Cdk2/cyclin A complex are very similar. The p27-KID^{wt} has a ΔG value of $-11.61 \text{ kcal mol}^{-1}$, which is very close to the $-11.7 \pm 0.27 \text{ kcal mol}^{-1}$ and $-12.71 \pm 0.17 \text{ kcal mol}^{-1}$ obtained for the p27-KID^{p21^{helix}} and the p27-KID^{p57^{helix}}, respectively. This shows that the three linker domains despite their differences in sequence contribute very similarly to the free energy of association of the Cip/Kip proteins with the P-Cdk2/cyclin A complexes. One difference observed in the thermodynamic parameters is the magnitude of the enthalpy of binding.

The chimeric mutants exhibit decreased value of the enthalpy of binding relative to that for p27-KID^{wt}. p27-KID^{p21^{helix}} and p27-KID^{p57^{helix}} exhibit enthalpies of $-27.94 \pm 1.26 \text{ kcal mol}^{-1}$ and $-31.15 \pm 0.51 \text{ kcal mol}^{-1}$, respectively, in comparison to p27-KID^{wt} which has an enthalpy of $-43.77 \text{ kcal mol}^{-1}$ on binding P-Cdk2/cyclin A. The kinase assay results show that these chimeric variants are potent inhibitors albeit slightly less potent than p27-KID^{wt}. This shows that D2 in the chimeric mutants binds to Cdk2 and is folded in a conformation very similar to

that of D2 of the p27-KID^{wt}. It is unlikely that D1 of the chimeric mutants is not binding since the free energies of association of the chimeric mutants with P-Cdk2/cyclin A are more comparable to the free energy of the association of p27-KID^{wt} and P-Cdk2/cyclin A than to the free energy of association of either the D2 peptide and the binary complex or p27-KID with monomeric Cdk2.¹⁹⁷ If D1 was not binding, the free energies of the association of the chimeric mutants and P-Cdk2/cyclin A should reflect binding only to Cdk2. This suggests that the enthalpy decrease observed in the ITC experiments is likely a result of the linker domains (since D1 and D2 bind) of the chimeric mutants not having exactly the same favorable interactions on binding P-Cdk2/cyclin A as p27-KID^{wt}. While the loss of enthalpy contributes unfavorably to the free energy of interaction, it is countered by a favorable decrease in the entropic penalty associated with the folding on binding. The $-T\Delta S$ values for the p27-KID^{p21helix} and the p27-KID^{p57helix} are $+16.24 \pm 1.29 \text{ kcal mol}^{-1}$ and $+18.44 \pm 0.35 \text{ kcal mol}^{-1}$ respectively, in comparison to the greater entropic cost of $+32.16 \pm 1.22 \text{ kcal mol}^{-1}$ of the interaction of p27-KID^{wt} with P-Cdk2/cyclin A. The proportional decrease in the entropic penalty effectively compensates for the enthalpic loss in the chimeric mutants resulting in free energy values very close to that of the p27-KID^{wt}. The chimeric mutants exhibit an entropy-enthalpy compensation phenomenon that makes their interaction with the P-Cdk2/cyclin A complex favorable.

It has been shown that the binding of the p27-KID^{wt} to the P-Cdk2/cyclin A complex greatly stabilizes the formed ternary complex.²¹⁶ The melting temperature of the P-Cdk2/cyclin A complex, which is only $54.7 \pm 0.6 \text{ }^{\circ}\text{C}$,

increases to 82.3 ± 0.5 °C when the p27-KID^{wt} binds. The binding of the chimeric mutants stabilize the binary complex to extents very similar to that of the p27-KID^{wt}. The ternary complexes of p27-KID^{p21helix} and p27-KID^{p57helix} have melting temperatures of 80.7 ± 0.2 °C and 80.1 ± 0.1 °C respectively. This suggests that the non-conserved residues in the linker domain contribute minimally to the stability of the ternary complexes formed when the Cip/Kip proteins bind the binary complex. The thermal denaturation data taken together with the ITC and kinase inhibition assay data suggest that the linker domains of the Cip/Kip proteins, despite their sequence differences, play very similar structural roles in binding to Cdk2/cyclin A.

5.3. The role of the conserved residues in the linker domain of Cip/Kip proteins

Only 3 (Glu39, Leu41 and Arg43) out of the 22 residues in the linker domain are conserved. I mutated these residues to glycines to probe their role in the interaction of the p27-KID with the P-Cdk2/cyclin A complex. The thermodynamic parameters obtained from ITC are very similar to those of the p27-KID^{wt}. There is a difference in the enthalpy of p27-KID^{ΔC} mutant and p27-KID^{wt}. The enthalpy of association of p27-KID^{ΔC} is -39.65 ± 0.13 kcal mol⁻¹, a decrease of 4.12 kcal mol⁻¹ relative to that of the p27-KID^{wt}. This indicates that the conserved residues contribute about 4 kcal mol⁻¹ to the enthalpy of interaction of the p27-KID^{wt} with P-Cdk2/cyclin A.¹⁰⁴ This difference is most likely caused by the mutation of Leu41 to Gly. In the p27/Cdk2/cyclin A structure,

Leu41 is buried into the p27-KID-cyclin A interface and mutating it to a smaller side chain, Gly, reduces the buried surface area resulting in a reduction of the amount of heat released. The other two conserved residues Glu39 and Arg43 are largely solvent exposed. Their mutation to Gly's probably does not have a great effect on the enthalpy. However, the Gly mutations may affect the dynamics of the linker helix by making it more flexible since the glycine side chain can access a broad range of conformational states. Like the chimeric mutants, this slight loss of enthalpy is compensated for and the free energy of association is the same as that of the p27-KID^{wt} within standard error. The free energy of association of p27-KID^{ΔC} with the binary complex is -11.89 Kcal mol⁻¹.

Even though the thermodynamic parameters associated with p27-KID^{ΔC} mutant are very similar to those associated with p27-KID^{wt}, the ternary complex this mutant forms on binding the binary complex is significantly less stable than that formed by p27-KID^{wt}. The binding of p27-KID^{ΔC} is associated with a T_m of 74.1 ± 0.6 °C, about 8 °C less than that associated with p27-KID^{wt}. This demonstrates that the conserved residues contribute significantly to the stability of the ternary complex.

The p27-KID^{ΔC} mutant is slightly less potent than the p27-KID^{wt}. The IC₅₀ of this mutant is 2.56 nM while the p27-KID^{wt} has an IC₅₀ value of 0.22 nM. This clearly indicates that the effects of mutations which are very close to D1 are propagated along the length of the helix and are able to affect the interaction D2 and Cdk2 resulting in the decrease in inhibition potency. This attests to the dynamic nature of the linker domain of the Cip/Kip proteins. In summary, the

conserved residues in the linker domains of the Cip/Kip proteins contribute to the enthalpy of binding of p27-KID to P-Cdk2/cyclin A, the kinase activity of p27-KID^{wt}, and also play a role in increasing the thermal stability of the ternary complex formed on when p27 binds P-Cdk2/cyclin A.

5.4 Biophysical and functional effects of increasing, decreasing, or eliminating the helical structure of the linker domain of p27

The nascent helical structure of the linker domain is conserved amongst members of the Cip/Kip family. I probed the consequences of perturbing this conserved structure on the biophysical and functional properties of p27 by studying the interactions of p27 mutants, in which the helical content of the linker domain was either increased (p27-KID^{+H}), decreased (p27-KID^{-H}), or completely eliminated (p27-KID^{loop}), with the P-Cdk2/cyclin A. The binding of p27 to Cdk2/cyclin A is associated with a great entropic cost. I hypothesized that the preformed helix in the linker domain of p27 decreases this entropic cost.

Results from the ITC analysis of the binding of p27-KID^{wt}, p27-KID^{+H}, and p27-KID^{-H} to P-Cdk2/cyclin A disprove this hypothesis. While the relative $-T\Delta S$ values for the binding p27-KID^{wt} and p27-KID^{+H} ($+32.16 \pm 1.22 \text{ kcal mol}^{-1}$ and $+22.3 \pm 2.46 \text{ kcal mol}^{-1}$ respectively) to P-Cdk2/cyclin A seem to support this hypothesis, an analysis of the values of the other thermodynamic parameters does not. If the p27-KID^{+H} bound to the P-Cdk2/cyclin A complex in the same way as the p27-KID^{wt}, its association with the binary complex would be more favorable than that of the p27-KID^{wt}. However, from my experimental

observation, this is not the case. The ΔG values of the association of p27-KID^{wt} and p27-KID^{+H} with P-Cdk2/cyclin A are $-11.61 \pm 0.3 \text{ kcal mol}^{-1}$ and $-10.42 \pm 0.15 \text{ kcal mol}^{-1}$, respectively (Table 4.1). The $\Delta\Delta G$ value of the association of p27-KID^{+H} and P-Cdk2/cyclin A with respect to the p27-KID^{wt} is $+1.2 \text{ kcal mol}^{-1}$. An explanation for the less favorable free energy of association of p27-KID^{+H} and P-Cdk2/cyclin A with respect to p27-KID^{wt} can be found in the values of the enthalpy that were observed for the two p27-KID variants. The enthalpies of association for the interaction of p27-KID^{wt} and p27-KID^{+H} with P-Cdk2/cyclin A are $-43.77 \pm 1.38 \text{ kcal mol}^{-1}$ and $-32.72 \pm 2.57 \text{ kcal mol}^{-1}$. The heat released by the binding of p27-KID^{+H} to P-Cdk2/cyclin A is less than that for the p27-KID^{wt}. This indicates the p27-KID^{+H} mutant does not make exactly the same contacts as does p27-KID^{wt} on binding P-Cdk2/cyclin A. Where are there differences in the contacts made by p27-KID^{+H} and p27-KID^{wt} on binding P-Cdk2/cyclin A?

It is unlikely the net enthalpy decrease in the binding of p27-KID^{+H} to P-Cdk2/cyclin A relative to p27-KID^{wt} is due to lack of binding or proper folding of either D1 or D2. If D1 did not bind, the binding affinity of the interaction of p27-KID^{+H} and P-Cdk2/cyclin A ($K_d = 23.41 \pm 6.09 \text{ nM}$) should be similar to the binding affinity of the interactions of p27 to Cdk2 ($70.3 \pm 23.1 \text{ nM}$)¹⁹⁷ or D2 peptide to P-Cdk2/cyclin A ($K_d = 82.96 \pm 17.89 \text{ nM}$), which it does not. The results from the P-Cdk2/cyclin A kinase inhibition assays (Table 4.3) show that D2 binds and folds, like the p27-KID^{wt}, into an inhibition competent conformation. This indicates that the reduced enthalpy observed for the binding of p27-KID^{+H} to P-Cdk2/cyclin A relative to p27-KID^{wt} stems from differences in the favorable

contacts within the linker domain of p27-KID^{+H}. Since the linker domain of the p27-KID^{+H} is already stabilized and does not need fold to the same extent as the p27-KID^{wt} on binding the P-Cdk2/cyclin A complex, the heat that is released during the folding of the linker domain helix of p27-KID^{+H} on binding P-Cdk2/cyclin A is reduced.

In addition to the thermodynamic studies of the p27-KID^{+H}, the studies of the p27-KID^{-H} also show that the nascent helix of p27 does not reduce the entropic cost associated with binding. The $-T\Delta S$ value of p27-KID^{-H} and p27-KID^{wt} on binding P-Cdk2/cyclin A are $+18.96 \pm 0.95 \text{ kcal mol}^{-1}$ and $+32.16 \pm 1.22 \text{ kcal mol}^{-1}$, respectively. . If the p27-KID^{-H} mutant folds in the same way as p27-KID^{wt} upon binding, its entropic cost should be higher than that for wild-type p27-KID. This suggests that the p27-KID^{-H} mutant does not fold to the same extent as p27-KID^{wt}. The ability of p27-KID^{-H} to inhibit P-Cdk2/cyclin A kinase activity shows that the conformation of D2 on binding the kinase complex is similar in both the p27-KID^{wt} and p27-KID^{-H}. The IC_{50} values for p27-KID^{wt} and p27-KID^{-H} are 0.3 nM and 0.35 nM respectively (Table 4.3). It is unlikely that D1 in the p27-KID^{-H} mutant is not binding since the dissociation constant of the binding of the D2 peptide to P-Cdk2/cyclin A ($82.96 \pm 17.89 \text{ nM}$) is much higher than that observed for the p27-KID^{-H} ($3.66 \pm 0.91 \text{ nM}$). This implies that the linker domain of this mutant does not fold on binding P-Cdk2/cyclin A. This is the most likely cause of the decreased enthalpy of binding observed for the association of the p27-KID^{-H} and the P-Cdk2/cyclin A relative to p27-KID^{wt}. The observed enthalpies for the binding of the p27-KID^{wt} and p27-KID^{-H} with the

P-Cdk2/cyclin A are $-43.77 \pm 1.38 \text{ kcal mol}^{-1}$ and -30.48 ± 0.82 respectively.

Like the p27-KID^{+H} and p27-KID^{-H} mutants, the ITC thermodynamic parameters and the P-Cdk2/cyclin A kinase inhibition assays indicate that the D1 and D2 domains of the p27-KID^{loop} mutant bind to P-Cdk2/cyclin A. The binding of the D2 domain of the p27-KID^{loop} mutant is shown by its ability to potently and completely inhibit P-Cdk2/cyclin A kinase activity ($IC_{50} = 16 \text{ nM}$). In Section 5.1 above, I discussed that complete kinase inhibition requires coupling of D1 and D2 by the linker domain. The ability of the p27-KID^{loop} mutant to completely inhibit kinase activity indicates that the D1 domain of this mutant binds to the cyclin A subunit of P-Cdk2/cyclin A. The $-T\Delta S$ value of this mutant ($+19.7 \pm 1.36 \text{ kcal mol}^{-1}$) therefore shows that its linker domain does not fold to the same extent as the p27-KID^{wt} ($-T\Delta S = +32.16 \pm 1.22 \text{ kcal mol}^{-1}$) on binding P-Cdk2/cyclin A. This is not surprising since this mutant was designed with residues that would minimize the formation of any helical structure. Expectedly, the enthalpy of the p27-KID^{loop} mutant on binding P-Cdk2/cyclin A is less than the enthalpy of the interaction p27-KID^{wt} and the same complex. The enthalpies of the interactions of p27-KID^{loop} and p27-KID^{wt} with P-Cdk2/cyclin A are $-29.12 \pm 1.19 \text{ kcal mol}^{-1}$ and $-43.77 \pm 1.38 \text{ kcal mol}^{-1}$, respectively. The results in the studies above suggest that the folding of the linker domain of the wild-type p27 may be an inherent feature of its sequence and is possibly the reason for the conservation of the p27 linker domain sequence in different species.

Stabilization of the linker domain affects the binding affinity of p27 for

P-Cdk2/cyclin A. The K_d values for the p27-KID^{wt} and p27-KID^{+H} were determined to be 3.34 ± 1.64 nM and 23.41 ± 6.09 nM respectively. The dissociation constants of the binding of the p27-KID^{-H} and p27-KID^{loop} mutants to the P-Cdk2/cyclin A complex were 3.66 ± 0.91 nM and 138.3 ± 43.05 nM respectively. The p27-KID^{-H} mutant has the same affinity for the P-Cdk2/cyclin A complex as the wild-type p27-KID, suggesting that flexibility of the linker domain may be important for binding to the P-Cdk2/cyclin A complex. On the other hand, restriction of the motions of the linker domain (p27-KID^{+H}) decreases the affinity of p27 for the P-Cdk2/cyclin A complex. However, complete elimination of the helical structure of the linker domain (p27-KID^{loop}) greatly decreases affinity for the P-Cdk2/cyclin A complex. Together these data suggest that the dynamics of the linker domain are necessary for tight binding of p27 to P-Cdk2/cyclin A.

The thermal denaturation results indicate that stabilization or destabilization of the helical structure of the linker domain of p27 decreases the thermal stability of the ternary complex formed when p27 binds P-Cdk2/cyclin A. Melting temperatures for the ternary complexes of the p27-KID^{wt}, p27-KID^{+H}, p27-KID^{-H}, and p27-KID^{loop} were 82.3 ± 0.5 °C, 80.3 ± 0.2 °C, 78.5 ± 0.6 °C, and 73.0 ± 0.3 °C respectively. The thermal stability of the ternary complexes is determined by the difference between the free energies of the native and unfolded states of these complexes. The more negative the difference obtained by subtracting the free energy of the native state from the free energy of the unfolded state, the more unstable the complex. The destabilization of the ternary complexes p27-KID^{+H} and p27-KID^{loop} mutants relative to that of p27-KID^{wt} is

therefore expected since the free energies of their native states are less favorable relative to that of the p27-KID^{wt}. The Gibbs free energy values of the native states of the p27-KID^{wt}, p27-KID^{+H}, and p27-KID^{loop} are -11.61 ± 0.30 kcal mol⁻¹, -10.42 ± 0.15 kcal mol⁻¹, and -9.37 ± 0.18 kcal mol⁻¹, respectively (Table 4.1). On the other hand, the free energy of the native state of the ternary complex of the p27-KID^{-H} mutant ($\Delta G = -11.52 \pm 0.14$ kcal mol⁻¹) has the same free energy as that for p27-KID^{wt} (-11.61 ± 0.30 kcal mol⁻¹) within standard error. The destabilization of this mutant therefore stems from the stabilization of its unfolded state. The destabilization of these mutants with structurally perturbed linker domains suggests that the structure of the wild-type linker domain of p27 is important for formation of a stable ternary complex.

Although p27-KID^{wt} is as potent as p27-KID^{-H} in inhibiting P-Cdk2/cyclin A kinase activity, it is more potent than either p27-KID^{+H} or p27-KID^{loop}. The observed IC₅₀ values for the p27-KID^{wt}, p27-KID^{-H}, p27-KID^{+H} and p27-KID^{loop} were 0.3 nM, 0.35 nM, 5 nM and 16 nM respectively. Being potent kinase inhibitors, I expected these mutants to be effective in causing cell cycle arrest. However, cell cycle arrest analysis showed that all of the p27 mutants do not cause cell cycle arrest. When flp27^{wt}, flp27^{+H}, flp27^{-H}, and flp27^{loop} are over-expressed in mouse fibroblasts, 14%, 18.2%, 28.1%, and 30.5% of the cells progressed to the S phase relative to a vector only control in which 28.4% of the cells progress to S phase (Table 4.4). These results show that while stabilizing the linker domain of p27 in cells impairs the cell cycle arrest function, destabilizing the linker domain or completely eliminating its helical structure

completely annuls the capacity of p27 to cause cell cycle arrest. This raises a question as to why the flp27^{-H} and flp27^{loop} mutants cannot cause cell cycle arrest even though they are potent inhibitors of P-Cdk2/cyclin A kinase activity *in vitro*. One explanation is that the structural changes in the linker domains of these mutants may alter their specificity. During G1/S cell cycle transition, Cdk4, Cdk6, Cdk1, and Cdk2 kinase complexes phosphorylate a number of downstream targets facilitating the progression of the cell cycle from the G1 to the S phase. The altered specificity of these mutants may decrease their binding to one or more of these Cdk complexes resulting in leaky inhibition of the G1/S transition. This may explain the results observed for the flp27^{-H} and flp27^{loop} mutants. The results of the cell cycle arrest analysis studies attest to the significance of the structure of the linker domain of p27 in biological function.

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Appendixes

A.1 Buffers

Pre-scission cleavage buffer (P)

50 mM Tris pH 7.0
150 mM NaCl
1 mM EDTA
1 mM DTT

Gel filtration buffer (GF)

20 mM HEPES pH 7.5
300 mM NaCl
5 mM DTT

Nickel binding buffer (Ni-A)

20 mM Tris pH 8.0
5 mM imidazole
500 mM NaCl

Nickel elution buffer (Ni-B)

20 mM Tris pH 8.0
500 mM imidazole
500 mM NaCl

GST binding buffer (GST-A) pH 7.3

140 mM NaCl
2.7 mM KCl
10 mM Na₂HPO₄
1.8 mM KH₂PO₄
5 mM DTT

GST elution buffer (GST-B)

50 mM Tris pH 8.0
10 mM reduced glutathione
5 mM DTT

Q-sepharose column binding buffer (Q-A)

20 mM HEPES pH 7.5

20 mM NaCl

5 mM DTT

Q-sepharose column elution buffer (Q-B)

20 mM HEPES pH 7.5

1.0 M NaCl

5 mM DTT

Kinase assay buffer (KA)

20 mM HEPES pH 7.3

25 mM sodium b-glycerolphosphate

15 mM MgCl₂

16 mM EGTA

0.5 mM activated Na₃VO₄

10 mM DTT

HPLC column binding buffer (HPLC-A)

0.1 % TFA (v/v) in H₂O

HPLC column elution buffer (HPLC-B)

0.1 % TFA in acetonitrile

Thrombin cleavage buffer for cyclin A (TcycA)

20 mM Tris pH 8.0

1.0 M NaCl

2.5 mM CaCl₂

Nickel binding buffer for D2 peptide (Ni-A/D2)

20 mM Na₂HPO₄·7H₂O pH 6.5

500 mM NaCl

5 mM imidazole

Nickel elution buffer for D2 peptide (Ni-B/D2)

20 mM Na₂HPO₄·7H₂O pH 6.5

500 mM NaCl

500 mM imidazole

Thrombin cleavage buffer (T)

20 mM Tris pH 8.0

50 mM NaCl

5 mM DTT

HPLC sample buffer (D)

10 mM Tris pH 7.2

5 mM NaCl

CD buffer

10 mM phosphate pH 7.0

5 mM DTT

Thermal denaturation buffer

1 mM phosphate pH 7.0

25 mM NaCl

1 mM DTT

A.2 Growth media

Luria-Bertani media (LB)

Tryptone	10 g
Yeast extract	5 g
NaCl	10 g
Antibiotic (kanamycin 30 mg/ml stock, ampicillin 50 mg/ml stock)	1 mL
H ₂ O	1 L

A.2 Tables and figures

Table A.1 Sequences of oligonucleotides used in preparing mutagenesis cassettes for the p27-KID mutants.

p27-KID mutant	Oligonucleotide sequences
p27-KID ^{p21helix}	5' GTAACGCATATGGAGCACCCCAAGCCCTCGGCCTGCAGGAACCTCT TCGGCCCCGGTGGACAGCGAACAGCTGAGCCGCGATTGCGATGCGCTGA TGGCGGGCTG 3'
	5' CCCC GCCTCGAGGGGTTTGTGATTCTGAAAATCGAAATTCCAGCGT TCGCGCGCTTCCTGAATGCAGCCCCGCCATCAGCGCATCGCAATCGCGG CTCAGCTGTT 3'
p27-KID ^{p57helix}	5' GTAACGCATATGGAGCACCCCAAGCCCTCGGCCTGCAGGAACCTCT TCGGCCCCGGTGGACCATGAAGAACTGAGCCGCGAACTGCAGGCGCGCC TGGCGGAACT 3'
	5' CCCC GCCTCGAGGGGTTTGTGATTCTGAAAATCGAAATTCCAGCGG TTCTGATCTTCCGCGTTTCAGTTCCGCCAGGCGCGCCTGCAGTTTCGCGG CTCAGTTCTT 3'
p27-KID ^{ΔC}	5' GCCTCCCATATGGAGCACCCCAAGCCCTCGGCCTGCAGGAACCTCT TCGGCCCCGGTGGACCACGGCGAGGGCACCGGCGAATTGGAGAAGCACT GCAGAGACAT 3'
	5' CTTGGGCTCGAGGGGTTTGTGATTCTGAAAATCGAAATTCCACTTG CGCTGGCTCGCCTCTTCCATGTCTCTGCAGTGCTTCTCCAAGTCGCCG GTGCCCTCGC 3'
p27-KID ^{A10}	5' GCCTCCCATATGGAGCACCCCAAGCCCTCGGCCTGCAGGAACCTCT TCGGCCCCGGTGGACCACGGCGGCGTTAACC GCGGACTTGGAGGCGCACT GCGCGGCGAT 3'
	5' CTTGGGCTCGAGGGGTTTGTGATTCTGAAAATCGAAATTCCACGCC GCCTGGCTCGCCGCTTCCATCGCCGCGCAGTGCGCCTCCAAGTCCGCG GTTAACGCCG 3'

(continued)

Table A.1 (continued).

p27-KID mutant	Oligonucleotide sequences
p27-KID^{A22}	5' GCCTCCCATATGGAGCACCCCAAGCCCTCGGCCTGCAGGAACCTCT TCGGCCCCGGTGGACGCGGCAGCAGCGGCCGAGCAGCAGCAGCGGCGGCAG CAGCCGCGGC 3'
	5' CTTGGGCTCGAGGGGTTTGTGATTCTGAAAATCGAAATTCCAGGCC GCGGCCGCGCTGCTGCCCCGCGGCTGCTGCCCCGCTGCTGCTGCTGCG GCCGCTGCTG 3'
p27-KID^{SL1}	5' GCCTCCCATATGGAGCACCCCAAGCCCTCGGCCTGCAGGAACCTCT TCGGCCCCGGTGGACCACCTGGAGCTGACCCTGGACTTGGAGCTGCACT GCCTGCTGAT 3'
	5' CTTGGGCTCGAGGGGTTTGTGATTCTGAAAATCGAAATTCCACTTG CGCTGGCTCAGCAGTTCCATCAGCAGGCAGTGCAGCTCCAAGTCCAGG GTCAGCTCCA 3'
p27-KID^{SL2 a}	5' GCCTCCCATATGGAGCACCCCAAGCCCTCGGCCTGCAGGAACCTCT TCGGCCCCGGTGGACCACGAGGAGCTGACCCTGGACTTGGAGCTGCACT GCCTGCTGAT 3'
	5' CTTGGGCTCGAGGGGTTTGTGATTCTGAAAATCGAAATTCCACTTG CGCTGGCTCAGCAGTTCCATCAGCAGGCAGTGCAGCTCCAAGTCCAGG GTCAGCTCCT 3'
p27-KID^{SL3} (p27-KID^{+H})	5' GCCTCCCATATGGAGCACCCCAAGCCCTCGGCCTGCAGGAACCTCT TCGGCCCCGGTGGACCACGAGGAGCTGACCAAGGACTTGGAGCTGCACT GCCTGCTGAT 3'
	5' CTTGGGCTCGAGGGGTTTGTGATTCTGAAAATCGAAATTCCACTTG CGCTGGCTCAGCAGTTCCATCAGCAGGCAGTGCAGCTCCAAGTCCTTG GTCAGCTCCT 3'

(continued)

Table A.1 (continued)

p27-KID mutant	Oligonucleotide sequences
p27-KID^{SL4}	5' GCCTCCCATATGGAGCACCCCAAGCCCTCGGCCTGCAGGAACCTCT TCGGCCCCGGTGGACCACGAGGAGCTGACCAAGGACTTGGAGCTGCACT GCAAGCTGAT 3'
	5' CTTGGGCTCGAGGGGTTTGTGATTCTGAAAATCGAAATTCCACTTC TTCTGGCTCAGCTCTTCCATCAGCTTGCACTGCAGCTCCAAGTCCTTG GTCAGCTCCT 3'
p27-KID^{SL5}	5' GCCTCCCATATGGAGCACCCCAAGCCCTCGGCCTGCAGGAACCTCT TCGGCCCCGGTGGACCACCTGGAGCTGACCCTGGACTTGGAGGAGCACT GCCTGAAGAT 3'
	5' CTTGGGCTCGAGGGGTTTGTGATTCTGAAAATCGAAATTCCACTTC AGCTGGCTCTCCAGTTCCATCTTCAGGCAGTGCTCCTCCAAGTCCAGG GTCAGCTCCA 3'
p27-KID^{SL6}	5' GCCTCCCATATGGAGCACCCCAAGCCCTCGGCCTGCAGGAACCTCT TCGGCCCCGGTGGACCACCTGCTGTTAACCCTGGACTTGGAGCTGCACT GCCTGCTGAT 3'
	5' CTTGGGCTCGAGGGGTTTGTGATTCTGAAAATCGAAATTCCACAGC AGCTGGCTCAGCAGTTCCATCAGCAGGCAGTGCACTCCAAGTCCAGG GTTAACAGCA 3'
p27-KID^H	5' GCCTCCCATATGGAGCACCCCAAGCCCTCGGCCTGCAGGAACCTCT TCGGCCCCGGTGGACCACGAAGAGTTAACCGAGGACTTGGAGGAGCACT GCGAGGAGAT 3'
	5' CTTGGGCTCGAGGGGTTTGTGATTCTGAAAATCGAAATTCCACTCC TCCTGGCTCTCCTCTTCCATCTCCTCGCAGTGCTCCTCCAAGTCCTCG GTTAACCTCTT 3'

(continued)

Table A.1 (continued)

p27-KID mutant	Oligonucleotide sequences
p27-KID ^{loop}	<p data-bbox="607 457 1373 541">5 ' GCCTCCCATATGGAGCACCCCAAGCCCTCGGCCTGCAGGAACCTCT TCGGCCCGGTGGACACCAACCCGGGCAACGGCGGCCATGGCGGCACCG GCGTGGGCGG 3 '</p> <p data-bbox="607 611 1373 695">5 ' CTTGGGCTCGAGGGGTTTGTGATTCTGAAAATCGAAATTCCAGCCA TGGCCGTTGCCGCCGTTGCCGCCACGCCGGTGCCGCCATGGCCGCCG TTGCCCGGGT 3 '</p>

Table A.2 Reagents for overlap extension PCR.

Reagent	Volume (μL)
Forward oligonucleotide (10 ng/μl)	1
Reverse oligonucleotide (10 ng/μl)	1
Forward end primer (125 ng/μl)	1
Reverse end primer (125 ng/μl)	1
Deoxynucleosides (dNTP's), 10 mM	1
PCR buffer	5
Enzyme ^a	1
Distilled, deionized water	40
Total Reaction Volume	50

^a The enzyme used was the expand high fidelity PCR system by Roche (Mannheim, Germany).

Table A.3 Thermocycler settings for the oePCR.

Step	Temperature	Time
1	95 °C	45 seconds
	Start cycle	
2	95 °C	45 seconds
3	55 °C	1 minute
4	72 °C	1 minute
	Go to step 2 for 20 cycles	
5	72 °C	10 minutes
6	End	

Table A.4 Sequences of oligonucleotides and primers used in preparing cloning cassettes for the Cip/Kip linker domain peptides.

Peptide	Sequences
p27 linker domain	<p>5' AAAAAACCGCGGGGTTCTTCTCATCATCATCATCAT TCTTCTGGTCTGGTTCCACGTGGTTCTCATATGCATGAAGA ACTGACTCGTGATCTGGA 3'</p> <p>5' TTTTTTGAATTCTTATTACCATTTACGTTGAGAAGCTTC TTCCATATCACGACAATGTTTTTCCAGATCACGAGTCAGTT CTTCATGCATATGAGAAC 3'</p> <p>5' AAAAAACCGCGGGGTTCTTCTCATCATC 3'</p> <p>5' TTTTTTGAATTCTTATTACCATTTACGT 3'</p>
p21 linker domain	<p>5' AAAAAACCGCGGGGTTCTTCTCATCATCATCATCATCAT TCTTCTGGTCTGGTTCCACGTGGTTCTCATATGAGCGAACA GCTGAGCCGCGATTGCGA 3'</p> <p>5' TTTTTTGAATTCTTATTACCAGCGTTTCGCGCGCTTCCTG AATGCAGCCCGCCATCAGCGCATCGCAATCGCGGCTCAGCT GTTGCTCATATGAGAAC 3'</p> <p>5' AAAAAACCGCGGGGTTCTTCTCATCATC 3'</p> <p>5' TTTTTTGAATTCTTATTACCAGCGTTTCG 3'</p>
p57 linker domain	<p>5' AAAAAACCGCGGGGTTCTTCTCATCATCATCATCATCAT TCTTCTGGTCTGGTTCCACGTGGTTCTCATATGCATGAAGA ACTGAGCCGCGAACTGCA 3'</p> <p>5' TTTTTTGAATTCTTATTACCAGCGGTTCTGATCTTCCGC GTTGAGTTCCGCCAGGCGCGCCTGCAGTTTCGCGGCTCAGTT CTTCATGCATATGAGAAC 3'</p> <p>5' AAAAAACCGCGGGGTTCTTCTCATCATC 3'</p> <p>5' TTTTTTGAATTCTTATTACCAGCGGTTTC 3'</p>

Table A.5 Cloning of the p27-KID linker domain mutants.

Variant	Vector(s)
p27-KID ^{wt}	pET28, pET28GSTHis
p27-KID ^{A10}	pET28, pET28GSTHis
p27-KID ^{A22}	pET28GSTHis
p27-KID ^{SL1}	pET28
p27-KID ^{SL2} ^a	
p27-KID ^{SL3} (p27-KID ^{+H})	pET28
p27-KID ^{SL4}	pET28
p27-KID ^{SL5}	pET28
p27-KID ^{SL6}	pET28, pET28GSTHis
p27-KID ^{-H}	pET28GSTHis
p27-KID ^{loop}	pET28GSTHis
p27-KID ^{p21helix}	pET28GSTHis
p27-KID ^{p57helix}	pET28, pET28GSTHis
p27-KID ^{ΔC}	pET28, pET28GSTHis

^a The cassette for the p27-KID^{SL2} mutant could not be successfully cloned into either the pET28 or the pET28GSTHis vector.

Table A.6 Expression and solubility of the p27-KID linker domain mutants.

Variant	Expression vector(s)	Solubility
p27-KID ^{wt}	pET28, pET28GSTHis	Soluble
p27-KID ^{A10}	pET28GSTHis	Soluble
p27-KID ^{A22}	pET28GSTHis	Insoluble
p27-KID ^{SL1}	pET28	Insoluble
p27-KID ^{SL3} (p27-KID ^{+H})	pET28	Partly soluble
p27-KID ^{SL4}	pET28	Soluble
p27-KID ^{SL5}	pET28	Insoluble
p27-KID ^{SL6}	pET28GSTHis	Insoluble
p27-KID ^{-H}	pET28GSTHis	Soluble
p27-KID ^{loop}	pET28GSTHis	Soluble
p27-KID ^{p21helix}	pET28GSTHis	Soluble
p27-KID ^{p57helix}	pET28GSTHis	Soluble
p27-KID ^{ΔC}	pET28GSTHis	Soluble

Figure A.1 A schematic representation of oePCR.

Figure A.2 Agarose gel analysis of the oePCR for the p27 linker domain peptide. The expected product size 159 bp.

TGTGAGCGGATACATTCCCCTCTANAAATCCNCTTTGTTTAAATTTAAGAAGGAGATATAC
CATGGGCAGCAGCCATCATCATCATCATCACAGCAGCGGCCTGGTGCCGCGCGGCAGCCAT
ATGGAGCACCCCAAGCCCTCGGCCTGCAGGAACCTCTTCGGCCCCGGTGGACCACGAAGAG
TTAACC CGG GACTTGGAGAAGCACTGCAGAGACATGGAAGAGGCGAGCCAGCGCAAGTGG
AATTTTCGATTTTTCAGAATCACAAACCCCTAGAGGGCAAGTACGAGTGGCAAGAGGTGGAG
AAGGGCAGCTTGCCCGAGTTCTACTACAGACCCCCGCGGCCCCCAAAGGTGCCTGCAAG
GTGCCGGCGCAGGAGTAATAAGGATCCGAATTCGAGCTCCGTCGACAAGCTTGCGGCCGC
ACTCGAGCACCACCACCACCACCCTGAGATCCGGCTGCTAACAAGCCCCGAAAGGAAGC
TGAGTTGGCTGCTGCCACCGCTGAGCAATAACTAGCATAACCCCTTGGGGCCTCTAAACG
GGTCTTGAGGGGTTTTTTTGCTGAAAGGAGGAACCTATATCCGGATTGGCGAATGGGACGCG
CCCTGTAGCGGNGCATTAAAGCGCGGCGGCTGTGGTGGTTACGCGCCANCGNGACCGCTAC
ACTTGCCAGCGCCCTAGCGACCGCTCCTTTTCGCTTTCTTCCCTTNCTTTCTCNCCACGTT
CGCCGGCTTTTCCTCNTCAAGCTCTAAATCGGGGGCTNCCCTTTAGGGNTCCGNTTTANT
GCTTTACGGCACCTCNACCC

Figure A.3 Sequencing results for the pET28p27-KID plasmid. This sequence is only for the p27-KID coding region.

AGAAGGAGATATACCATGGGCAGCAGCCATCATCATCATCACAGCAGCGGCCTGGTG
CCGCGCGGCAGCCATATGTCAAACGTGCGAGTGTCTAACGGGAGCCCTAGCCTGGAGCGG
ATGGACGCCAGGCAGGCGGAGCACCCCAAGCCCTCGGCCTGCAGGAACCTCTTCGGCCCCG
GTGGACCACGAAGAGTTAACCCGGGACTTGGAGAAGCACTGCAGAGACATGGAAGAGGCG
AGCCAGCGCAAGTGGAAATTTGATTTTCAGAATCACAAACCCCTAGAGGGCAAGTACGAG
TGGCAAGAGGTGGAGAAGGGCAGCTTGCCCGAGTTCTATTACAGACCCCGCGGCCCCC
AAAGGTGCCTGCAAGGTGCCGGCGCAGGAGAGCCAGGATGTCAGCGGGAGCCGCCCGGCG
GCGCCTTTAATTGGGGCTCCGGCTAACTCTGAGGACACGCATTTGGTGGACCCAAAGACT
GATCCGTCGGACAGCCAGACGGGGTTAGCGGAGCAATGCGCAGGAATAAGGAAGCGACCT
GCAACCGACGATTCTTCTACTCAAAACAAAAGAGCCAACAGAACAGAAGAAAATGTTTCA
GACGGTTCCCCAAATGCCGGTTCTGTGGAGCAGACGCCCAAGAACCTGGCCTCAGAAGA
CGTCAAACGTAATAAGGATCCGAATTCGAGCTCCGTCGACAAGCTTGCGGCCGCACTCGA

Figure A.4 Sequencing results for the pET28flp27 plasmid. This sequence is only for the flp27 coding region.

Figure A.5 Vector map for the MSCV-I-GFP plasmid. The highlighted EcoRI site was the cloning site for the flp27 linker domain mutants.

MSCV-I-GFPflp27^{wt}

CTTCTCTAGGCGCCGAATTCATGTCAAACGTGCGAGTGTCTAACGGGAGCCCTAGCCTGGAGCGGATGGACGCCAGGC
AGGCGGAGCACCCCAAGCCCTCGGCCTGCAGGAACCTCTTCGGCCCGGTGGACCACGAAGAGTTAACCCGGGACTTGGA
GAAGCACTGCAGAGACATGGAAGAGGCGAGCCAGCGCAAGTGGAAATTTTCGATTTTCAGAATCACAAACCCCTAGAGGGC
AAGTACGAGTGGCAAGAGGTGGAGAAGGGCAGCTTGCCCGAGTTCTATTACAGACCCCCGCGGCCCCCCAAAGGTGCCT
GCAAGGTGCCGGCGCAGGAGAGCCAGGATGTACAGCGGGAGCCGCCCGGGCGGCGCCTTTAATTGGGGCTCCGGCTAACTC
TGAGGACACGCATTTGGTGGACCCAAAGACTGATCCGTTCGGACAGCCAGACGGGGTTAGCGGAGCAATGCGCAGGAATA
AGGAAGCGACCTGCAACCGACGATTTCTTCTACTCAAAAACAAAAGAGCCAAACAGAACAGAAAGAAATGTTTCAGACGGTT
CCCCAAATGCCGGTTCTGTGGAGCAGACGCCCAAGAAGCCTGGCCTCAGAAGACGTCAAACGTATCCGTATGATGTGCC
GGATTATGCGTAATAAGAATTCGTTAACCTCGAGCGGGATCAATTCCGCCCCCCCCCTAACGTTACTGGCCGAAGC

MSCV-I-GFPflp27^{SL3} (MSCV-I-GFPflp27^{+H})

GCGCCGGANTTCATGTCAAACGTGCGAGTGTCTAACGGGAGCCCTAGCCTGGAGCGGATGGACGCCAGGCAGGCGGAGC
ACCCCAAGCCCTCGGCCTGCAGGAACCTCTTCGGCCCGGTGGACCACGAGGAGCTGACCAAGGACTTGAGACTGCACTG
CCTGCTGATGGAAGTCTGAGCCAGCGCAAGTGGAAATTTTCGATTTTCAGAATCACAAACCCCTCGAGGGCAAGTACGAG
TGGCAAGAGGTGGAGAAGGGCAGCTTGCCCGAGTTCTACTACAGACCCCCGCGGCCCCCCAAAGGTGCCTGCAAGGTGC
CGGCCGAGGAGAGCCAGGATGTACAGCGGGAGCCGCCCGGGCGGCGCCTTTAATTGGGGCTCCGGCTAACTCTGAGGACAC
GCATTTGGTGGACCCAAAGACTGATCCGTTCGGACAGCCAGACGGGGTTAGCGGAGCAATGCGCAGGAATAAGGAAGCGA
CCTGCAACCGACGATTTCTTCTACTCAAAAACAAAAGAGCCAAACAGAACAGAAAGAAATGTTTCAGACGGTTCCCCAAATG
CCGGTTCTGTGGAGCAGACGCCCAAGAAGCCTGGCCTCAGAAGACGTCAAACGTATCCGTATGATGTGCCGGATTATGC
GTAATAAGAATTCGTTAACCTCGAGCGGGATCAATTCCGCCCCCCCCCTAACGTTACTGGCCGAAGCCGCTTGAATAA
NGCCGGTGTGCGTTTGTCTATATGTTATTTTCCACCATATTG

MSCV-I-GFPflp27^H

CCCTTTATCCAGCCCTCACTCCTTCTCTAGGCGCCGAATTCATGTCAAACGTGCGAGTGTCTAACGGGAGCCCTAGCC
TGGAGCGGATGGACGCCAGGCAGGCGGAGCACCCCAAGCCCTCGGCCTGCAGGAACCTCTTCGGCCCGGTGGACCACGA
AGAGTTAACCGAGGACTTGAGGAGCACTGCGAGGAGATGGAAGAGGAGAGCCAGGAGGAGTGGAAATTTTCGATTTTCAG
AATCACAAACCCCTCGAGGGCAAGTACGAGTGGCAAGAGGTGGAGAAGGGCAGCTTGCCCGAGTTCTACTACAGACCCC
CGCGGCCCCCCAAAGGTGCCTGCAAGGTGCCGGCGCAGGAGAGCCAGGATGTACAGCGGGAGCCGCCCGGGCGGCGCCTTT
AATTGGGGCTCCGGCTAACTCTGAGGACACGCATTTGGTGGACCCAAAGACTGATCCGTTCGGACAGCCAGACGGGGTTA
GCGGAACAATGCGCAGGAATAAGGAAGCGACCTGCAACCGACGATTTCTTCTACTCAAAAACAAAAGAGCCAAACAGAACAG
AAGAAATGTTTCAGACGGTTCCCCGAATGCCGGTTCTGTGGAGCAGACGCCCAAGAAGCCTGGCCTCAGAAGACGTCA
AACGTATCCGTATGATGTGCCGGATTATGCGTAATAAGAATTCGTTAACCTCGAGCGGGATCAATTCCGCCCCCCCCCT
AACGTTACTGGCCGAAGCC

MSCV-I-GFPflp27^{loop}

GAATTCATGTCAAACGTGNGAGTGTCTAACGGGAGCCCTAGCCTGGAGCGGATGGACGCCAGGCAGGCGGAGCACCCCA
AGCCCTCGGCCTGCAGGAACCTCTTCGGCCCGGTGGACACCAACCCGGGCAACGGCGGCCATGGCGGCACCGGCGTGGG
CGGCAACGGCGGCAACGGCCATGGCTGGAATTTTCGATTTTCAGAATCACAAACCCCTCGAGGGCAAGTACGAGTGGCAA
GAGGTGGAGAAGGGCAGCTTGCCCGAGTTCTACTACAGACCCCCGCGGCCCCCCAAAGGTGCCTGCAAGGTGCCGGCGC
AGGAGAGCCAGGATGTACAGCGGGAGCCGCCCGGGCGGCGCCTTTAATTGGGGCTCCGGCTAACTCTGAGGACACGCATTT
GGTGGACCCAAAGACTGATCCGTTCGGACAGCCAGACGGGGTTAGCGGAGCAATGCGCAGGAATAAGGAAGCGACCTGCA
ACCGACGATTTCTTCTACTCAAAAACAAAAGAGCCAAACAGAACAGAAAGAAATGTTTCAGACGGTTCCCCAAATGCCGGTT
CTGTGGAGCAGACGCCCAAGAAGCCTGGCCTCAGAAGACGTCAAACGTATCCGTATGATGTGCCGGATTATGCGTAATA
AGAATTCGTTAACCTNGAGCGGGATCAATTCCGCCCCCCCCCTAACGTTACN

Figure A.6 Sequences of the MSCV-I-GFP plasmids. Only the region coding for the flp27 variants is shown.

Figure A.7 Flow cytometry analysis of GFP positive cells. The percentages of GFP positive cells for are circumscribed with red ellipses.

Figure A.8 Cell cycle arrest analysis of the p27-KID mutants.

Vita

Steve Biko Otieno was born on March 7, 1979 in Rachuonyo District, Kenya. He attended Maseno National School where he completed his studies in 1996. Steve received a Bachelor of Science degree in Chemistry from Rust College in Holly Springs, MS in 2001 and joined the Molecular Sciences program at The University of Tennessee Health Science Center in the fall of 2001. He began his doctoral work in the laboratory of Dr. Richard Kriwacki at St. Jude Children's Research Hospital in the summer of 2002. Steve expects to graduate with a Ph.D. in Molecular Sciences in May 2007.