Apoptotic Functions of PDCD10/CCM3, the Gene Mutated in Cerebral Cavernous Malformation 3

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Background and Purpose—Mutations in the Programmed Cell Death 10 (PDCD10) gene cause autosomal dominant familial cerebral cavernous malformations (CCM3). To date, little is known about the function of this gene and its role in disease pathogenesis.

Methods—We examined the effects of overexpression of wild-type and 2 human disease-causing variants of PDCD10 on cell death using 3 different methods (TUNEL and MTT assays and caspase-3 activation). We analyzed expression of CCM3, activated caspase-3, and p38 in endothelial cell lines using the serum deprivation model of apoptosis induction. Finally, we assayed the effects of siRNA-mediated inhibition of endogenous PDCD10 expression on cell death in endothelial cell cultures.

Results—Overexpression of wild-type CCM3, but not disease-linked mutant forms, induced apoptosis as confirmed by TUNEL and increased levels of activated caspase-3. Serum starvation of endothelial cells, an inducer of apoptosis, led to increased expression of CCM3 and activation of p38 and ultimately activated caspase-3. siRNA-mediated inhibition of CCM3 expression resulted in decreased levels of p38 and activated caspase-3, and decreased cell death.

Conclusions—CCM3 is both necessary and sufficient to induce apoptosis in vitro in well-defined cell culture systems. Even though it is currently unclear whether this effect on apoptosis is direct or indirect through modulation of cell cycle, these results led to the novel hypothesis that CCM lesions may form as a consequence of aberrant apoptosis, potentially altering the balance between the endothelium and neural cells within the neurovascular unit. (Stroke. 2009;40:1474-1481.)

Key Words: cerebral cavernous malformations • CCM3 • PDCD10 • apoptosis • p38 • caspase-3

Cerebral cavernous malformations (CCM) are vascular lesions affecting almost exclusively the central nervous system (CNS) and the retina. Grossly, they are a collection of enlarged sinusoidal vascular channels almost resembling a raspberry. Ultrastructurally, these channels are lined by a single layer of endothelium, lack normal vessel wall elements such as smooth muscle and are devoid of intervening normal CNS parenchyma. These lesions can occur sporadically, or as a familial form attributable to mutations in 3 different genes: Krev1 Interaction Trapped 1 (KRIT1; alternative name: CCM1), CCM2 (malcavernin) and Programmed Cell Death 10 (PDCD10; alternative name: CCM3). Overall, mutations in these 3 genes account for approximately 96% of familial cases. Despite the identification and characterization of these 3 CCM genes, several questions regarding this complex disorder remain unanswered. These include a sophisticated understanding of the function of the CCM genes and how mutations lead to formation of lesions that almost exclusively affect the CNS and are focal in nature. It has been suggested that the CCM lesions may be the consequence of hemorrhage-induced proliferation, a process by which abnormal vascular beds cause reactive angiogenesis with new vessel formation and coalescence, but mechanistic insight into lesion formation is lacking.

Recent molecular biological studies on the CCM1 and 2 proteins shed some light into their functions. CCM1 was shown to bind to microtubules in vitro and to interact individually with Krev1 and Rap1, ICAP1α (integrin cytoplasmic domain associated protein 1 alpha), β-catenin, and CCM2 in various in vitro systems. CCM2 binds CCM1 and MEKK3 in a ternary complex and acts as a scaffolding protein signaling through p38 after extracellular stimulation. The p38 pathway is known to be important in diverse physiological processes ranging from proliferation to differentiation to apoptosis. More recently, CCM3 was shown to be a part of this complex in vitro.

Despite this recent progress, little is known about the biological functions of the CCM3 protein. It was first identified through a screen of genes expressed during the induction of apoptosis in a premeloblast cell line. Recent evidence suggests...
that CCM3 binds to and is phosphorylated by serine/threonine kinase 25 (STK25) and is dephosphorylated by binding to the phosphatase domain of Fas-associated phosphatase-1.27

Based on these limited data, we sought to investigate the biological function of CCM3 and specifically focused on its potential involvement in apoptotic pathways. We modulated CCM3 expression using cell culture based assays in vitro and examined its effects on cell death. Our results indicate that CCM3 has a proapoptotic function in the cell culture models tested and led to the novel hypothesis that CCM lesions potentially form as a result of aberrant apoptosis in the vasculature within the neurovascular unit.

**Methods**

**Cell Culture**

HeLa cells were plated in high glucose DMEM supplemented with 10% FBS, 1% penicillin/streptomycin, and 1% L-glutamine. Primary human umbilical vein endothelial cells (HUVECs) were obtained from the Vascular Biology and Transplantation Program at Yale University (New Haven, Conn). HUVECs were grown in M199 medium with 20% fetal bovine serum (FBS) and 1% endothelial cell growth supplement (ECGS), on gelatin-coated (0.1%) plates (J.T. Baker Inc). Cells used in this study were obtained between passages 3 and 6. For serum starvation assays, HUVECs were seeded on 0.1% gelatin coated culture slides and after 48 hours of attachment in standard medium, cells were washed in 0.2% FBS medium, followed by an additional 24-hour incubation in 0.2% FBS.

**Expression Vector Construction**

A PCR-amplified human PDCD10/CCM3 (hCCM3) cDNA fragment was subcloned into Xhol and HindIII sites of the pEGFP-C3 vector (Clontech) or a pRFP-C3 vector containing monomeric red fluorescent protein, resulting in a C-terminal GFP- or RFP-tagged CCM3, respectively. Both expression plasmids carrying two human disease causing PDCD10/CCM3 point mutations (c.283C>T and c.192delA)12 were generated from the pEGFP-C3-hCCM3 plasmid using a standard mutagenesis kit (Stratagene). All constructs were verified by direct sequencing.

**Transfections**

HeLa cells were seeded (1×10⁵) in 6-well plates. The pEGFP-C3-hCCM3 or pRFP-C3-CCM3 and pEGFP-C3 or pRFP-C3 (empty) plasmids were transfected into the cell lines using the lipofectAMINE 2000 (Invitrogen Corp) according to the manufacturer’s protocols. After overnight incubation at 37°C to ensure attachment, the cells were transfected with one of the previously mentioned plasmids encoding either the GFP-tagged wild-type hCCM3, hCCM3 with a point mutation (c.283C>T), or a single base-pair deletion (c.192delA).

**Western Blotting**

Cells were harvested and total protein from the cells was extracted in a lysis buffer (50 mmol/L HEPES, pH:7.4, 150 mmol/L NaCl, 10% glycerol, 1% Triton X-100, 1.5 mmol/L MgCl₂-6H₂O, 1 mmol/L EGTA, 100 mmol/L NaF, 10 mmol/L sodium pyrophosphate and protease inhibitors, 1 mmol/L Na₃V₀₄, 10 mg/mL leupeptin, 10 mg/mL aprotinin and 4 mmol/L PMSF). Cell lysates were centrifuged to collect supernatant, and equal amounts of protein, 50 µg/lane, (Bio-Rad protein assay system; Bio-Rad Laboratories) were gelled to collect supernatant, and equal amounts of protein, 50 µg/lane, (Bio-Rad protein assay system; Bio-Rad Laboratories) were separated with 10% or 4% to 20% sodium dodecyl sulfate-polyacrylamide gel electrophoresis (Bio-Rad Laboratories) and transferred onto a PVDF membrane (Bio-Rad Laboratories). The membranes were incubated with primary antibodies: anticleaved caspase-3 (Cell Signaling Technology Inc), antiphospho-p38 or antip38 (Cell Signaling Technology), anti-GAPDH (Santa Cruz), or anti-GFP (Sigma Aldrich) followed by the appropriate fluorescence-conjugated (immunohistochemistry) or HRP-conjugated (Western blot) secondary antibodies.

**Apoptosis Assays**

Cells were permeabilized with 0.1% Triton X-100 in 0.1% sodium citrate for 2 minutes on ice and the terminal deoxynucleotidyl transferase-mediated dUTP nick end-labeling (TUNEL) reaction mixture (In Situ Cell Death Detection Kit, TMR-red, Roche) was applied for 1 hour at 37°C. Cell counts were performed at 10 different fields in 5 separate transfection experiments.

**Cell Viability Assays (MTT)**

Cell viability was measured using the 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl tetrazolium (MTT) assay. Briefly, 72 hours after transfection, MTT was added to each well at a final concentration of 0.5 mg/mL for 4 hours; the cells were subsequently lysed in dimethyl sulfoxide (DMSO) and the amount of MTT formazan was quantified by determining the absorbance at 550 nm, using a microplate reader. Cell viability is expressed as a percentage of the control culture value (GFP plasmid transfection group).

**PDCD10 siRNA Inhibition**

HUVECs were seeded on 6-well plates (5×10⁴ cells per well). After attachment overnight, cells were washed in PBS, and transfected with 3 sets of PDCD10 Stealth select siRNAs (25 bp; Unigene ID: Hs.478150), following the manufacturer’s protocol (Invitrogen Corporation). As control siRNA, stealth RNAi negative control (Invitrogen) was used. After 72 hours of incubation, HUVECs were washed and exposed to low serum (0.2% FBS). At different time points, HUVECs were harvested for Western blotting. The membranes were incubated with the following antibodies: antiphospho-p38 (Cell Signaling Technology), anti-CCM3,29 or anti-GAPDH (Santa Cruz).

To quantify cell death occurring in the cultures, terminal deoxynucleotidyltransferase-mediated UTP end labeling (TUNEL) analysis was performed as above. The percentage of apoptotic cells was calculated from the positively labeled cells in each field (400×) from at least 5 fields for each group.

**Statistical Analyses**

Data are presented as means±SE of absolute values or percent of control and were analyzed using either Student t test (Figures 1 and 2) or 1-way ANOVA (Figures 3 to 5). We initially performed a global ANOVA test to check for significance and when proven, proceeded with pair-wise comparisons to examine which groups were significantly different from the baseline. Bonferroni correction was applied to correct for multiple testing unless specified otherwise and P<0.05 was considered significant. The global ANOVA probability values and Bonferroni corrected probability values for pair-wise ANOVA tests are reported.

**Results**

**CCM3 Induces Apoptosis in HeLa Cells**

We analyzed the effects of CCM3 overexpression in cell culture by transfecting HeLa cells with the expression plasmid pEGFP-C3-hCCM3 (referred as GFP-CCM3 henceforth). GFP-CCM3 (wild-type) overexpression resulted in increased levels of apoptosis compared to vector alone (Figure 1). A significantly larger proportion of apoptotic cells with fragmented nuclei was observed in the population of GFP-positive (and, therefore, CCM3 expressing) cells transfected with GFP-CCM3 compared to those from control transfections with empty vector (Figure 1b). Protein expression of GFP and GFP-CCM3 was confirmed by immunoblotting with anti-GFP (Figure 1c).

We then investigated whether CCM3-mediated cell death is associated with caspase-3 activation. Caspase-3 has previously
been established as a terminal executioner of the apoptotic cell death pathway. Activated caspase-3 was detected in apoptotic cells with fragmented/condensed nuclei expressing CCM3 (Figure 2). These data indicate that CCM3 overexpression in HeLa cells results in caspase-3 activation and cell death.

Disease-Causing Mutated CCM3 Fails to Induce Apoptosis

We had previously reported 4 novel disease-causing PDCD10/CCM3 mutations in CCM families. We introduced 2 of these point mutations (c.283C>T [M1], c.192delA [M2]) into the GFP-CCM3 expression vector. Both mutations lead to production of truncated proteins (Figure 3a). We tested the effects of overexpression of mutated forms of CCM3 in HeLa cells. As expected, overexpression of wild-type CCM3 resulted in increased cleaved caspase-3 levels, whereas overexpression of either mutant form of CCM3 (c.283C>T, c.192delA) had no effect on cleaved caspase-3 expression (Figure 3b). In addition, overexpression of either CCM3 mutant resulted in reduced cell loss by the MTT assay compared to wild type CCM3 (Figure 3c), further suggesting that mutant forms of CCM3 fail to induce apoptosis.

Serum-Deprivation of Endothelial Cells Results in Increased CCM3 Levels

Expanding on a previously established assay for the induction of apoptosis in endothelial cells, we cultured human umbilical vein endothelial cells (HUVECs) in low-serum media. Serum-deprived HUVECs demonstrated an increase in cleaved caspase-3 protein levels by Western blotting after 3 hours as expected (Figure 4a). Interestingly, CCM3 expression increased on serum-deprivation starting at approximately 30 minutes, before the increase in cleaved caspase-3 levels (Figure 4b). After serum starvation, reincubation of cells in normal media (20% FBS) allows for cell recovery and restores (reduces) CCM3 expression to normal levels (Figure 4c). To establish the mechanism of this increase in CCM3 expression, we incubated the cells with different concentra-
In summary, serum deprivation of endothelial cells results in increases in CCM3 and phosphorylated p38 protein levels followed by caspase-3 activation.

siRNA Inhibition of CCM3 Expression Reduces Endothelial Cell Apoptosis During Serum Deprivation

Using the same serum-starvation model, we then reduced CCM3 protein levels with siRNAs as described in Methods. HUVECs treated with CCM3 siRNA demonstrated decreased levels of cell death, as shown by the TUNEL assay (Figure 5a and 5b). We then evaluated whether the reduction of cell death correlated with reduction of caspase-3 activation. Serum-starved HUVECs also demonstrated decreased activated caspase-3 levels after treatment with CCM3 siRNA (Figure 5c). This decrease in CCM3 and activated caspase-3 levels correlates with decreased p38 activation after CCM3 siRNA transfection compared to cells treated with siRNA controls (stealth RNAi negative control; Figure 5d).

Discussion

Because CCM lesions in all 3 inherited forms of the disease are clinically and pathologically nearly identical, we had hypothesized over 10 years ago that genes causing familial cavernous malformations CCM1, 2, and 3 (KRIT1, CCM2, and PDCD10, respectively) all participate in a common or parallel pathways,7,12,31 and thus disruption of signaling would lead to similar disease manifestations. To test this hypothesis, we previously investigated the expression patterns of CCM1, 2, and 3 and showed that these proteins spatially and temporally show similar expression patterns.29,31–33 These proteins are specifically expressed throughout the endothelium in many tissues and organs and within the neurovascular unit of the CNS that consists of astrocytes, neurons, and vascular endothelium.29,31–33 Consistent with this observation, recent data from yeast and mammalian systems demonstrate that CCM1, 2, and 3 physically interact,18,24,27,28 confirming findings in zebrafish which demonstrate that CCM1 and 2 act in a common pathway34 with ccm1 and ccm2 mutants displaying comparable vascular defects.35 Similarly, in vivo studies in the mouse based on Ccm1 and Ccm2 knockout models have revealed their nonredundant, vital roles in angiogenesis36,37 but failed to reveal any breakthrough insights into their function.

On the other hand, multiple levels of in vitro evidence suggest that the CCM genes might play a role in a variety of biological processes linking extracellular signals, including adhesion (through β1 integrin22,23 or β-catenin21 interactions) or stress (including osmo- or mechano-stress) to various biological responses, including changes in morphology and proliferation.38 In this article, we present data that link CCM signaling to apoptotic pathways.

Indeed, in vitro work has shown that the CCM1 protein, KRIT1, binds to microtubules17,18 in a folded state.19 Both Rap1a and ICAP1α can disrupt the association of KRIT1 with microtubules localizing it to endothelial cell–cell junctions.19,21 Interestingly, only ICAP1α appears to be able to unfold KRIT1.19 After localization to the cell membrane through its FERM domain, KRIT1 binds to β-catenin and plays a role in stabilizing adherens junctions and regulates...
endothelial permeability. Through its NPXY domains, KRIT1 also binds CCM2, potentially recruiting CCM2 to these junctions. CCM2 has been identified as a human paralog of the mouse osmosensing scaffold protein for MEKK3 (OSM), which, after binding with CCM1 and potentially this complex, might participate in the p38 mitogen-activated protein kinase (p38 MAPK) stress signaling pathway. More recent evidence suggests that CCM3, at least in certain biological states, might be part of this complex. Furthermore, on activation of this complex, CCM1 shuttles from the cytoplasm and the cell membrane into the nucleus and therefore might interact with other nuclear molecules, including transcription factors, which might influence basic cellular functions such as proliferation and apoptosis. Consistent with this hypothesis is the observation in Caenorhabditis elegans, in which kri-1, a CCM1 ortholog, interacts with DAF-16, a transcription factor, such that mutations in kri-1 lead to an increase in life span.

Based on these observations, we investigated the biological functions of CCM3 using overexpression or siRNA-mediated inhibition of PDCD10/CCM3 expression. Previously published studies showed CCM3 to be a part of a cohort of proteins upregulated on growth factor deprivation and apoptosis induction in the TF-1 premeloid cell line as well as in a fibroblast cell line. Based on these data which suggested that CCM3 might play a role in apoptosis signaling, we focused our attention on understanding the capacity of CCM3 to induce apoptosis in the absence of exogenous proapoptotic stimuli. Overexpression of CCM3 in HeLa cells was in fact sufficient to induce increased levels of cell death as evidenced by both nuclei morphology as well as caspase-3 activation, a well established common end pathway protein in apoptosis. This increase in apoptosis and activated caspase-3 levels was abrogated when cells were transfected with biologically inactive forms of CCM3, identified as mutations causing CCM in patients with the familial form of the disease.

We then focused on whether CCM3 was induced in the presence of proapoptotic stimuli. With the induction of serum starvation in HUVECs, CCM3 levels increased concomitantly with increased levels of activated caspase-3 and p38. Interestingly, this increase in the percentage of apoptotic cells and activated caspase-3 and p38 expression levels was diminished when endogenous CCM3 was blocked by siRNA, suggesting that CCM3 might be necessary for induction of apoptosis in this in vitro model. Furthermore, although HUVECs represent a well-characterized model of endothelial cell responses in culture, it is always possible that in the context of the organism, endothelial cells behave differently. Thus, in vivo experiments are needed to extend and confirm these findings. In addition, even though some of the molecules involved in CCM signaling are known, many remain anonymous. Identification of these molecules along with in vivo experiments are needed to dissect the biological role of CCM signaling. Our results demonstrating CCM3 to be both necessary and sufficient for programmed cell death in cell culture models provide a novel insight into this complex pathophysiology.

The effects of CCM3 on apoptosis could be direct or indirect through cell cycle modulation. Previous evidence in the literature suggests that interference with cell cycle progression can lead to apoptosis. If true, CCM signaling can
affect both proliferation and apoptosis in a cell type-dependent manner, potentially explaining observed effects of CCM3 on these biological processes. Further work will be needed to answer these questions. Our results, however, lead to the novel hypothesis of attenuated apoptosis affecting endothelial cells within the neurovascular unit as a molecular mechanism underlying cavernous malformations. If proven to be true in vivo, this might lead to a mechanistic understanding of the role of apoptosis and cell cycle control in CCM pathophysiology, potentially having significant implications in designing new therapies.

Conclusions
CCM3 is both necessary and sufficient to induce apoptosis in well-defined cell culture systems. These results led to a novel hypothesis that CCM lesions may form because of aberrant apoptosis, potentially altering the balance between the endothelium and neural cells within the neurovascular unit. Even though it is currently unclear whether this effect on apoptosis is direct or indirect through modulation of cell cycle, the current data provide a testable hypothesis about CCM lesion development and has potentially important implications for a mechanistic understanding of CCM pathophysiology.

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Disclosures

None.

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