Newcastle Disease Virus Exerts Oncolyis by both Intrinsic and Extrinsic Caspase-Dependent Pathways of Cell Death

Subbiah Elankumaran, Daniel Rockemann, and Siba K. Samal*

Virginia-Maryland Regional College of Veterinary Medicine, University of Maryland, College Park, Maryland 20742

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Newcastle disease virus (NDV), an avian paramyxovirus, is tumor selective and intrinsically oncolytic. Here, we present evidence that genetically modified, recombinant NDV strains are cytotoxic to human tumor cell lines of ecto-, endo-, and mesodermal origin. We show that cytotoxicity against tumor cells is due to multiple caspase-dependent pathways of apoptosis independent of interferon signaling competence. The signaling pathways of NDV-induced, cancer cell-selective apoptosis are not well understood. We demonstrate that NDV triggers apoptosis by activating the mitochondrial/intrinsic pathway and that it acts independently of the death receptor/extrinsic pathway. Caspase-8-methylated SH-SYSY neuroblasticoma cells are as sensitive to NDV as other caspase-8-competent cells. This demonstrates that NDV is likely to act primarily through the mitochondrial death pathway. NDV infection results in the loss of mitochondrial membrane potential and the subsequent release of the mitochondrial protein cytochrome c, but the second mitochondrion-derived activator of caspase (Smac/DIABLO) is not released. In addition, we describe early activation of caspase-9 and caspase-3.

In contrast, cleavage of caspase-8, which is predominantly activated by the death receptor pathway, is a TNF-related, apoptosis-inducing ligand (TRAIL)-induced late event in NDV-mediated apoptosis of tumor cells. Our data, therefore, indicate that the death signal(s) generated by NDV in tumor cells ultimately converges at the mitochondria and that it acts independently of the death receptor pathway. Our cytotoxicity studies demonstrate that recombinant NDV could be developed as a cancer virotherapy agent, either alone or in combination with therapeutic transgenes. We have also shown that trackable oncolytic NDV could be developed without any reduction in oncolytic efficacy.

Naturally occurring or engineered oncolytic viruses (OVs) are emerging as novel tools for selective growth in and killing of a variety of tumor cells. OVs are multimodality therapeutics that can be engineered to have the tumor specificity of a small molecule, the potent cell killing of a chemotherapeutic, the ability to arouse the host immune system against tumor antigens, and an innate capacity to stimulate the production of host cytokines that have potential anticancer activity (3, 28, 36). Among the OVs, Newcastle disease virus (NDV) is considered to be a very promising oncolytic agent (34–35, 44, 47). NDV has been in a clinical setting as an experimental oncolytic agent for more than 30 years (10, 51). Among the OVs, Newcastle disease virus (NDV) is considered to be a very promising oncolytic agent (34–35, 44, 47).

NDV contains a single-stranded, negative-sense, nonsegmented RNA genome and belongs to the genus Avulavirus in the family Paramyxoviridae (37). The genomic RNA is 15,186 nucleotides in length (27) and contains six genes that encode at least seven proteins (50). The envelope of NDV contains two glycoproteins, the hemagglutinin-neuraminidase (HN) and fusion (F) proteins: the HN protein mediates attachment of the virus to the cell, and the F protein mediates fusion of the viral envelope with cellular membranes (8). In common with other paramyxoviruses, NDV produces two additional proteins, V and W, from the P gene by alternative mRNAs that are generated by RNA editing (50).

Although much preclinical work establishing that NDV could serve as a cancer therapeutic has been carried out, the mechanisms governing cytotoxicity remain to be fully characterized. NDV strains are known to evoke cellular apoptosis (30, 58). The reported apoptosis signaling pathways induced by NDV in tumor cells are inconsistent and conflicting. It has been suggested that tumor necrosis factor alpha (TNF-α) might be involved in the tumoricidal activity of NDV-activated murine macrophages and human peripheral blood mononuclear cells (36, 65). It is also claimed that the cell-to-cell contact killing of tumor cells by NDV-stimulated macrophages is mediated by TNF-related, apoptosis-inducing ligand (TRAIL) (58). Most of these studies tested the apoptotic response of NDV-activated human cells on human tumor cells. On the other hand, direct infection studies with NDV strain MTH/68 in PC12 rat pheochromocytoma cells indicated that major mitogen-activated protein kinase pathways (including the stress-inducible c-Jun N-terminal kinase pathway and the p38 pathway) or mechanisms regulated by reactive oxygen species have no role in virus-induced apoptotic cell death (12, 52).

Apoptosis is a multistep, multipathway cell death program that is inherent in every cell of the body. The apoptotic pathways leading to cell death can generally be divided into two nonexclusive signaling cascades involving death receptors (extrinsic pathways) or mitochondria (intrinsic pathway) (21). In both pathways, cysteine aspartyl-specific proteases (caspases) that cleave cellular substrates are activated, and this leads to the biochemical and morphological changes characteristic of apoptosis. The death receptor family (the TNF receptor superfamily) includes CD95 (Fas/APO-1), TNF-R1, DR3, DR4 (TRAIL-R1), and DR5 (TRAIL-R2) receptors (20, 25, 39). Apoptosis initiated via death receptors involves the adaptor molecule Fas-associating protein with a death domain.

* Corresponding author. Mailing address: Virginia-Maryland Regional College of Veterinary Medicine, University of Maryland, College Park, MD 20742. Phone: (301) 314-6813. Fax: (301) 314-6855. E-mail: ssamal@umd.edu.
brane potential, followed by release of cytochrome c, second mitochondrion-derived activator of caspase (Smac) or direct inhibitor of apoptosis binding protein with low pI (DIABLO) (11, 56), apoptosis-inducing factor, and endonuclease G in the cytosol. Cytosolic cytochrome c triggers the formation of a multimeric Apaf-1/cytochrome c/dATP/procaspase-9 protein complex termed the apoptosome and leads to the activation of caspase-9 (21). Caspase activation and the activity of already active caspases can be inhibited by the inhibitor of apoptosis proteins (IAPs). In turn, IAPs can be inactivated by Smac/DIABLO or Omi/HtrA2, two regulatory proteins that are released from mitochondria. Both the intrinsic and the extrinsic pathways converge on downstream “executor” caspases, mainly caspase-3 and caspases-6 and -7, which are responsible for the cleavage of structural cytoplasmic and nuclear proteins, with consequent cell collapse and death (46). Activation of the death receptor pathway and that of the mitochondrion-associated death pathway are not mutually exclusive, and these pathways may interact (cross talk) at many levels.

With the availability of the reverse genetics system for NDV, it is now possible to manipulate the genome of NDV, engineer additional genes, and retarget the virus to specific receptors (4, 18, 26). It also affords a viable system for studying the mechanistic basis of oncolysis by NDV. To this end, we tested several recombinant NDVs (rNDVs), with and without transgenes, and studied their mechanisms of apoptotic cell death in a variety of tumor cell lines. This study has incorporated the following NDV strains with different properties: (i) a strain that is low pathogenic to the natural host (chickens) that replicates only in the presence of exogenous trypsin in cell culture, (ii) a low-pathogenic strain with a mutation in the F protein to be unable to interact with gangliosides Gb3 and Gb4, (iii) a moderately pathogenic strain with intact interferon (IFN)-antagonistic function, (iv) a moderately pathogenic strain with a defect in IFN antagonism, and (v) a moderately pathogenic strain with an enhanced green fluorescent protein (EGFP) insert. These strains were chosen to detect the differences in cell death pathways induced by lowly pathogenic and moderately pathogenic NDV strains in tumor cells. The IFN-sensitive virus lacks the expression of the antia apoptotic and IFN-antagonistic V protein (19, 42) and replicates only in cells that lack a functional IFN system. The rNDV that expresses EGFP was used to proved that oncogenic rNDV is a trackable virus in vivo and could be exploited to study the mechanism of cell death in vitro.

We found that rNDV mediates oncolysis by the direct induction of apoptosis through multiple caspase-dependent and interferon-independent pathways. We demonstrate here that NDV acts independently of death receptor signaling to induce apoptosis. NDV primarily triggers the activation of the intrinsic mitochondrial death pathway, leading to programmed cell death. Additionally, we have demonstrated that engineering a reporter gene in rNDV to track the virus in vivo does not compromise oncolytic efficacy and can serve as an aid to understanding the mechanistic basis of oncolysis. We have also shown that IFN-sensitive rNDV that replicates in tumor cells, but not in normal cells, is a better oncolytic agent than its parental type.

Materials and methods

Cells. DF1 chicken embryo fibroblast, HeLa, HepG2, CaCo2, and HuTu80 cells were grown in Dulbecco’s modified Eagle’s medium (DMEM) with 10% fetal calf serum (FCS), 100 μg/ml penicillin, and 0.1 μg/ml streptomycin (In-vitrogen, Grand Island, NY). TS4 colon cancer and SH-SY5Y neuroblastoma cells were grown in a 1:1 mixture of DMEM and Ham’s F12 medium with 10% FCS and antibiotics. THP-1, CCRF-CEM, PC3, SW 620, MCF-7, Colo 205, HT29, and HT1080 cells were grown in RPMI 1640 medium supplemented with 10% FCS and antibiotics. The cells were grown at 37°C with 5% CO2 in a humidified incubator.

Viruses. The plasmid pNDVII expressing the full-length antigenome of moderately pathogenic NDV strain Beaudette C (BC) and avirulent NDV strain LaSota have been described previously (18, 26) and were used to construct mutants, chimera viruses, or viruses with additional transgenes. The construction and recovery of the P gene editing mutant (rBC-Edit), recombinant LaSota (rLaSota), and recombinant LaSota with a Virulent Fusion protein cleavage site (rLaSota V.F.) have been described in detail elsewhere (18–19, 41). The plasmid pNDVNDV/NDV Beaudette C expressing the full-length antigenome of NDV was used to construct the P gene editing mutant. The Ascl–SacII fragment containing the P gene editing site from pNDVII was subcloned into pGEM-TZ (+) (Promega, Madison, Wis.) between XbaI and HindIII for the construction of a low-pathogenic strain with a mutation in the F protein to be unable to interact with gangliosides Gb3 and Gb4, and (v) a moderately pathogenic strain with intact interferon (IFN) antagonism, and (vi) a moderately pathogenic strain with a defect in IFN antagonism, and (v) a moderately pathogenic strain with an enhanced green fluorescent protein (EGFP) insert. These strains were chosen to detect the differences in cell death pathways induced by lowly pathogenic and moderately pathogenic NDV strains in tumor cells. The IFN-sensitive virus lacks the expression of the antia apoptotic and IFN-antagonistic V protein (19, 42) and replicates only in cells that lack a functional IFN system. The rNDV that expresses EGFP was used to proved that oncogenic rNDV is a trackable virus in vivo and could be exploited to study the mechanism of cell death in vitro.

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polyacrylamide gel electrophoresis and transferred to a nitrocellulose membrane for immunoblotting. Blots were then probed with the specific antibodies. Proteins were visualized using the trimethylbenzidine membrane substrate (Kirkegaard and Perry Laboratories, Gaithersburg, MD). The following antibodies were used for immunoblotting: Stat1a (sc-345; Santa Cruz Biotechnology, Inc., Santa Cruz, CA), Stat2 (sc-476; Santa Cruz), mononuclear anti-human TRAIL/TNFSF10 (MAB687; R&D Systems, Minneapolis, MN), anti-cytochrome c (sc-13561; Santa Cruz), NDV-MCA (mononuclear antibody cocktail directed against the NDV HN protein) (31), and actin (sc-8432; Santa Cruz).

**WST-1 assay.** A colorimetric assay for analyzing cell viability is based on the cleavage of the tetrazolium salt WST-1 by mitochondrial dehydrogenases. In each experiment, the test cell line was seeded into 96-well plates at 1 × 10⁴ cells/well in growth medium (DMEM + 10% FCS or RPMI + 10% FCS; Invitrogen). Following overnight incubation (37°C and 5% CO₂), media were removed by aspiration, and to each well 20 µl of virus-containing OptiMEM (Invitrogen) was added, with the MOI ranging in 10-fold increments from 0.01 to 10 or with no virus for the negative control medium. Each virus dose was tested in replicates of six. After 60 min of incubation to allow virus attachment, 80 µl of growth medium was added to each well, and the plates were incubated for another 48 h. Cell viability was measured by adding 10 µl of WST-1 reagent (Roche)/well. An expansion of the number of viable cells increases the overall activity of mitochondrial dehydrogenases. This leads to an increase in the amount of formazan dye that is formed and directly correlates with the number of metabolically active cells in the culture. The formazan dye produced by metabolically active cells was quantified 2 h after the addition of WST-1 reagent by measurement in a scanning multwell spectrophotometer at 450 nm. Background absorbance was subtracted using the control medium and WST-1 reagent.

**Annexin V staining and DNA ladder.** Nuclei of rNDV-infected tumor cells were visualized after cell fixation in 4% paraformaldehyde, permeabilization with 0.1% Triton X-100, and incubation with a 90/10 mixture of ice-cold acetone and water, and DNA staining with 1.0 µg of 4',6'-diamidino-2-phenylindole (DAPI) (Sigma-Aldrich, St. Louis, Missouri). Externalization of phosphatidylserine from the inner to the outer leaftlet of the cell membrane in rNDV-infected tumor cells was detected using an annexin V-fluorescein isothiocyanate kit (BD Biosciences, San Jose, CA) per the manufacturer’s instructions at 6 and 14 h postinfection (p.i.). Propidium iodide was used as the vital dye to differentiate live, dead, and apoptotic cells by fluorescence microscopy (Axiovert 200; Carl Zeiss, Thornwood, NY). Intraunec- osomal DNA fragmentation in infected cells was detected by using an apoptotic DNA ladder kit (Roche). A kinetic assay of DNA fragmentation was performed in HuTu80 cells at 8, 10, 12, 24, and 48 h p.i.

**TNFα and TRAIL.** TNFα production by rNDV in SV-HUC1 and tumor cell lines was tested at 48 h p.i. using a human TNFα Quantikine enzyme-linked immunosorbent assay (ELISA) kit (HSTA00C; R&D Systems). The time course of TNFα induction by rNDV strains (MOI, 10) was monitored in HuTu80 cells at 6, 10, 12, 20, 24, 48, and 72 h p.i. by using the above-mentioned ELISA kit. A TRAIL ActivELISA kit (Imgenex, San Diego, CA) was used to detect the soluble form of TRAIL (sTRAIL) by a sandwich ELISA protocol according to the manufacturer’s recommendations. Culture supernatants (48 h p.i.) from rNDV-infected (MOI, 10) DF1 or tumor cells or standard dilutions of recombinant sTRAIL were tested in triplicate using a TRAIL ActivELISA kit. sTRAIL was captured on the microtiter plate by using an anti-TRAIL antibody. The amount of bound TRAIL was then detected by TRAIL-specific detecting antibody, followed by incubation with alkaline phosphatase-conjugated secondary antibody and color reaction with p-nitrophenyl phosphate. Optical density was measured at 405 nm and compared with those for the standard dilutions.

**Caspase assay.** ApoAlert Caspase fluorescent assay kits (BD Biosciences) for caspase-3, caspase-8, and caspase-9 were employed to detect the activation of different caspases in infected cell lysates. DF1 cells and various tumor cell lines were infected with rNDV strains (MOI, 10), harvested at 48 h p.i., and tested for activated caspases using fluorogenic substrates per the manufacturer’s instructions. In some cases, with HuTu80 cells, 1 × 10⁶ cells/time point were used. Cells were centrifuged at 200 × g, supernatants were removed, and the cell pellets were frozen at −70°C until cells from all of the time points were collected. Assays were performed in 96-well plates and analyzed using a fluorescence plate reader (PerkinElmer, Boston, MA). The caspase-3 and caspase-8 fluorogenic assay kits detect the enzyme shift of 7-amino-4-trifluoromethyl coumarin (AFC). The AFC-substrate conjugate usually emits blue light (λmax = 400 nm). However, cleavage of the substrate by the appropriate caspase liberates AFC, which fluoresces yellow-green (λmax = 505 nm). Similarly, the caspase-9 fluorogenic assay kit detects the enzyme shift of 7-amino-4-methyl coumarin (AMC). The LEHD-AMC conjugate emits light in the UV range (λmax = 380 nm); however, free AMC fluorescence blue-green at 440 nm upon liberation by caspase-9. The amount of fluorescence detected is directly proportional to the amount of caspase activity. Results of all experiments are reported as means ± standard errors of the mean (SEM).

**Caspase inhibition.** The selective inhibitors acetyl-Asp-Glu-Val-Asp-aldehyde (Ac-DEVd-CHO; inhibitor of caspase-3, -7, and -8), benzylxycarbonyl-Ille-Glu-Thr-Asp-fluoromethylketone (Z-LETd-FMK; inhibitor of caspase-6, -8, -9, and -10), and acetyl-Leu-Glu-His-Asp-aldehyde (Ac-LEHD-CHO; inhibitor of caspase-9) were used for caspase inhibition experiments. For apoptosis inhibition assays, HuTu80 cells were infected with 600 µM Ac-DEVd-CHO, 600 µM Ac-LEHD-CHO, or 120 µM Z-LETd-FMK for 3 h or with 100 µM benzylxycarbonyl-Val-Ala-Asp-fluoromethylketone (Z-VDd-FMK) for 1 h prior to infection with rBCEGFP (MOI of 1), rLaSota V.F., rBC, or rBCEdit viruses. Control cultures were treated with the appropriate amount of dimethyl sulfoxide (0.1%, final concentration), used as a solvent for peptide inhibitors. After virus adsorption for 1 h, the media were removed and replaced with OptiMEM. The infected cells were monitored for either EGFP expression or CPE and virus production at 6, 8, 10, 12, 14, 16, 20, 24, 38, 48, and 72 h p.i. Apoptosis induction in these cells was evaluated using an apoptotic DNA ladder kit (Roche) for intranecosomal DNA fragmentation at the time points indicated above.

**Mitochondrial membrane integrity.** To detect mitochondrial membrane integrity, tumor cells were cultured on 24-well cell culture plates. Fifteen minutes before fixation, the MitoTracker Red (750 nM) (CMX-Ros; Molecular Probes, Eugene, OR) dye was added to the culture medium. Accumulation of the dye was allowed to occur for 20 min at 37°C. Then, the cells were fixed with 4% paraformaldehyde for 15 min and permeabilized with 0.01% Triton X-100, used as a solvent for peptide inhibitors. After virus infection for 4 h, the media were removed and replaced with OptiMEM. The infected cells were monitored for either EGFP expression or CPE and virus production at 6, 8, 10, 12, 14, 16, 20, 24, 38, 48, and 72 h p.i. Apoptosis induction in these cells was evaluated using an apoptotic DNA ladder kit (Roche) for intranuclosomal DNA fragmentation at the time points indicated above.

**RESULTS**

**Rescue of Newcastle disease viruses from cloned cDNAs and expression vectors.** By reverse genetics technology, we have recovered several rNDVs expressing additional genes as transcriptional units (18, 26), including one carrying EGFP for this study. The recombinant viruses used in this study are listed in Table 1. The recovered recombinant viruses exhibited all of the biological phenotypes of their parental natural isolates. The rBCEGFP virus stably expressed EGFP even after 10 passages in 9- to 11-day-old embryonated chicken eggs and 15 passages in DF1 cells.

**Recombinant NDV strains are highly lytic for members of the National Cancer Institute panel and other cancer cell lines.** To assess the oncolytic properties of the rNDV strains, representative cell lines from the NCI-60 human tumor cell panel and other tumor cell lines (14 cell lines from a spectrum of malignancies) were infected with either rLaSota, rLaSota V.F., rBC, rBCEGFP, or rBCEdit viruses and assayed for metabolic cell death by the WST-1 assay 48 h later. The rNDVs were able to infect and induce cytolysis in a wide range of.

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**TABLE 1. Recombinant Newcastle disease virus strains used in this study**

<table>
<thead>
<tr>
<th>Virus*</th>
<th>Additional, replaced, or mutated gene</th>
</tr>
</thead>
<tbody>
<tr>
<td>Recombinant LaSota</td>
<td>None</td>
</tr>
<tr>
<td>Recombinant LaSota V.F.</td>
<td>Virulen Fusion (V.F.) gene cleavage site</td>
</tr>
<tr>
<td>Recombinant Beaudette C</td>
<td>None</td>
</tr>
<tr>
<td>Recombinant Beaudette C-Edit</td>
<td>V protein editing site mutant</td>
</tr>
<tr>
<td>Recombinant Beaudette C-EGFP</td>
<td>Enhanced green fluorescent protein (EGFP)</td>
</tr>
</tbody>
</table>

* All viruses were recovered entirely from cloned cDNAs of their respective antigenome by our established reverse genetics techniques.
TABLE 2. Recombinant NDV strains are highly lytic to tumor cells in vitro

<table>
<thead>
<tr>
<th>Cell line</th>
<th>rLaSota*</th>
<th>rLaSota V.F.*</th>
<th>rBC*</th>
<th>rBC-EGFP*</th>
<th>rBC-Edit*</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Sensitivity (%)</td>
<td>MOI</td>
<td>Sensitivity (%)</td>
<td>MOI</td>
<td>Sensitivity (%)</td>
</tr>
<tr>
<td>THP-1</td>
<td>0 &lt; 10</td>
<td>100 (1/1)</td>
<td>0.0001</td>
<td>100 (1/1)</td>
<td>0.003</td>
</tr>
<tr>
<td>CCRF-CEM</td>
<td>0 &gt; 10</td>
<td>100 (2/2)</td>
<td>0.142</td>
<td>50 (1/2)</td>
<td>0.003</td>
</tr>
<tr>
<td>Leukemia cell lines</td>
<td>0 &gt; 10</td>
<td>100 (1/1)</td>
<td>0.0005</td>
<td>100 (1/1)</td>
<td>0.001</td>
</tr>
<tr>
<td>Prostate cancer (PC3)</td>
<td>0 &gt; 10</td>
<td>100 (1/1)</td>
<td>0.028</td>
<td>100 (1/1)</td>
<td>0.02</td>
</tr>
<tr>
<td>HT1080</td>
<td>0 &gt; 10</td>
<td>100 (1/1)</td>
<td>3.565</td>
<td>100 (1/1)</td>
<td>0.07</td>
</tr>
<tr>
<td>HuTu80</td>
<td>0 &gt; 10</td>
<td>100 (1/1)</td>
<td>0.21725</td>
<td>100 (1/1)</td>
<td>0.07</td>
</tr>
<tr>
<td>Cervical cancer (HeLa)</td>
<td>100 (1/1)</td>
<td>0.01</td>
<td>100 (1/1)</td>
<td>0.019</td>
<td>100 (1/1)</td>
</tr>
<tr>
<td>Liver cancer (HepG2)</td>
<td>100 (1/1)</td>
<td>0.0164</td>
<td>100 (1/1)</td>
<td>0.019</td>
<td>100 (1/1)</td>
</tr>
<tr>
<td>Caco-2</td>
<td>0 &gt; 10</td>
<td>100 (1/1)</td>
<td>0.0573</td>
<td>100 (1/1)</td>
<td>0.008</td>
</tr>
<tr>
<td>CoLo 205</td>
<td>0 &gt; 10</td>
<td>100 (1/1)</td>
<td>0.0076</td>
<td>100 (1/1)</td>
<td>0.008</td>
</tr>
<tr>
<td>SW 620</td>
<td>0 &gt; 10</td>
<td>100 (1/1)</td>
<td>0.03</td>
<td>100 (1/1)</td>
<td>0.098</td>
</tr>
<tr>
<td>HT29</td>
<td>0 &gt; 10</td>
<td>100 (1/1)</td>
<td>4.648</td>
<td>0 &gt; 10</td>
<td>0 &gt; 10</td>
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<tr>
<td>T84</td>
<td>0 &gt; 10</td>
<td>0 &gt; 10</td>
<td>0 &gt; 10</td>
<td>0 &gt; 10</td>
<td>100 (1/1)</td>
</tr>
<tr>
<td>Colon cancer lines</td>
<td>0 &gt; 10</td>
<td>80 (4/5)</td>
<td>1.9357</td>
<td>20 (1/5)</td>
<td>0.988</td>
</tr>
<tr>
<td>SH-SYSY neuroblastoma</td>
<td>0 &gt; 10</td>
<td>100 (1/1)</td>
<td>0.002</td>
<td>100 (1/1)</td>
<td>0.025</td>
</tr>
<tr>
<td>Breast carcinoma (MC7-7)</td>
<td>0 &gt; 10</td>
<td>0 (0/1) &gt; 10</td>
<td>0 (0/1) &gt; 10</td>
<td>0 (0/1) &gt; 10</td>
<td>0 (0/1) &gt; 10</td>
</tr>
<tr>
<td>All cell lines tested</td>
<td>14 (2/14) &gt; 10</td>
<td>86 (12/14)</td>
<td>0.84</td>
<td>57 (8/14)</td>
<td>0.22</td>
</tr>
</tbody>
</table>

In parentheses are numbers of highly susceptible cell lines out of the number of cell lines tested. A cell line was considered highly susceptible if the EC50 was ≤ MOI of 1 following a 48-h infection. The MOI represents the average EC50 (MOI) of susceptible cell lines obtained from two independent experiments. All of the tested viruses are recombinant viruses derived by our established reverse genetics techniques.

A. Boldface type indicates combined results for indicated cell lines.

B. The sensitivity is the percentage of cancer cell lines by tumor type deemed highly sensitive to virus-mediated killing.
cer cells, such as HT29 and CaCo2 cells, apoptosis was induced by rNDV. Surface expression of TRAIL was detected in many of the cancer cell lines at 48 h p.i., suggesting that human TRAIL is essentially a membrane-bound protein (data not shown). Time course studies by Western blotting in HuTu80 cells indicated that TRAIL expression commenced only at 14 h p.i. (Fig. 3D), which suggests that TRAIL-mediated apoptosis is a late event in NDV-infected cells. Treatment of rBC-EGFP-infected HuTu80 cells with anti-TRAIL antibody (MAB687; R&D Systems) inhibited apoptosis but not the virus replication of rBC-EGFP virus, demonstrable by EGFP expression and viral plaque assay (data not shown).

Caspase-8 is activated but dispensable for apoptosis induction after NDV infection of tumor cells. Death receptor signaling by ligand binding at the cell surface leads to the formation of death-inducing signaling complex and activation of caspase-8. Therefore, we examined the activation of caspase-8 in NDV-infected cells. NDV infection activated caspase-8 in many of the tumor cell lines at 48 h p.i. (Fig. 4A). Caspase-8 was activated independently of IFN responsiveness, indicating that cells respond to NDV with apoptosis either in the presence or in the absence of IFN signaling (59). Caspase-8 was activated in a biphasic manner commencing at 12 h, with peak activation occurring at 32 h p.i. for the rBC virus in HuTu80 cells. Interestingly, rLaSota V.F. and rBC-Edit viruses induced caspase-8 late in the infection cycle (Fig. 4B), following a similar biphasic activation, correlating with the onset of CPE. This biphasic activation indicated that other pathways might amplify caspase-8-induced pathways or that caspase-8 is induced by additional pathways operating independently of death receptor signaling. Further support for this assumption stemmed from the fact that rapid CPE and apoptotic DNA laddering was evident after rNDV infection in TRAIL-sensitive, caspase-8-methylated SH-SY5Y neuroblastoma cells (Fig. 1 and 2D) and TRAIL-resistant CaCo2 and HT29 colon carcinoma cells (Fig. 2E). In CaCo2 and HT29 cells, apoptosis was induced by NDV without any caspase-8 activation. Loss of caspase-8 expression by gene methylation is suggested to be the reason for TRAIL resistance in neuroblastoma cells (53). Caspase-8 activation in neuroblastoma cells commenced only after TRAIL expression was evident in NDV-infected cells (Fig. 4C). Recent evidence suggests that SH-SY5Y neuroblastoma cells become TRAIL sensitive and FADD positive and induce caspase-8 upon stimulation with TRAIL and IFN-γ (22). As our results suggested that caspase-8 is probably not the initiator caspase, we looked for evidence as to whether NDV infection activated intrinsic, mitochondrion-associated...
apoptotic signaling pathways that may operate earlier than the death receptor pathway.

Mitochondrial membrane potential is not maintained following NDV infection. Cell death at the mitochondrial level is initiated by perturbation of the mitochondrial membrane leading to the release of various proapoptotic factors. To identify the drop in mitochondrial membrane potential ($\Delta \psi_{\text{mit}}$), mock- and NDV-infected cells were stained with MitoTracker Red CMX-Ros dye. The CMX-Ros dye is taken up only by actively respiring mitochondria with intact $\Delta \psi_{\text{mit}}$. The DNA-binding DAPI fluorophore was then used to delineate the nuclear morphology. NDV-infected cells which had a disruption of the $\Delta \psi_{\text{mit}}$ and were undergoing apoptosis were detected by the diffuse cytoplasmic pattern of CMX-Ros (Fig. 5A to I) with condensed chromatin. Cells with intact mitochondrial transmembrane potential displayed punctate staining with CMX-Ros (Fig. 5J to L). Following a drop of $\Delta \psi_{\text{mit}}$, cytochrome $c$ can be released from mitochondria through the opened mitochondrial pores. Therefore, we investigated the localization of cytochrome $c$ after NDV infection. Mitochondrial and cytosolic extracts from mock- and virus-infected cells were prepared by subcellular fractionation and analyzed by Western blotting. In NDV-infected cells, the level of cytochrome $c$ in cytosol increased twofold compared to the level observed in mock-infected cells (data not shown). These results indicate that the intrinsic mitochondrial pathway is initiated after infection with NDV.

Smac/DIABLO is not released from the mitochondria following NDV infection. Inhibitor of apoptosis proteins (IAPs) inhibit the enzymatic activity of caspases (55). IAP-mediated inhibition of apoptosis is countered by Smac/DIABLO, and Smac protein is secreted from mitochondria into the cytosol during apoptosis (11, 56). Synthetic Smac-mimicking molecules have been shown to potentiate TRAIL- and TNF-$\alpha$-mediated cell death (14, 32). We therefore examined the secretion of Smac into the cytosol after infecting HuTu80 cells with rNDV. Mitochondrion-free lysates were prepared from both mock- and NDV-infected cells at specified time points and analyzed by Western blotting for the presence of cytosolic Smac/DIABLO. We were not able to detect Smac expression in NDV-infected cells at various time points up to 48 h p.i. in HuTu80 cells.

Caspase-9 is activated early after NDV infection. In the cytosol, cytochrome $c$ forms an apoptosome with Apaf-1 and procaspase-9, leading to the activation of caspase-9. Caspase-9 was found to be activated by a fluorogenic substrate assay in
most of the tested cell lines (Fig. 6A) after rNDV infection. Significant levels of caspase-9 were induced as early as 6 h p.i. in HuTu80 cells, and caspase-9 also followed a biphasic pattern of activation (Fig. 6B). rLaSota V.F. and rBC-Edit viruses initiated low levels of caspase-9 between 8 and 10 h p.i. followed by another wave of significant induction after 48 h p.i. Similarly, in SH-SY5Y neuroblastoma cells, caspase-9 activation occurred by 12 h after infection with rNDV (Fig. 6C). These results suggest that apoptosis in NDV-infected tumor cells probably commences intrinsically through the double-stranded RNA intermediates and possibly through endoplasmic reticulum stress, leading to mitochondrial membrane destabilization and caspase-9 activation.

**Effector caspase activation.** Having shown that NDV infection resulted in the activation of both death receptor and mitochondrion-associated pathways, we tested the activation of caspase-3. As shown in Fig. 7A, caspase-3 activation was detected by a fluorogenic substrate assay in many of the cell lines associated with apoptosis. Caspase-3 followed a biphasic activation similar to that of caspase-9 (Fig. 7B). In caspase-3 null MCF-7 breast cancer cells, rNDV strains replicated, although at reduced efficiency, but failed to exert cytolysis (EC50 > 10), consistent with the idea that caspase-mediated cell death is the predominant mechanism of oncolysis by NDV. In caspase-8-methylated SH-SY5Y neuroblastoma cells, caspase-3 was detected as early as 8 h p.i., which indicates that other initiator pathways are involved in effector caspase activation (Fig. 7C). IFN-sensitive rBC-Edit virus induced caspase-3 in HuTu80 cells at 32 h p.i., while it induced caspase-9 as early as 10 h p.i. but caspase-8 only at 48 h p.i., reinforcing the view that intrinsic apoptotic pathways operate early in NDV-infected cells (Fig. 7B). Classically, caspase-8 has been viewed as an initiator caspase involved in death receptor signaling (62). However, our results demonstrated that caspase-8 was activated only after caspase-9 and caspase-3 were activated. This time course suggested that caspase-8 was not the primary initiator caspase involved in NDV-mediated apoptosis. These results also suggest that caspase-8 activation probably occurs as an indirect effect of activation of caspase-9 and caspase-3 (61) or through the death receptor pathway late in the infection. It is probable that other initiator caspases might be involved, depending on the cell type and virus strain in effector caspase activation.
**NDV-induced apoptosis is caspase dependent.** To identify whether rNDV-induced apoptosis is entirely dependent on caspase activation, HuTu80 epithelial cells were pretreated with the broad-specificity caspase inhibitor Z-VAD-FMK or with relatively specific inhibitors of caspase-9 (Z-LEHD-FMK), caspase-3 (Z-DEVD-FMK), and caspase-8 (Z-IETD-FMK) and infected with rBC-EGFP. The expression of EGFP was monitored by epifluorescence microscopy, and virus content in the supernatant was determined by a plaque assay on DF1 cells at specified intervals. The broad-specificity caspase inhibitor Z-VAD-FMK was able to inhibit rNDV-triggered cytolysis completely until 48 h p.i., indicating that NDV-mediated cytotoxicity is caspase dependent. But after 48 h p.i., CPE appeared in the form of syncytia and cell rounding, indicating that rNDV can also induce caspase-independent lysis, perhaps as a direct result of viral replication. An intranucleosomal DNA fragmentation assay of infected cells confirmed apoptosis. Furthermore, despite the block in apoptosis, titers of infectious virus in Z-VAD-FMK-treated cells were similar to those in untreated cells (Fig. 7D), which implies that apoptosis is not required by the virus as a mechanism to facilitate its replication. Caspase-8 and caspase-9 inhibitors also suppressed morphological and biochemical alteration of NDV-infected cells until 48 h p.i. But none of them were able to individually protect HuTu80 cells from rNDV-induced apoptosis, which suggests that rNDV triggers multiple caspase-dependent pathways in these cells. From the above results, we infer that NDV triggers apoptosis by a rapid dissociation of the mitochondrial membrane potential, the release of cytochrome c, and the activation of caspase-9. Further, we detected caspase-8 late in the infection cycle, suggesting that multiple pathways that lead to the death of NDV-infected cells could be acting after an initial proapoptotic signal subsequent to the entry of the virus.

**DISCUSSION**

The genetic malleability and inherent tumor selectivity of NDV afford the opportunity to enhance its therapeutic potential. With the advent of reverse genetic systems for NDV, it is now possible to refine and optimize the oncolytic potency,
specificity, and therapeutic index, besides specifically retargeting it to the desired tumor cells. However, the exact modes of tumor cell killing by NDV are still incompletely understood. Understanding these mechanisms is crucial to improving oncolytic efficacy. Keeping this in mind, we explored the molecular mechanisms of oncolysis by rNDV. In order to achieve this, we examined the oncolytic potential of rNDVs with deficiencies in replicative abilities, cell and tissue tropism, and interferon antagonism. In addition, we engineered an rNDV with a reporter (EGFP) gene to make it possible to monitor virus infection. We have demonstrated that human tumor cells of neuroectodermal, mesenchymal, and epithelial origins could be infected and lysed by rNDV.

A nonpathogenic NDV (rLaSota V.F.) with a modified fusion protein cleavage site showed cytotoxicity against a wide range of tumor cells (86%), while a moderately pathogenic NDV (rBC) was cytolysic in only 57% of the tested tumor cells. However, the rBC virus efficiently lysed tumor cells at a relatively lower MOI. Engineering an additional transgene in rBC virus did not lower its oncolytic efficacy, as the cytotoxic range and effective concentration to lyse tumor cells of rBC and rBC-EGFP viruses were similar. But, the rLaSota V.F. and rBC viruses differed in their cytotoxic abilities against tumor cells, despite having similar F protein cleavage site sequences. We have previously shown that they also differ in virulence in chickens, the natural host for these viruses (41). Therefore, the observed differences in tumor cell cytotoxicity might be a reflection of the differences in the other major surface glycopro-

FIG. 5. Disruption of mitochondrial membrane potential in rBC-EGFP-infected tumor cells was examined by staining with DAPI and MitoTracker Red CMX-Ros. NDV-infected cells which had a disruption of the $\Delta\phi_m$ and were undergoing apoptosis were shown by the diffuse cytoplasmic pattern of CMX-Ros with condensed chromatin. Tumor cells were infected with rBC-EGFP virus, treated 24 h postinfection with MitoTracker Red CMX-Ros for 2 h, and fixed later. Syncytium formation, EGFP expression, and mitochondrial membrane disruption in rBC-EGFP-infected cells are shown. (A) CaCo2 colon carcinoma cells, bright field; magnification, $\times40$. (B) Epifluorescence microscopy, $\times40$. (C) Diffuse staining of cytoplasm with MitoTracker Red merged with fluorescent image, $\times40$. (D) HEpG2 hepatocarcinoma cells, bright field, $\times40$. (E) Epifluorescence, $\times40$. (F) Diffuse cytoplasmic staining of MitoTracker Red with EGFP expression, $\times40$. (G) PC3 prostate cancer cells, bright field, $\times40$. (H) Epifluorescence, $\times40$. (I) Diffuse cytoplasmic staining of MitoTracker Red with EGFP expression, $\times40$. (J) HuTu80 cells, uninfected control cells, bright field, $\times40$. (K) Epifluorescence, $\times40$. (L) Punctuate cytoplasmic staining with MitoTracker Red, $\times40$. 

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tein, the HN protein. The HN protein has been shown to mediate apoptosis in NDV-infected cells (63). There are 17 amino acid differences in the HN protein between rLaSota and rBC viruses that might contribute to these differences. Since all strains of NDV enter all types of cells using sialic acid receptors, the observed differences in cytotoxicity against tumor cells by these viruses are likely due to other HN protein functional differences that do not alter receptor specificity. It is highly likely that these differences could be due to a difference in receptor avidity resulting from the amino acid differences in the HN protein. It is also possible that other viral proteins, such as the V and L proteins, may be responsible for these differences. The rBC-Edit virus, on the other hand, showed a wider spectrum of cytotoxicity with a very low MOI. This could be due to the absence of expression of the V protein, which is reported to be an antiapoptotic protein (42).

NDV has been reported to induce apoptosis in infected cells (12, 27, 63), which is considered to be mediated by TRAIL (58). Mitogen-activated protein kinase pathways or reactive oxygen species are not associated with apoptosis induced by NDV (12). However, the exact cellular pathways of NDV-induced apoptosis and the mechanistic basis of oncolysis are largely unknown. By using an array of tumor cells defective in components of apoptotic machinery and IFN signaling, we have shown that NDV mediates tumor cell killing independently of IFN signaling by using multiple caspase-mediated apoptotic pathways. The inability to execute cell death in caspase-3 null MCF-7 cells reinforces our claim that NDV is oncolytic only by caspase-dependent apoptosis. The process of mitochondrial outer membrane permeabilization leading to the leakage of cytochrome c and probably other proteins, including Smac, from the intermembrane space (IMS) and the factors that regulate this process following rNDV-induced apoptosis remain to be discerned. The redistribution of these particular IMS proteins may be triggered by events upstream of the mitochondria, such as Bax recruitment to the outer mitochondrial membrane. Additionally, feed-forward loops external to mitochondria, such as those involving Ca2+ release.
from endoplasmic reticulum, may act to coordinate the entire population of mitochondria in one cell to release cytochrome c and other IMS proteins (5, 33).

The most proximal event in NDV-mediated apoptosis is the activation of caspase-9, followed by the activation of caspase-3, and ultimately the activation of caspase-8. Given that caspase-8 was originally identified as an initiator caspase triggered by death receptors, its involvement as a late event in NDV-induced apoptosis was unanticipated. The sequence of caspase activation we observed suggests activation of caspase-8 via caspase-3, which in turn may be triggered by the mitochondrial pathway via caspase-9. Once activated, caspase-8 may participate in an amplification mechanism, further activating effector caspases to accelerate NDV-mediated cell death. Indeed, recent studies have shown the ability of chemotherapeutic drugs to elicit the processing of caspase-8 in the absence of CD95 ligand-receptor interaction, and caspase-8 activation can occur downstream of caspase-9 and caspase-3 in response to anticancer drugs (60–61). In this regard, pathways of apoptosis induced by NDV overlap those induced by DNA-damaging chemotherapeutic drugs. NDV causes the loss of mitochondrial membrane potential both in wild-type cells and in cells devoid of functional caspase-8. Inhibition of caspase-8 activation, as in neuroblastoma cells, or TRAIL resistance, as in HT29 and CaCo2 cells (29), did not alter the apoptotic events in these cells. Despite TRAIL resistance, we were able to demonstrate that the rBC-Edit virus and the rLaSota V.F. virus were apoptotic in HT29 and CaCo2 cells, but the rBC virus was nontoxic in these cells, probably due to the potent antiapoptotic activity of the V protein. These results confirm our findings that NDV-induced TRAIL potentiates only the intrinsic apoptotic pathway.
As reported for vesicular stomatitis virus (1, 15), we also found that caspase-8 and caspase-9 inhibitors did not completely abrogate the signs of apoptosis in NDV-infected cells and that the presence of the general caspase inhibitor Z-VAD-FMK during the infection cycle prevented apoptosis and delayed cell death but could not inhibit CPE induced by the virus after 48 h p.i. In addition, we found that the activation of caspases in NDV-infected cells is not a requirement for viral replication, since caspase inhibitors had no effect on virus replication. These data show that NDV kills cancer cells primarily by activation of the mitochondrial death pathway, and furthermore, they indicate that the caspase-8/Bid-dependent signal amplification loop is not important for NDV-induced death. Viruses that induce death receptor-dependent apoptosis include human immunodeficiency virus (38), measles virus (57), influenza A virus (40), Sindbis virus (16), reovirus (9), lyssavirus (23), and hepatitis C virus (2). A number of viruses have been found to cause relocalization of proapoptotic mitochondrial proteins into the cytosol. Among these are human immunodeficiency virus (13), influenza A virus (7), herpes simplex virus type 1 (64), herpes simplex virus type 8 (49), hepatitis B virus (54), reovirus (24), and West Nile virus (43).

Our data identify the mechanistic events leading to apoptosis when tumor cells are infected with NDV. We have demonstrated that apoptosis is initiated by rNDV through the mitochondrial intrinsic pathway, leading to the activation of caspase-9 and effector caspase-3. A second round of caspase activation occurs when TRAIL-induced death receptor-mediated or caspase-3-mediated caspase-8 activation commences. Caspase-8 amplifies the cycle, allowing full expression of apoptosis leading to oncolysis. While caspase-8 may play a role in amplifying effector caspase activation, this appears to be unnecessary for NDV-induced apoptosis. The delay in TRAIL expression also supports this view. Reoviruses induce apoptosis primarily through the death receptor pathway, and cytochrome c release and subsequent caspase-9 activation were not critical mitochondrial events in apoptosis induction (24). NDV, on the other hand, depends primarily on the mitochondrial apoptotic pathway with no dependence on the death receptor pathway, which suggests that the molecular mechanisms of apoptosis may depend on the virus strain and cell type.

We have also shown that reporter genes could be engineered in rNDV to track the sojourn of oncolytic virus in the host without diminishing its oncolytic efficacy. The rLaSota V.F. and rBC-Edit viruses reported here have a number of properties which make them suitable as oncolytic viruses for tumor therapy. (i) These viruses are nonpathogenic to humans. We therefore hypothesize that they will be associated with fewer side effects. In fact, our recent studies in nonhuman primates showed that rNDV undergoes only a limited replication in the lungs with no clinical effects (6). (ii) Multiple serologically defined types exist for avian paramyxoviruses (APMV); different serotypes (APMV-2 to APMV-9) can therefore be engineered to deliver therapeutic transgenes. Alternatively, rNDV (APMV-1) with different antigenic specificity could be constructed by exchanging the surface glycoproteins of the virus from serotypes APMV-2 to -9. Therefore, immunity of the host after the first round of rNDV therapy can be circumvented by choosing rNDV with appropriate surface glycoproteins or recombinant APMV of differing serotypes, allowing repeated administration. (iii) It is possible to construct rNDVs with multiple transgenes to enhance oncolytic efficacy, as NDV has been shown to accommodate up to 4 kb of foreign genes. (iv) These viruses could be engineered for targeting to specific sites.

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REFERENCES
