Functional genetic screening reveals the role of mitochondrial cytochrome b as a mediator of FAS-induced apoptosis

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Functional selection of genetic suppressor elements (GSEs), engineered gene fragments that interfere with the function of a particular gene product, was used to identify regulators of FAS-induced apoptosis. Chicken DF-1 cells expressing human FAS receptor and susceptible to FAS-induced apoptosis were infected with a GSE library consisting of randomly fragmented normalized chicken cDNAs in a replication-competent avian retroviral vector. Virus-producing cells were subjected to several rounds of selection using FAS agonistic antibodies, resulting in isolation of a set of GSEs conferring resistance to FAS-induced apoptosis. Surprisingly, one of the isolated GSEs encoded a 42 amino acid-long polypeptide derived from the C-terminal half of cytochrome b (Cyt b) encoded by the mitochondrial genome. Subsequent experiments showed that caspase 8-dependent cleavage of mitochondrial Cyt b and translocation of its C-terminal half into the cytoplasm occurred during FAS-induced apoptosis in both chicken and human cells. Ectopic cytoplasmic expression of either full-length Cyt b or its C-terminal half in several human cell lines induced apoptosis, which could be suppressed by the isolated GSE, but not by Bcl2 over-expression or Apaf-1 or cytochrome c knock-down. These results reveal a cytochrome c-independent branch of FAS-induced apoptosis involving cleavage and cytoplasmic release of mitochondrial Cyt b.

FAS is a member of the death domain-containing tumor necrosis factor receptor family of proteins that regulate apoptosis (1–3). FAS-induced apoptosis occurs upon binding of its physiological ligand, FAS ligand (FASL), or experimental FAS agonistic antibodies (4). The major physiological function of this conserved apoptotic pathway is to support homeostasis of the immune system, directing elimination of self-reacting lymphocytes and proliferation and maturation of some lymphocyte populations (5). Deregulation of the FAS pathway has been implicated in various malignancies and diseases (4). Tumor cells often escape immune attack by suppressing the FAS pathway or by using it to their advantage (3, 6, 7). For example, some tumors up-regulate expression of FASL to induce apoptosis in infiltrating lymphocytes (4). Some viruses promote their propagation by expressing FAS-inhibitory proteins (8, 9). Mutations in FAS and some haplotypes of FAS splice variants are associated with human diseases (4).

Binding of FASL or FAS agonistic antibodies to FAS on the cell surface triggers the extrinsic pathway of apoptosis through formation of a death-inducing signaling complex (DISC) (10). DISC contains a number of adaptor proteins, caspases 8 and 10 and c-Flip, a negative regulator of caspase 8 (11) that is used by some tumor cells and viruses to acquire FAS resistance (12). DISC formation is the main mechanism of FAS-mediated killing in cells designated “type I” (H9, SKW6.4, human lymphocytes). However, in “type II” cells (Jurkat, CEM, HeLa, human hepatocytes), FAS-mediated apoptosis occurs via the intrinsic pathway (13). Type II cells do not form enough DISC to induce apoptosis directly; instead, DISC-activated caspase 8 cleaves the Bcl2 family member Bid (14), which triggers the mitochondrial pathway of apoptosis (15) involving cytochrome c (Cyt c) release and activation of the Apaf1/procaspase 9 apoptosisosome (16). Other mitochondrial components have recently been defined as apoptosis mediators, including AIF, SMAC/Diablo, and Omi (16–20); however, their relative impact on FAS-mediated death remains unclear.

In this study we sought to identify different components and modulators of the FAS pathway through an unbiased genetic suppressor element (GSE) (21) library screen. This method allows identification of genes associated with specific cellular phenotypes by functional selection of cells expressing GSEs, engineered gene fragments constructed to encode either antisense RNAs or dominant negative partial proteins. Developed in 1992 (22), GSE technology has been used to identify unique genes involved in tumor suppression, drug sensitivity, apoptosis, and growth regulation (21, 23, 24). In past studies, GSE libraries were constructed in mammalian retroviral vectors that required packaging cells to produce viral stocks of the GSE library and selected clones and a second cell type for functional selection. Because clone rescue between rounds of selection was laborious and somewhat unreliable, we have modified the method to use a GSE library constructed in an RCAS avian retroviral vector (25). Because RCAS vectors are replication competent, functional selection can be performed directly on the virus-producing cells.

We used this method to isolate GSEs conferring resistance to FAS apoptosis in chicken cells expressing human FAS. Several of the isolated GSEs corresponded to genes with known relevance to apoptosis. However, one corresponded to the cytochrome b (Cyt b), a mitochondrial DNA-encoded component of complex III of the mitochondrial electron-transport chain (26–28), which has not previously been linked to apoptosis or any other nonmitochondrial activity. We were intrigued by how a GSE-encoded fragment of a mitochondrial protein could affect FAS apoptosis when expressed in the cytoplasm. This apparent paradox was resolved by our work demonstrating a different role for a processed Cyt b protein as a cytoplasmic mediator of FAS-induced apoptosis.

Results

Identification of GSEs Suppressing FAS-Mediated Apoptosis. We designed a modified GSE screening approach that allows direct phenotypic selection of cells producing GSE-encoding retroviruses (see Materials and Methods for details). This was made possible by the use of a GSE library prepared in a replication-competent avian avian retroviral vector | Bcl-2 | caspase | cytochrome c | genetic suppressor element


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retroviral vector [RCAS series (25); see supporting information (SI) Fig. S1] comprised of randomly fragmented and normalized chicken cDNAs ligated to an adaptor enabling translation of the inserts in all three reading frames (a gift of E. Feinstein, see SI Text for details).

To use this library for identification of mediators of FAS-dependent apoptosis, we established a chicken cell line permissive to RCAS replication that is susceptible to apoptosis induced by human FAS agonistic antibodies. Chicken DF-1 cells transfected with an RCAS construct directing expression of human FAS (CD95) (Fig. 1A) became uniformly susceptible to FAS agonistic antibodies (Fig. 1B). This indicates that FAS was expressed in all of the DF-1FAS cells and that human FAS can function with chicken DISC components.

The scheme of isolation of GSEs conferring FAS resistance is shown in Fig. 1D. Briefly, DF-1FAS cells were infected with virus produced in DF-1 cells transfected with library DNA. Supernatant of DF-1FAS cells with the library was possible because the FAS cDNA and GSE library were cloned into RCAS vectors differing in their env genes (29–31). The protocol used for library transduction ensured delivery of the library in its full complexity with each of the 1 x 10^6 clones delivered to at least 10 cells.

The library-transduced DF-1FAS cells were treated with a low concentration of FAS antibody that allowed survival of one out of 10^6 untransduced cells. These conditions were chosen to allow isolation of relatively weak GSEs and quantitative assessment of their biological effects. The primary selection was performed on five plates of 10^7 library-transduced cells and a single plate of 10^7 cells transduced with insert-free virus. Secondary and tertiary rounds of selection were performed similarly. Each of the five branches of library selection was processed separately to reduce the risk of losing weak GSEs because of competition with stronger ones.

The first round of selection revealed an equivalent frequency of FAS resistance in the library- and insert-free vector-transduced populations (~500 clones were isolated from each of the five branches of the library experiment and ~100 clones were isolated in the control experiment). However, in the secondary and tertiary rounds of selection, the selected viruses produced an increase in FAS resistance as compared to the empty vector (Fig. 1C). The degree of FAS resistance in each branch remained at the same level after further rounds of selection, suggesting that a plateau had been reached that presumably reflects the biological efficacy of the selected viruses. It is possible that the strength of the effects of the isolated virus populations might be reduced by competition of FAS- and GSE-encoding RNAs during viral packaging. However, our successful isolation of confirmed bioactive GSEs (see below) indicates that such competition did not affect selection. Presumably GSE-encoding RNA has a packaging advantage because it is significantly shorter than the FAS-transcript.

Identification of the Genes Corresponding to FAS-Inhibitory GSEs. To identify the genes corresponding to selected GSEs, total RNA was isolated from the FAS-resistant populations of cells obtained after tertiary selection. RT-PCR was performed with a GSE-specific primer corresponding to the sense strand of the universal GSE adaptor (see Fig. S1B). The PCR products were cloned back into the RCAS(A) vector and sequenced. The isolated sequences are listed in Table S1. Some of them represented fragments of genes with known relevance to FAS-mediated apoptosis, including c-FLIP (12, 32), NK receptor KIR/NKB (33, 34), and protein kinase A (PKA) (35). The remainder corresponded to genes that have not been previously associated with apoptosis. A subset of these represented highly abundant transcripts (fibronectin and actin) or RNA sequences with no recognized relation to mammalian genes.

Because the same universal adaptor flanked each library insert at both ends, the orientation of the recovered GSEs that was effective in conferring FAS resistance was not known. We therefore prepared two constructs for each GSE differing in the orientation of the insert in the RCAS(A) vector and assessed their effect on the FAS sensitivity of DF-1FAS cells. Viruses grown in DF-1 cells were used to infect DF-1FAS cells, which were then treated with FAS-agonistic antibodies. For the majority of tested sequences, one of the two tested constructs was effective in conferring FAS resistance (Fig. 1E). No FAS resistance was associated with isolated fibronectin- and actin-derived sequences cloned in either orientation, confirming that they represent background (data not shown).

Seven out of nine GSEs with confirmed activity demonstrated the FAS protective effect only when cloned in their sense orienta-
Cyto b is cleaved during FAS-induced apoptosis and its C-terminal part is released into the cytoplasm. (A) Intracellular distribution of Cyt b in DF-1 cells left untreated (control) or treated with FAS-agonistic antibody in the absence or presence on the proteosomal inhibitor MG-132 (PI). Cytoplasmic and mitochondrial fractions prepared either 1 or 2 h after treatment were analyzed by Western blotting with antibodies against Cyt b, proliferating cell nuclear antigen (PCNA) (cytoplasmic marker) and Cyt c. Full-length (30 kDa) and cleaved (15 kDa) Cyt b are indicated by arrows. (B) Mitochondrial and cytoplasmic fractions were prepared from HeLa cells left untreated (control) or treated with FAS antibody at the indicated times after treatment (0.5–4 h). Fractions were analyzed by Western blotting for Cyt b (full-length, 30 kDa, and cleaved, 15 kDa), GRP75 (mitochondrial marker), PCNA (cytoplasmic marker), and Cyt c. (C) Western blot analysis of HeLa cells expressing full-length Cyt b, Cyt b512–1146 (C-term), or the portion of Cyt b corresponding to GSE F21 in the cytoplasm. All recombinant proteins contained a C-terminal FLAG tag and were detected using anti-FLAG antibodies. (D) Full-length Cyt b and Cyt b512–1146 are toxic when expressed in the cytoplasm. HeLa cells were infected with lentiviruses directing expressing of EGFP (negative control), full-length Cyt b, the C-terminal half of Cyt b (Cyt b512–1146), the N-terminal half of Cyt b or the portion of Cyt b corresponding to GSE F21 as indicated below each bar. Cell survival was measured by methylene-blue staining 72 h after infection and is shown relative to the EGFP control (set at 100%). Here and in (E), the error bars indicate standard deviation of three independent experiments. (E) Quantitation of cells with subG1 DNA content. FACS analysis was performed on HeLa cells 72 h after infection with lentiviral Cyt b constructs as in (D). (F) Time course of caspase induction in HeLa cells expressing cytoplasmic Cyt b. Activation of caspases 3/7 and 9/6 were measured using the fluorogenic substrates DEVD-AMC (filled circles) and LEHD-AMC (open circles), respectively. AMC fluorescence resulting from substrate cleavage was measured at the indicated times after substrate addition.

### Cleavage and Release of a C-Terminal Portion of Cyt b into the Cytoplasm During FAS-Mediated Apoptosis

While Cyt b resides inside mitochondria, its corresponding GSE (F21) is expressed in the cytoplasm. To investigate whether GSE F21 has a chance to act against its endogenous target, we monitored the integrity and intracellular localization of Cyt b during FAS apoptosis (Fig. 2A and B).

Mitochondrial and cytoplasmic fractions were prepared from DF-1FAS cells and HeLa cells expressing endogenous FAS at various times after treatment with FAS agonistic antibodies. The quality of separation was assessed by detection of fraction-specific marker proteins (e.g., PCNA as a cytoplasmic marker and GRP75 as a mitochondrial marker). Cyt c was used as a classic example of a protein that undergoes mitochondria-to-cytoplasm translocation during apoptosis. Western blot analysis using P49 antibodies (37) generated against a C-terminal peptide of Cyt b (location of epitope is illustrated in Fig. S2) showed that in untreated (nonapoptotic) cells, Cyt b (30 kDa) was restricted to the mitochondrial fraction. However, 0.5 to 1 h after FAS antibody treatment, a new band corresponding to a 15-kDa protein was detected primarily in the cytoplasmic fraction. Appearance of the 15-kDa Cyt b band in the cytoplasmic fraction was accompanied by a decrease in the intensity of the 30-kDa Cyt b band in the mitochondrial fraction. This profile of Cyt b expression and localization was observed in both chicken (DF-1FAS) and human (HeLa) cells.

Antibodies recognizing an N-terminal epitope of Cyt b also revealed the decrease in full-length Cyt b in the mitochondrial fraction of FAS antibody-treated cells, but failed to identify the 15-kDa cytoplasmic band (data not shown). This result suggests that the 15-kDa protein is the result of cleavage of mitochondrial Cyt b and export of a C-terminal portion (roughly half) to the cytoplasm. The sequence corresponding to GSE F21 lies within this C-terminal portion (see Fig. S2). Neither cleavage nor translocation of Cyt b to the cytoplasm were detected in the same cells (DF-1FAS and HeLa) undergoing apoptosis because of treatment with staurosporin (data not shown), indicating that this phenomenon is not an essential part of any mitochondria-mediated apoptosis and might be specific for FAS.

The identity of the protease involved in Cyt b cleavage during FAS-induced apoptosis, as well as the exact cleavage site, remain to be determined; however, it should be noted that cleavage of Cyt b was not blocked by the proteasomal inhibitor MG-132 (see Fig. 2A).

### Cytoplasmic Expression of Cyt b or the C-Terminal Fragment Cyt b512–1146 Induces Apoptosis

Because Cyt b is encoded by the mitochondrial genome and translated by mitochondrial ribosomes from mitochondrial RNA, we could not use gene knock-out or knock-down techniques to test the role of Cyt b in FAS-induced apoptosis
directly. As an alternative, we tested whether ectopic expression of full-length Cyt b or a C-terminal portion predicted to correspond to the 15-kDa Cyt b fragment (Cyt b1-511) in the cytoplasm could initiate apoptosis in the absence of FAS activation.

Because of differences in the nuclear and mitochondrial genetic codes, we could not use the Cyt b coding sequence taken from mitochondrial DNA for cytoplasmic expression of the protein. The mitochondrial sequence of the Cyt b ORF contains 11 tryptophan residues encoded by codons that are interpreted as stop codons in the nuclear code. Therefore, we generated an artificial human Cyt b gene sequence encoding the same amino acids but using codons of the nuclear code that are most frequently used in the mammalian genome to optimize translation (see Fig. S2A). It should be noted that GSE F21 does not contain any tryptophan codons.

In addition to full-length Cyt b, three Cyt b derivatives were also generated from the recoded synthetic gene (see Fig. S2A). Two of these corresponded to the N- and C-terminal halves of the protein, roughly corresponding to the putative cleavage products of Cyt b generated during FAS-mediated apoptosis (Cyt b1-511 and Cyt b512-1146, respectively). The third encoded the 42 amino acid-long portion of Cyt b corresponding to GSE F21. To simplify delivery of these constructs into avian and mammalian cells, they were cloned into a lentiviral vector. The ability of the constructs to drive expression of FLAG-tagged proteins of the expected sizes in HeLa cells was analyzed by Western blot (Fig. 2C).

The effect of the generated Cyt b expression constructs on the viability of HeLa cells is shown in Fig. 2D. Similar results were obtained in DF-1 cells (data not shown). No cytotoxicity was associated with expression of either the N-terminal portion of Cyt b or the GSE F21-derived sequence. In contrast, expression of either full-length Cyt b or Cyt b512-1146 caused significant cell death, which could be blocked by coexpression of the GSE F21-derived sequence. This confirms that the apoptosis-inhibiting effect of GSE F21 is because of specific interference of the GSE with the biological activity of its cognate gene product, Cyt b. The fact that the original GSE F21 (mitochondrial sequence) and the recoded version (nuclear sequence) were equally effective in suppressing FAS-induced apoptosis indicates that they act as dominant negative peptides rather than inhibitory RNA molecules.

The apoptotic nature of the observed cell death was confirmed by FACS analysis of cellular DNA content, showing that ectopic expression of Cyt b or Cyt b512-1146 in HeLa cells results in appearance of a large population of cells with subG1 DNA content that can be blocked by coexpression of GSE F21 (Fig. 2E, Fig. S3A). The proportion of cells with subG1 DNA content in cell cultures infected with different Cyt b constructs correlated with their cytotoxicity (see Fig. 2D and E). In addition, ectopic expression of Cyt b resulted in caspase activation as illustrated by caspase enzymatic activity in cell lysates (Fig. 2F) and poly(ADP-ribose) polymerase (PARP) cleavage (Fig. S3B). The proapoptotic activity of Cyt b and Cyt b512-1146 was evident in several different cell lines (including PC3 and CWR22R, Fig. S3 C and D).

**Caspase 8 and Bid Mediate Cyt b Cleavage During FAS Apoptosis.**

Caspase 8 activation is known to occur immediately downstream of DISC assembly during FAS-mediated apoptosis (38). To determine whether Cyt b release from mitochondria depends upon caspase 8 activation, we monitored Cyt b content in the mitochondria and cytoplasm of cells treated with FAS agonistic antibodies in the presence or absence of the specific caspase 8 inhibitor Z-IETD-FMK. As expected, Z-IETD-FMK treatment of HeLa cells resulted in inhibition of FAS-induced cell death (Fig. 3A). In addition, we found that the presence of Z-IETD-FMK resulted in reduced Cyt b cleavage (Fig. 3B). Caspase 8 mediates the mitochondrial branch of FAS apoptosis by proteolytic activation of Bid (1). We found that Bid knock-down by specific shRNA not only makes HeLa cells resistant to FAS agonistic antibody treatment, but also prevents Cyt b cleavage (Fig. S4B). Thus, activation of both caspase 8 and Bid is required for Cyt b-dependent apoptosis.

**Bcl-2 and Cyt b-Mediated Apoptosis.** Blockade of caspase activation by the pan-caspase inhibitor Z-VAD resulted in inhibition of Cyt b-mediated apoptosis, as well as FAS-mediated apoptosis (Fig. 3 C and E). This provides further evidence that Cyt b mediates an apoptotic pathway and suggests that it functions in a manner similar to Cyt c, through induction of the caspase cascade upon release into the cytoplasm. To investigate how Cyt b fits into the current paradigm of mitochondrial apoptosis, we analyzed the effects of known regulators of mitochondrial apoptosis, Bcl-2 and Cyt c, on apoptosis induced by FAS-agonistic antibodies or cytoplasmic Cyt
b expression. Bcl-2 is an antiapoptotic member of a large family of proteins that control apoptosis, at least in part, by regulating mitochondrial outer membrane permeability. Activated Bcl-2 counteracts proapoptotic family members and prevents mitochondrial release of apoptogenic factors such as Cyt c, which activates caspases in the cytoplasm (39). Bcl-2 overexpression effectively blocks FAS-induced apoptosis in HeLa cells (see Fig. 3C), indicating that, as for other Type II cells, they die following FAS ligation predominantly via a mitochondria-mediated mechanism. Analysis of endogenous Cyt b in cytoplasmic and mitochondrial fractions of cells overexpressing Bcl-2 following treatment with FAS agonistic antibodies showed that Cyt b cleavage and cytoplasmic release of its C-terminal portion were completely blocked (Fig. 3D). These data suggest that Bcl-2 might block a specific signal originating in the cytoplasm upon FAS activation that is required to trigger mitochondrial events leading to Cyt b cleavage. Bcl-2 could presumably prevent such a signal from passing through the mitochondrial membrane. This hypothesis is supported by our finding that over-expression of Bcl-2 did not alleviate apoptosis induced by ectopic cytoplasmic expression of full-length Cyt b or Cyt b512-1146 (see Fig. 3E).

Cyt c in Cyt b-Mediated Apoptosis. The failure of Bcl-2 over-expression to prevent Cyt b-induced apoptosis suggests that FAS apoptosis mediated by Cyt b might involve a different intrinsic pathway independent of Cyt c/Apaf-1. To test this hypothesis, specific shRNAs were used to knock down expression of Apaf-1 (Fig. 4A) or Cyt c (Fig. 4B) in HeLa cells. Cells were then treated with FAS agonistic antibodies. Knock-down of either component of the apoptosome resulted in a similar significant delay in the rate of FAS-induced cell death; however, the overall extent of cell death 48 h after FAS antibody treatment was the same, regardless of Cyt c or Apaf-1 presence or absence (Fig. 4C). At 4 h after FAS antibody treatment, nearly 90% of cells with intact Cyt c/Apaf-1 were dead, as compared to only ~50% of cells lacking Cyt c or Apaf-1. This could not be accounted for by incomplete gene knock-down, because shCyt c completely blocked apoptosis induced by staurosporine, a known inducer of intrinsic apoptosis (Fig. S5A). These observations indicate that the Cyt c/Apaf-1 pathway can accelerate but is not essential for FAS-mediated apoptosis. In addition, FAS-induced disappearance of Cyt b from mitochondria (see Fig. 4B) and apoptosis induced by cytoplasmic expression of Cyt b were unaffected by knock-down of Cyt c (Fig. 4D). Moreover, expression of GSE F21 increased the proportion of surviving cells to a level that did not depend upon Cyt c knock-down (see Fig. 4C). Taken together, these data suggest that Cyt b acts independently of Cyt c during FAS apoptosis (Fig. S6).

Discussion

In this study, we used a previously unused GSE method based upon RCAS replication competent avian retroviral vectors that allow immediate rescanning of enriched virus subpopulations after each round of functional selection. In addition to expedited screening, this system also offers improved safety, as chicken retroviruses cannot replicate in mammalian cells. The concern of biological differences between avian and mammalian cells is not a serious drawback when the method is applied to evolutionarily conserved pathways.

The RCAS-based GSE method was validated by our isolation of a number of GSEs capable of protecting both chicken and human cells from FAS-mediated apoptosis. The known involvement of several of the identified genes in apoptosis provided confidence in our method and suggested that isolation of a GSE corresponding to Cyt b was meaningful. Our investigation into the relevance of Cyt b to FAS-induced apoptosis revealed that cleavage and cytoplasmic release of a C-terminal portion of Cyt b occurs during FAS-induced apoptosis and that cytoplasmic expression of Cyt b is sufficient to induce apoptosis.

Several lines of evidence indicate that FAS-induced cleavage of Cyt b occurs inside the mitochondria before its release into the cytoplasm. First, ectopically expressed cytoplasmic Cyt b was not cleaved upon treatment with FAS agonistic antibodies (data not shown). Second, in our cell fractionation experiments full-length Cyt b was only observed in mitochondrial fractions. In addition, our finding that cytosolic cytochrome c expression of either full-length Cyt b or Cyt b512-1146 induced apoptosis supports the idea that intramitochondrial cleavage of Cyt b is needed not for activation of its proapoptotic activity, but rather for its release from the mitochondria. If cleavage of Cyt b is a prerequisite for its release to the cytoplasm, then the antiapoptotic activity of GSE F21 might be because of its acting as an inhibitor of Cyt b cleavage. However, although GSE F21-expressing cells are resistant to FAS-dependent apoptosis, FAS antibody treatment still results in cleavage and cytoplasmic release of Cyt b in these cells (data not shown). This, together with the proapoptotic effect of cytoplasmic Cyt b and inhibition of this effect by GSE F21, indicate that the GSE blocks some proapoptotic cytoplasmic function of Cyt b (see below).

Our data suggest the following hypothetical sequence of events occurring upon FAS activation (see Fig. 4): Caspase 8 activation following DISC formation results in generation of a signal of unknown nature that triggers intra-mitochondrial cleavage of Cyt b. This signal is Bid-dependent and can be blocked by over-expression of Bcl-2, suggesting that it involves a factor passing through the mitochondrial pore. It is unlikely that Bcl-2 acts at the point of the Cyt b C terminus leaving the mitochondria, because Bcl-2 over-

Fig. 4. Limited dependence of FAS apoptosis on Cyt c and Apaf-1. (A) RT-PCR detection of Apaf-1 and actin RNA in HeLa cells expressing shRNAs targeting Apaf-1 or EGFP (control). (B) Suppression of Cyt c does not affect Cyt b processing during FAS apoptosis. HeLa cells expressing shCyt c or shLuc (negative control) were left untreated (nt) or treated with FAS agonistic antibody and CHI (Fas ab). Western blot analysis of Cyt b and Cyt c was performed on mitochondrial (Upper) and cytoplasmic (Lower) fractions 8 h after treatment. Cytochrome Oxidase Subunit IV (OK Phos Comp IV) and β-actin were used as mitochondrial and cytoplasmic markers, respectively. (C) Time course of FAS apoptosis in control HeLa cells and HeLa cells expressing the indicated constructs. Methylene-blue staining was performed at the indicated times after exposure to FAS antibody plus CHI. Here and in (D), three independent experiments were performed; error bars indicate standard deviations. (D) Knock-down of Cyt c does not prevent Cyt b-induced apoptosis. Methylene-blue staining was performed in control HeLa cells and HeLa cells expressing cytoplasmic Cyt b or Cyt c (shCyt c) or both. Survival was measured 3 days after lentiviral infection.
expression prevents Cyt b cleavage. The C-terminal half of cleaved Cyt b then appears in the cytoplasm. Given the high hydrophobicity of Cyt b and its tight binding to the inner mitochondrial membrane (27), it will be important to identify mediators of this process. One hypothesis regarding the mechanism of action of the C-terminal Cyt b peptide is that it induces activation of the caspases that execute apoptosis through interaction with a cellular factor, a process which might be blocked by GSE F21. The identity of such cellular mediators of Cyt b-induced caspase activation and of the protease responsible for Cyt b cleavage remains to be determined. These are likely to be unique factors, as (i) the classic Cyt c/Apaf-1 apoptosis is not responsible for Cyt b-induced apoptotic function, and (ii) there are no conventional protease cleavage sites recognizable within Cyt b (although there is a cluster of three overlapping imperfect caspase-recognition sites in the middle of Cyt b) (see Fig. S2). Complete elucidation of the Cyt b apoptosis pathway will be important to resolve the apparent discrepancy between the near complete block of FAS apoptosis observed with Bcl-2 over-expression (see Fig. 3C) and the lack of complete inhibition seen upon knock-down of Cyt c or Apaf-1 (see Fig. 4), suggesting that FAS signaling only partially goes through the canonical Cyt c path.

Preliminary observations (see Fig. S4 A and B) suggest that a similar Cyt b-mediated mechanism is involved in apoptosis initiated by TRAIL and TNFα, thus extending the generality of the described phenomenon beyond FAS apoptosis to all death receptor-mediated programmed cell death.

Materials and Methods

GSE Library Screening. To convert library DNA into viral stock, 2.5 × 10⁵ DF-1 cells were plated on each of five 15-cm plates and transfected the next day with 50-μg library DNA per plate using Lipofectamine Plus (Invitrogen). A plate of DF-1 cells was transfected with empty RCASBP(A) vector for use as a negative control. Transfection conditions were optimized using an RCASBP(A)-based GFP expression construct to achieve virus titers of at least 10⁶ IU/ml. The virus from each of the five library-transfected plates was kept separate, generating five parallel experimental branches. DF-1 cells were seeded at 2 × 10⁵ cells per plate (i.e., 2 × 10⁴ cells per well) were infected with library virus (five plates) or negative control virus (one plate) overnight with 4-μg/ml polybrene. Two days later, when the cells were nearly confluent (10⁷ cells per plate), they were treated overnight with FAS agonistic antibody (1 μg/ml). Cells were allowed to recover for 1 week and those surviving formed clonal colonies (dead cells were removed by aspiration during medium changes). Virus-containing media collected from the pooled clones surviving from each plate was used to infect fresh DF-1 cells. The secondary and tertiary selections were performed as above, except in a six-well plate format (2 × 10⁵ cells per well). Five wells were each infected with one of the five independent virus subpopulations generated by the primary selection and one well was infected with negative control virus (original virus stock). Two days later, the nearly confluent cells (about 10⁶ cells/well) were again subjected to FAS antibody selection. Following tertiary selection, surviving clones from each well were pooled and expanded. Total RNA was isolated and used for RT-PCR with a primer specific for the adaptor flanking each library insert: 5′-AAATCATGATCATGGGT-CATGGTCTAGG-3′. Recovered GSES were sequenced and identified using NCBI BLAST.

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chicken p53, myc, and relative abundance of different mRNA sequences. We used the Southern hybridization according to standard protocol for the PCR amplified, blotted onto nylon membranes, and analyzed by chromatography. The aliquots of isolated ds- and ss-DNA were normalized (2). For this, aliquots of library DNA were denatured and renanned for different time intervals. DNA from each time point was subjected to HAP (hydroxylapatite) chromatography. The aliquots of isolated ds- and ss-DNA were PCR amplified, blotted onto nylon membranes, and analyzed by Southern hybridization according to standard protocol for the relative abundance of different mRNA sequences. We used the chicken p53, myc, and β-actin cDNAs as reference probes for low, middle, and high abundance messages, respectively. The fraction that contained similar proportions of those cDNA sequences was used for library preparation. The pool of normalized cDNA fragments was digested with Clai and ligated into the Clai site of the RCASBP(A) vector. The complexity of the obtained library was ~10^6 clones, sufficient for isolation of cDNAs (GSEs) corresponding to most genes expressed.

Cells, Cell Culture, and Induction of Apoptosis. The immortalized CEF cell line DF-1 was obtained from the American Type Culture Collection (ATCC number CRL-12203) and cultured in Dulbecco’s modified minimal essential medium (DMEM), supplemented with 2-mM glutamine, 100-U/ml penicillin and streptomycin, and 10% FCS (FCS, Life Technology). HeLa, 293, PC3, and CWR22R cells were cultured in DMEM supplemented with 2-mM glutamine, 100-U/ml penicillin and streptomycin, and 10% FBS (FBS, Life Technology). The purified stock of FAS agonistic antibody [hybridoma IPO-4 (3)] had a concentration of 1 mg/ml and was used at a final concentration of 1 µg/ml unless indicated otherwise. CHI (0.5 µg/ml) was used in all FAS treatment experiments involving human cells (it was not used with avian cells) by addition along with FAS antibody to a final concentration of 1 µM. CHI blocks protein synthesis, and at this concentration is presumed to counter the antiapoptotic effect of NF-κB activation during receptor-mediated apoptosis. We found that in HeLa cells, FAS-induced apoptosis was much more efficient in the presence of CHI (data not shown). Staurosporine was used at a concentration of 0.1 µM. The proteasomal inhibitor MG-132 was used at concentration of 10 µM (Calbiochem). The caspase inhibitors Z-VAD-FMK (Sigma, V116) and Z-IETD-FMK (Sigma, C1230) were used at concentration of 10 µM. Caspase 9 inhibitor Z-LEHD-FMK (Sigma, C1355) was used at concentration of 10 µM. Recombinant TRAIL was used at a final concentration of 0.1 µg/ml (a kind gift of Janet Houghton, Cleveland Clinic, Cleveland, OH). TNF-α was used at a final concentration of 0.5 µg/ml (Peprotech, cat # 300-01A). TRAIL and TNF-α were used in combination with CHI (0.5 µg/ml).

Generation of Cyt b-Encoding cDNA in a Nuclear Code. The artificial (nuclear coded for cytoplasmic expression) Cyt b construct was synthesized by GENEART. The Cyt b gene sequence was optimized using Proprietary GeneOptimizer software, allowing optimal expression. The nuclear recoded Cyt b was supplied with a FLAG tag sequence at its C terminus. Briefly, synthesis of the artificial gene by GENEART involved generation of DNA oligonucleotides covering the entire gene sequence. The oligonucleotides are annealed to form a full-length contig, the gaps in which are filled by DNA polymerase followed by ligation and cloning into a GENEART-provided shuttle vector. Upon receiving the construct from GENEART, the insert was recloned into the pLM-CMV-PL3 vector, a kind gift of Dr. Peter Chumakov (Cleveland Clinic, Cleveland, OH). The inserts were amplified by PCR with specific primers, digested with appropriate restriction enzymes, and cloned under the CMV promoter. All shRNA constructs were prepared using the pLSP lentiviral vector containing the puromycin resistance gene. Expression of shRNA was driven by the H1 promoter.

Supporting Information

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SI Materials and Methods

Chicken GSE Library. The GSE cDNA library was provided by Elena Feinstein (Weizmann Institute of Science, Rehovot, Israel). It was constructed according to provided guidelines (1) but with some modifications. PolyA plus RNA was prepared from chick embryo fibroblast (CEF) cells (7-day embryos, around 35 population doublings). To achieve equal representation of 5' and 3' mRNA sequences in random fragment cDNA preparation, cDNA synthesis was carried out on fragmented mRNA using random oligonucleotides as primers. The resulting cDNA was ligated to the “universal linker” shown in Fig. S1B. cDNA fragments ranging from 100 to 400 bp were selected by agarose gel fractionation for further PCR amplification using the sense oligonucleotide of the universal linker. To achieve equal representation of different abundance, the library was normalized (2). For this, aliquots of library DNA were denatured and renanned for different time intervals. DNA from each time point was subjected to HAP (hydroxylapatite) chromatography. The aliquots of isolated ds- and ss-DNA were PCR amplified, blotted onto nylon membranes, and analyzed by Southern hybridization according to standard protocol for the relative abundance of different mRNA sequences. We used the chicken p53, myc, and β-actin cDNAs as reference probes for low, middle, and high abundance messages, respectively. The fraction that contained similar proportions of those cDNA sequences was used for library preparation. The pool of normalized cDNA fragments was digested with Clai and ligated into the Clai site of the RCASBP(A) vector. The complexity of the obtained library was ~10^6 clones, sufficient for isolation of cDNAs (GSEs) corresponding to most genes expressed.

Vectors and Constructs. The RCASBP(A) and RCASBP(B) vectors (4) were a kind gift from Stephen H. Hughes (National Cancer Institute, Frederick MD). pBSfi I, a modified version of pBluescript in which the multicloning site is flanked by two SfiI sites, was a kind gift from Masa Aoki (The Scripps Research Institute, La Jolla, CA). RCASBP(A)SfiI and RCASBP(B)SfiI were prepared by inserting the SfiI adaptor with multiple cloning sites from pBSfi I into the Clai site of the RCASBP(A) and (B) vectors. For expression of human FAS in chicken DF-1 cells, the full-length human FAS cDNA clone was obtained from Invitrogen. The FAS ORF was cloned into the pBSfi vector. It was then cut out with SfiI and directionally cloned into RCASBP(B)SfiI. For generation of the chicken GSE library, GSE inserts (see Chicken GSE Library section, above) were cloned into the Clai site of RCASBP(A). Recovered individual GSEs, as well as all other constructs, were prepared using the RCASBP-P(A)SfiI vector. Recovered individual GSEs were amplified by PCR using oligonucleotide primers specific for the corresponding genes, cloned into pBSfi I, and subsequently transferred into the RCASBP(A)SfiI vector. Sense and antisense constructs were identified by sequencing. All lentiviral constructs for overexpression experiments were prepared using the pLM-CMV-PL3 vector, a kind gift of Dr. Peter Chumakov (Cleveland Clinic, Cleveland, OH). The inserts were amplified by PCR with specific primers, digested with appropriate restriction enzymes, and cloned under the CMV promoter. All shRNA constructs were prepared using the pLSP lentiviral vector containing the puromycin resistance gene. Expression of shRNA was driven by the H1 promoter.

Retroviral and Lentiviral Transduction. All chicken retroviral transductions using RCASBP(A)- and (B)-based constructs were done as follows: plasmid DNA was transfected into DF-1 cells according to the standard Lipofectamin Plus (Invitrogen) pro-
tocol (10 μg of plasmid DNA per 10 cm plate; 50 μg per 15 cm plate). The virus was allowed to spread in the cell population for 4 days in the presence of +4/μg/ml polybrene. The culture supernatant was then collected and used to infect a fresh population of cells. For lentiviral transduction, human A293 kidney cells (BD Biosciences Clontech) were plated subconfluently on a 10-cm tissue culture plate. The next day, the cells were transfected with 3.3-μg lentiviral vector DNA, 3.3-μg pVsV-G expression plasmid and 3.3-μg packaging vector pCMV-DeltaR8.2 (kind gifts of Dr. Peter Chumakov, Cleveland Clinic, Cleveland, OH) using Lipofectamine Plus (Invitrogen) according to the manufacturer’s recommendations. The culture medium was changed 16 h after transfection and the virus-containing medium was collected either 48 or 72 h after transfection. The medium-containing virus was then used to infect target cells in the presence of polybrene (4 μg/ml, Sigma). Neither the RCASBP vectors nor the lentiviral vectors contained selectable markers. The efficiency of infection was monitored by RT-PCR or Western blot analysis with specific antibodies to expressed proteins. All shRNA constructs were designed using Dharmacon Web site software. Oligonucleotides corresponding to sense and antisense strands of the shRNA were ordered through Integrated DNA Technologies, Inc. 5’-phosphorylated shRNA oligos were PAGE-purified, annealed, and cloned into the pLSLP lentiviral vector (a kind gift of Dr. Peter Chumakov, Cleveland Clinic) under the H1 promoter. HeLa cells transduced with pLSLP-based shRNA constructs were selected with puromycin (1 μg/ml) for at least 3 days. Down-regulation of the targeted genes was assessed by RT-PCR with specific primers or by Western blotting with specific antibodies.

**Cell Viability and Apoptosis Assays.** Methylene-blue staining of cells remaining attached to the plate after induction of apoptosis was performed as an initial assessment of cell viability. Floating cells were removed from plates and the remaining attached cells were fixed with methanol for 15 min and stained with methylene-blue (1% solution in PBS) for 1 h. Plates were then rinsed with water to remove excess dye and allowed to dry completely. Cell numbers were quantitated by methylene-blue elution and spectrophotometric measurement at 630 nm using a Beckman Coulter DU8000 instrument. The dye was eluted from dried plates with 0.1 N HCl at room temperature. For caspase-activation assays, HeLa, PC3, and CWR22R cells were plated in 96-well plates at 7,000 cells per well in DMEM/10% FCS and treated 24 h after plating with lentiviral isolates (where indicated, in the presence of polybrene). The medium-containing virus was then used to infect target cells in the presence of polybrene (4 μg/ml, Sigma). Neither the RCASBP vectors nor the lentiviral vectors contained selectable markers. The efficiency of infection was monitored by RT-PCR or Western blot analysis with specific antibodies to expressed proteins. All shRNA constructs were designed using Dharmacon Web site software. Oligonucleotides corresponding to sense and antisense strands of the shRNA were ordered through Integrated DNA Technologies, Inc. 5’-phosphorylated shRNA oligos were PAGE-purified, annealed, and cloned into the pLSLP lentiviral vector (a kind gift of Dr. Peter Chumakov, Cleveland Clinic) under the H1 promoter. HeLa cells transduced with pLSLP-based shRNA constructs were selected with puromycin (1 μg/ml) for at least 3 days. Down-regulation of the targeted genes was assessed by RT-PCR with specific primers or by Western blotting with specific antibodies.

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For amplification of human FAS from the Invitrogen clone for further cloning, the following primers were used:

- hFAS-5’-Xba: 5’-CCG TCT AGA GAT TGC TCA ACA ACC ATG CTGGGC A-3’
- hFAS-3’Sal: 5’-CCG GTC GAC CTA GAC CAA ACT TTG GAT TTCATT T-3’

For generation of Cyt b derivatives (C-terminal portion, N-terminal portion and GSE) from the full-length artificial (nuclear recoded) Cyt b construct, the following primers were used for amplification:

- FLAG-Cyt b-Xba-5’
- 5’-cg ccg tct aga gcc acc atg tat cag gat gac gat gct atg gga ggc tac tcag tag aca gtt-3

Cyt b-Sal-3: 5’-ccg ccg gtc gac tca att tga gta ttt ttt cca caa tca-3’

- FLAG-Cytb-GSE-5’: 5’-CC GCC ACC ATG GAT TAC AAG GAT GAC GAC GAT AAG GAT GGC CTC ATC CCC ACC AAC-3’

- Cytb-GSE-3’: 5’-C TCA TGG GTC ATC GGG GAG-3’

Gene-specific primers used for amplification of recovered GSEs are available upon request.

**Western Blot Analysis and Antibodies.** Cells were lysed in Nonidet P-40 lysis buffer (50-mM Tris, pH 8.0; 150-mM NaCl; 1% Nonidet P-40) containing protease inhibitor mixture at a 500-fold dilution (Sigma) for 30 min on ice. The lysates were centrifuged at 14,000 × g at 4°C and the supernatant was collected and used for Western blot analysis. The protein concentration was measured using DC Protein Assay kit (Bio-Rad). Equal amounts of protein were electrophoresed on gradient 4 to 20% precast gels (Novex). For immunoblotting, the proteins were transferred to a PVDF membrane (Perkin-Elmer Life Sciences) by electroblotting and the membranes were blocked in 5% milk in Tris-buffered saline with Tween 20 (TBST; 20-mM Tris-Cl (pH 8.0), 150-mM NaCl, 0.1% Tween 20), probed with the indicated primary antibody in TBST, and visualized by enhanced chemiluminescence (Perkin-Elmer Life Sciences). All primary antibodies were used at a dilution of 1:200. The goat anti-rabbit and anti-mouse HRP-conjugated secondary antibodies (Santa Cruz Biotechnology) were used at a dilution of 1:10,000. The immunoblots were stripped [4% SDS, 62.5-mM Tris-HCl (pH 6.8), 100 mM 2-mercaptoethanol] at 55°C for 45 min before reprobing. The following primary antibodies were used:

- anti-Cyt b rabbit polyclonal P-49 raised against the C-terminal half of the protein was described earlier (5), anti-FLAG M2 mouse monoclonal (Sigma), anti-caspase 8 SC-56070 mouse monoclonal IgG (Santa Cruz Biotechnology), anti-PARP-H-250 rabbit polyclonal (Santa Cruz Biotechnology), anti-GRP75 C-19 goat polyclonal (Santa Cruz Biotechnology), anti-PCNA F-2 mouse monoclonal (Santa Cruz Biotechnology), anti-cytochrome c mouse monoclonal (BD Pharmingen, catalogue number 556433), anti-hFAS C-20 rabbit polyclonal (Santa Cruz Biotechnology), anti-β-actin HRP-conjugated (Sigma A3854), and anti-OxPhos comp I-IV mouse monoclonal (Molecular Probes, 20E8).

**Cell Fractionation and Isolation of Mitochondria.** Mitochondrial and cytoplasmic cell fractions were prepared as described in detail elsewhere (Benchtop Mitochondria Isolation Protocol, Mitosciences, Inc.). In brief, cells were plated on 10-cm plates and treated as described. At the end of the experiment, the floating apoptotic cells were collected from the media and the remaining
attached cells were collected by trypsinization. The floating and attached cells were pooled, washed several times with PBS, and incubated in 1-ml hypotonic buffer supplemented with protease inhibitors for 15 min (10-mM NaCl, 1.5-mM MgCl₂, 10-mM Tris-HCl, pH 7.5, Protease Inhibitor Mixture, Sigma). Cell membranes were disrupted by mechanical shearing (2-ml Dounce homogenizer), and 400 μl of Sucrose 2.5× MS Buffer was added (2.5× MS Buffer: 525-mM Mannitol, 175-mM Sucrose, 125-mM Tris-HCl, pH 7.5, 2.5 mM EDTA, pH 7.5). The cellular lysate containing mitochondria was separated from cell debris and nuclei by centrifugation at 500 × g. The supernatant was then subjected to sucrose gradient centrifugation at 30,000 × g. The resulting mitochondrial pellet was resuspended in RIPA buffer supplemented with protease inhibitors and the supernatant containing the cytoplasmic fraction was directly loaded onto SDS/PAGE gels for Western blot analysis. All reactions were performed on ice and all centrifugation steps were performed at 4°C. Protein concentrations were determined for equal loading.

A

Fig. S1. (A) Scheme of GSE library generation in RCASBP(A) vector (details in text). (B) Structure of GSE adaptors, carrying three start ATG codons, three stop TGA codons, and Kozak sequence (underlined).
According to our results, the C-terminal part of Cyt b is cleaved and released into the cytoplasm. The positions of the GSE sequence and the P-49 antibody epitope are indicated.

Fig. S2. (A) Scheme of generated artificial (nuclear recoded) Cyt b construct showing its humanized DNA sequence based on the protein amino acid content. This construct was also supplied with FLAG sequence at the C-terminal. GSE-, C-terminal, and N-terminal parts are shown. White box indicates a cluster of overlapping imperfect caspase-recognition sites. (B) According to our results, the C-terminal part of Cyt b is cleaved and released into the cytoplasm. The positions of the GSE sequence and the P-49 antibody epitope are indicated.
Fig. S3.  (A) FACS analysis of the cell cycle of HeLa cells overexpressing Cyt b. Cells were collected in 48 h after infection with lentiviral constructs, and their DNA content was measured by staining with propidium iodide. Cyt b expression induces apoptosis as judged by the appearance of a population of cells with subG1 DNA content. (B) Western blot analysis of apoptosis-specific PARP cleavage during initiation of cell death by Cyt b. (C) Cyt b induces caspase activation when overexpressed in the cytoplasm of PC3 and CWR22R cells. The cells were treated with lentiviral supernatants in the range from undiluted to 1:64, and cell death was measured by methylene-blue staining of surviving cells 48 h after infection. (D) Cyt b induces caspase-dependent apoptosis when overexpressed in the cytoplasm of PC3 and CWR22R cells. The cells were treated with lentiviral supernatants in the range from undiluted to 1:64. At 48 h after infection the supernatant was replaced with DEVD-AMC to measure caspase 3/7 activation.
Fig. S4. Cyt b cleavage and translocation to the cytoplasm is a common property of death receptor-mediated apoptosis induced by different ligands. (A) HeLa cells were treated with FAS, TRAIL, or TNFα (all in combination with 1 μg/ml cycloheximide) as described in SI Materials and Methods. Survival was measured by methylene-blue staining 24 h after treatment. Bars represent averages of three independent experiments, with the error bars indicating standard deviations. (B) All three receptor-mediated apoptotic ligands induce cleavage and translocation of Cyt b to cytoplasm. Mitochondrial and cytoplasmic fractions were prepared from HeLa cells, untreated (nt) or incubated with FAS, TRAIL, or TNFα at the indicated times of treatment. Fractions were analyzed by Western blotting for Cyt b (left panel: full-length, 30 kDa; right panel: cleaved, 15 kDa), GRP75 (mitochondrial marker), PCNA (cytoplasmic marker), and Cyt c. Asterisks indicate high statistical significance (P < 0.05).
Fig. S5. (A) Down-regulation of Cyt c blocks staurosporine-induced apoptosis. HeLa cells were infected with lentiviruses directing expression of shCyt c or shLuc (negative control) and left untreated (nt) or treated with staurosporine (stau, 0.1 μM). Cell survival was measured on day 3 after treatment. (B) siRNA-mediated knockdown of Bid completely blocks FAS apoptosis in HeLa cells. siRNA against Bid (a mixture of five synthetic gene-specific siRNAs) and control scrambled siRNA was obtained from Dharmacon and applied to the cells according to manufacturer’s recommendations. Three days later cells were treated with FAS agonistic antibodies and cell survival was measured 24 h later by methylene-blue staining. The quantification data represent averages of three independent experiments, with the error bars indicating standard deviations. Asterisks indicate high statistical significance (P < 0.05).
Fig. S6. Scheme of Cyt b action in FAS-induced apoptosis. FAS engagement at DISC induces a signal leading to activation of Bid and mitochondrial Bax/Bak pore opening. In addition, a second signal originates in the cytoplasm, which results in Cyt b inducing caspase-dependent apoptosis and is downstream of proapoptotic factors released from mitochondria (details in text).
Table S1. List of isolated GSEs

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<th>GSE</th>
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<th>Chicken gene name</th>
<th>Similarity to human gene, %</th>
<th>Known relation to apoptosis</th>
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