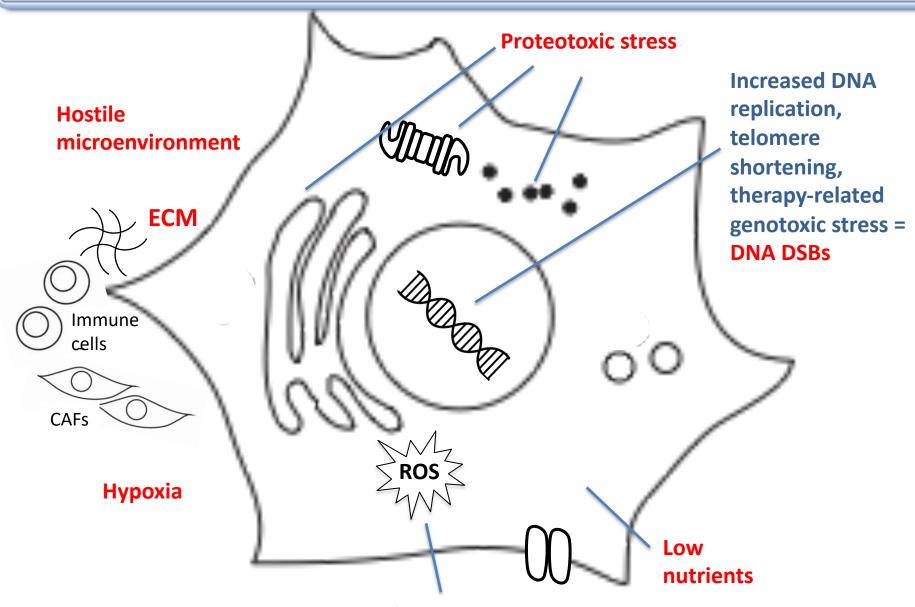
HALLMARK #4: EVASIONE DAI MECCANISMI ONCOSOPPRESSIVI INTRINSECI

RESISTENZA ALLA MORTE CELLULARE

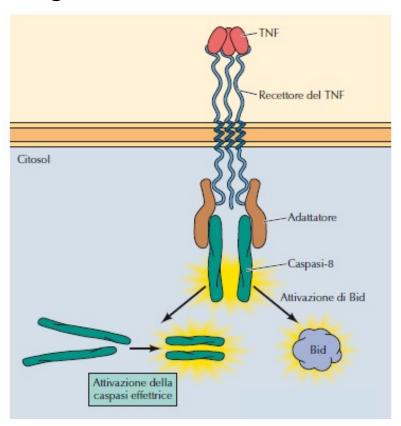
Le cellule tumorali sono sottoposte a molteplici stress



Le strategie terapeutiche mirate all'induzione di morte cellulare devono essere mirate ai meccanismi di morte che sono attivi nelle cellule tumorali

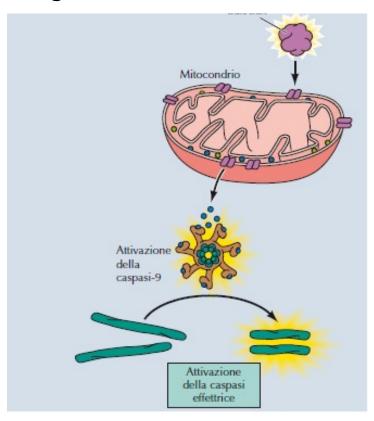
Apoptosi: via estrinseca e via intrinseca

Segnale di morte extracellulare



Es. TRAIL

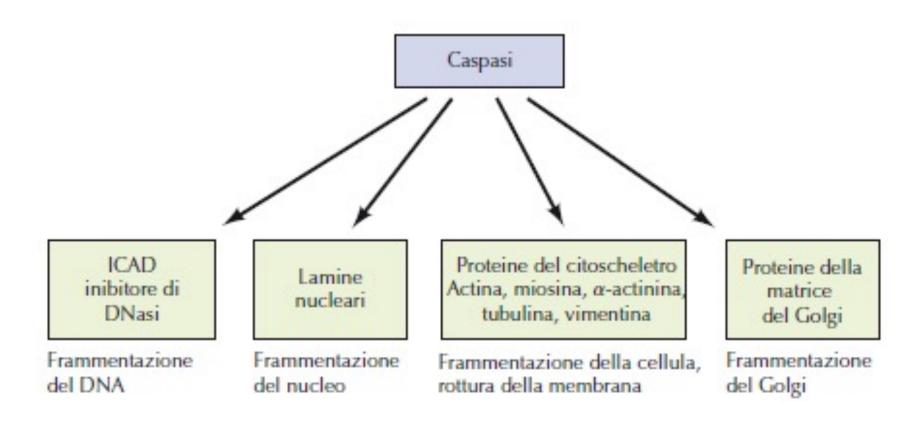
Segnale di morte intrinseco



Es.
Danno al DNA non riparabile
ER stress cronico,
Ipossia
Stress metabolico

Effettori dell'apoptosi: le CASPASI

Cistein-proteasi che tagliano a monte di un residuo di Asp



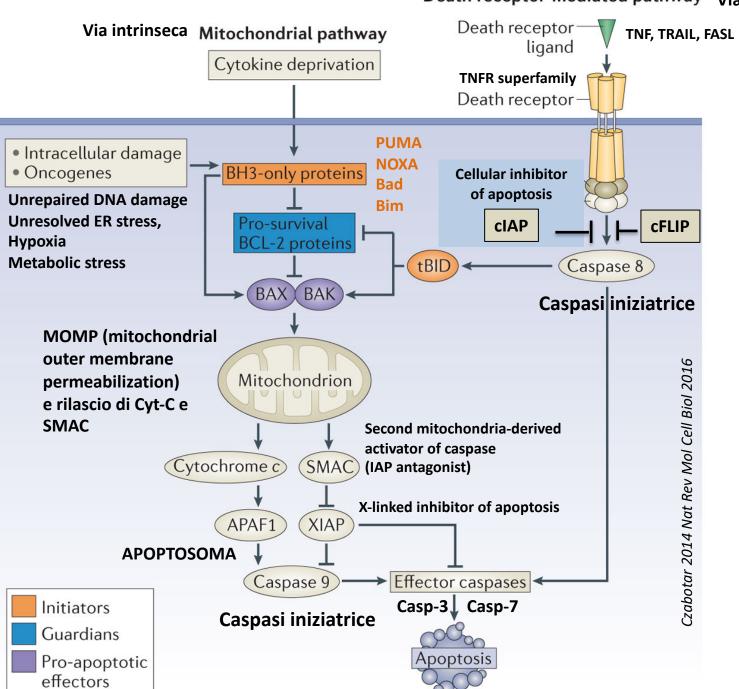
Regolatori dell'apoptosi

➤ Membri della famiglia di Bcl-2 (B-cell lymphoma 2)

Inhibitori dell' apoptosi (IAPs)

Antagonisti di IAP (SMAC/DIABLO)

Death receptor-mediated pathway Via estrinseca

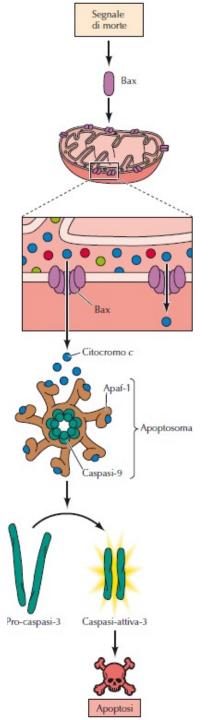


Regolatori dell'apoptosi

➤ Membri della famiglia di Bcl-2 (B-cell lymphoma 2)

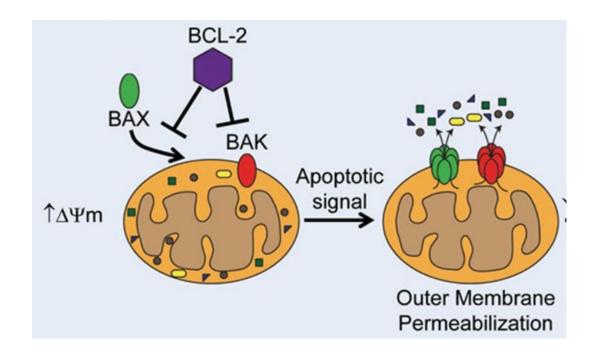
Inhibitori dell' apoptosi (IAPs)

Antagonisti di IAP (SMAC/DIABLO)

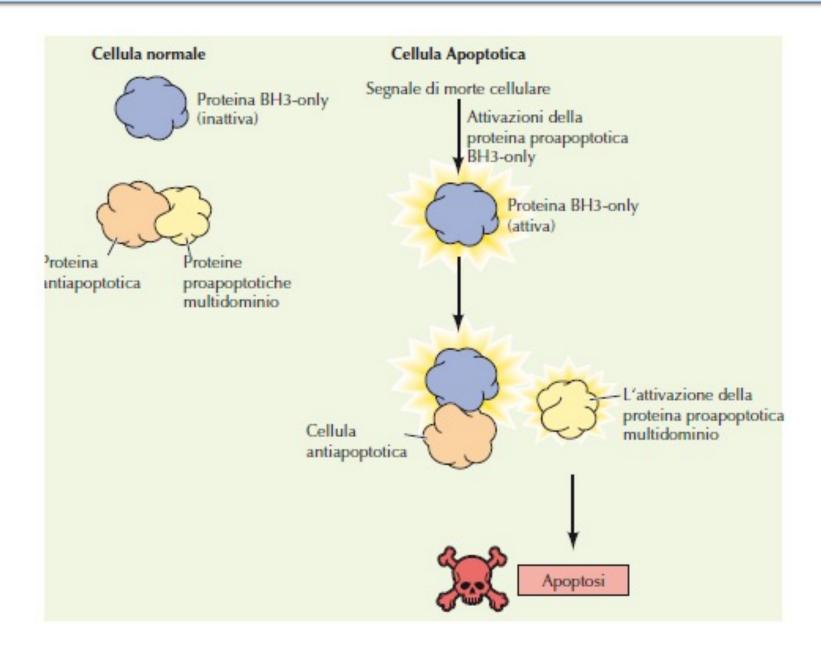


Le proteine proapoptotiche multidominio:

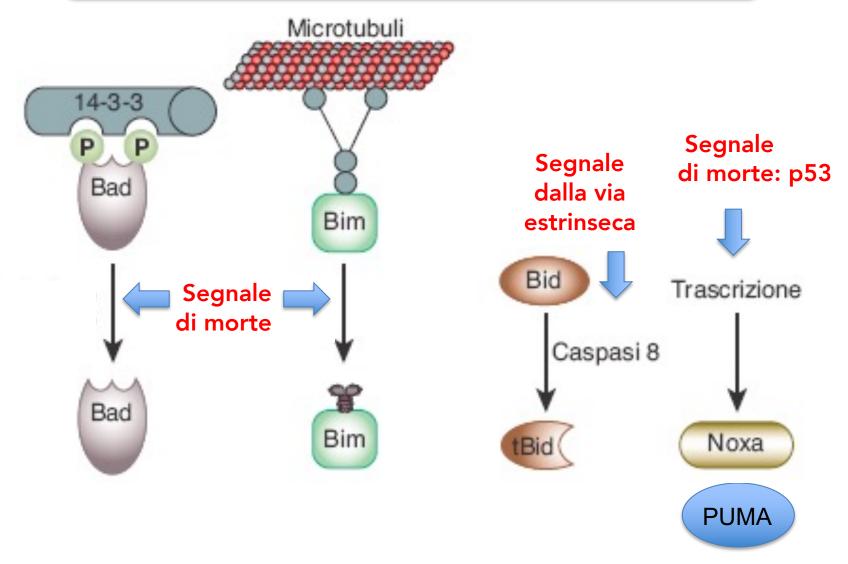
EFFETTORI della permeabilizzazione mitocondriale (MOMP)



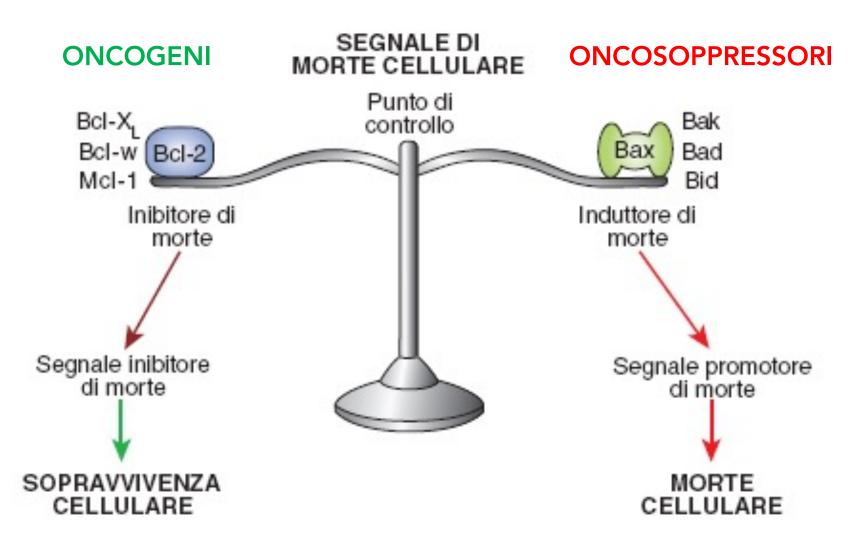
Interazioni regolatorie tra i membri della famiglia di Bcl-2



L'attivazione della via intrinseca è mediata dalle piccole proteine BH3-only



I membri della famiglia di Bcl2 regolano l'equilibrio tra sopravvivenza e morte cellulare



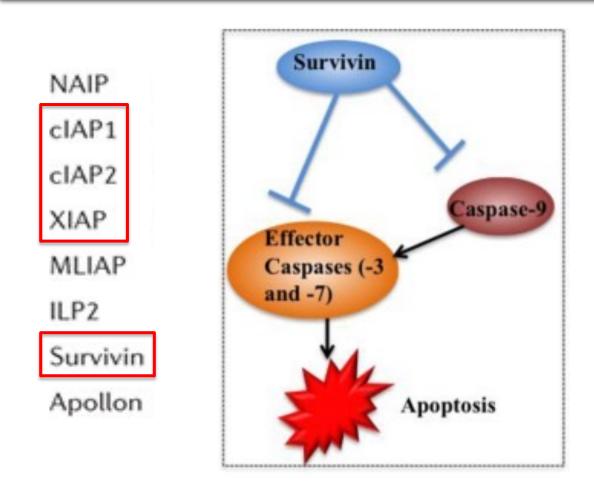
Regolatori dell'apoptosi

➤ Membri della famiglia di Bcl-2 (B-cell lymphoma 2)

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Antagonisti di IAP (SMAC/DIABLO)

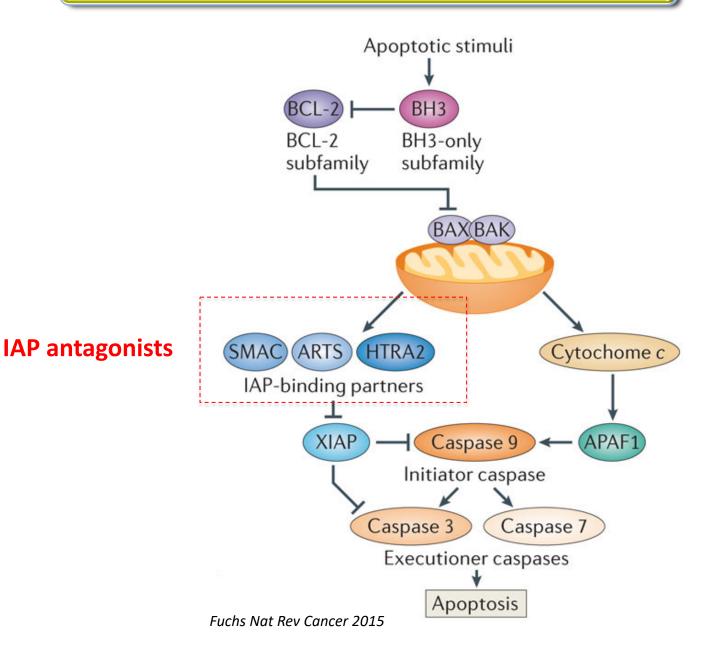
Inhibitor of apoptosis proteins IAPs

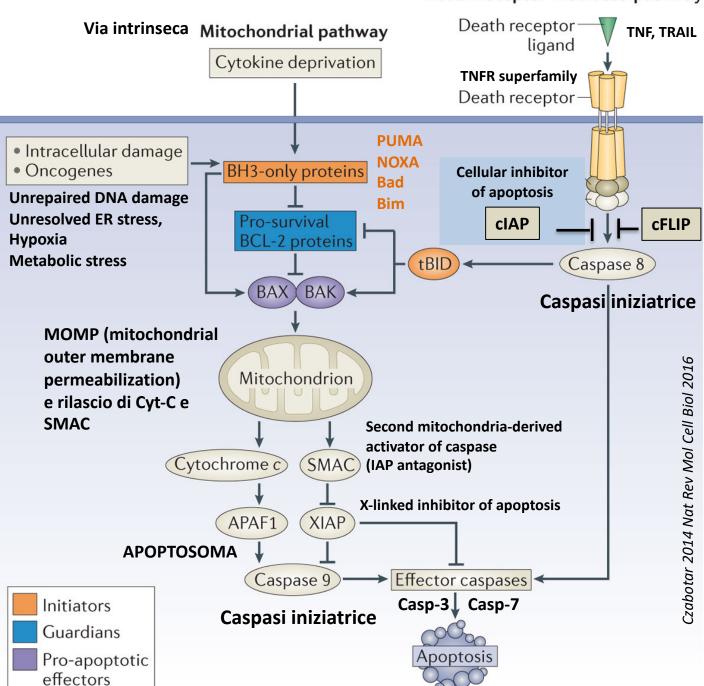


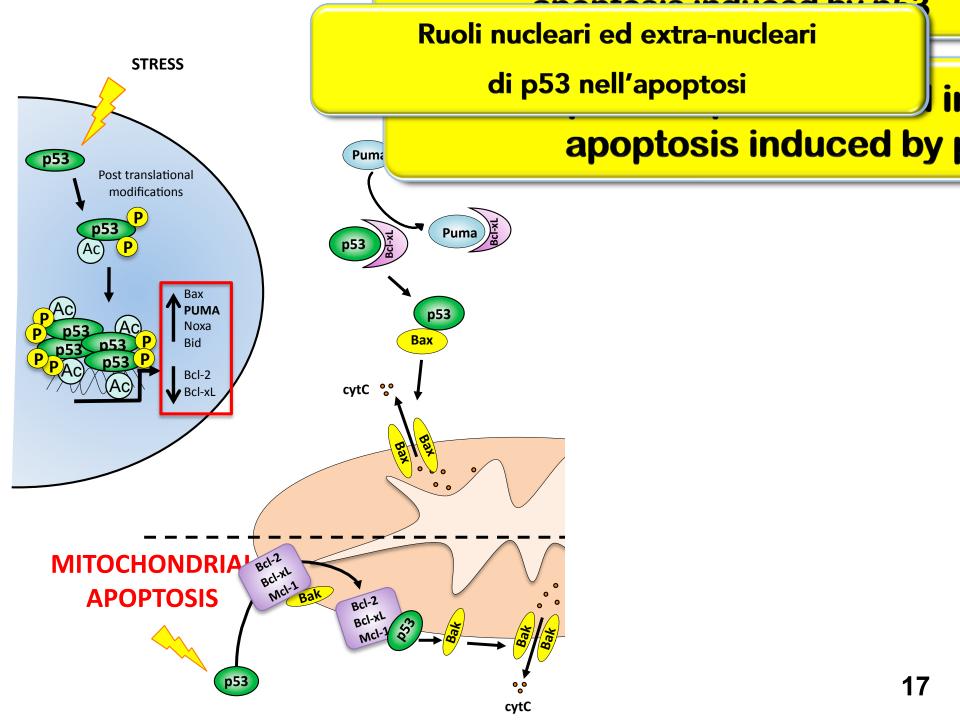
IAPs legano direttamente le CASPASI

RING (really interesting new gene) domain conferisce attività ubiquitin protein ligase (E3) causando degradazione di proteine proapoptotiche 14

IAP antagonists





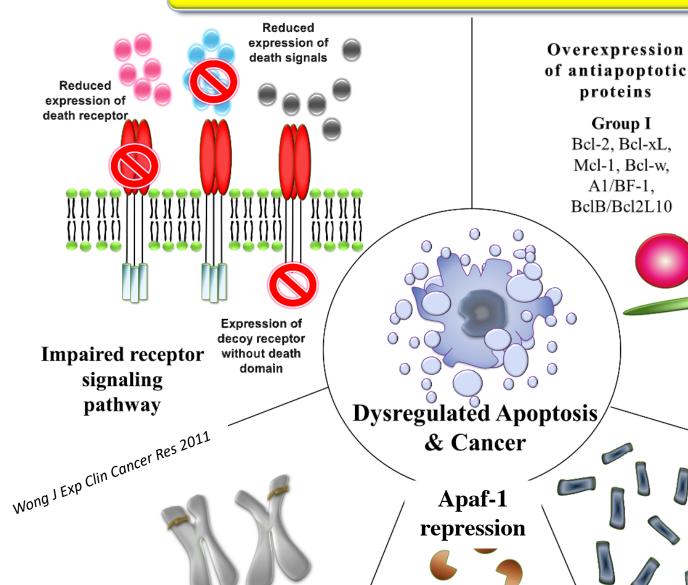


Meccanismi di evasione dall'apoptosi

Reduced

expression of

caspases



Defects/ mutations

in p53

Underexpression expression of proapoptotic proteins

Group II

Bid, Bim, Bax, Bak,

Puma, Bok/Mtd

Noxa,Bad,

Bmf, Hrk, Bik

Disrupted balance of Bcl-2 family of proteins

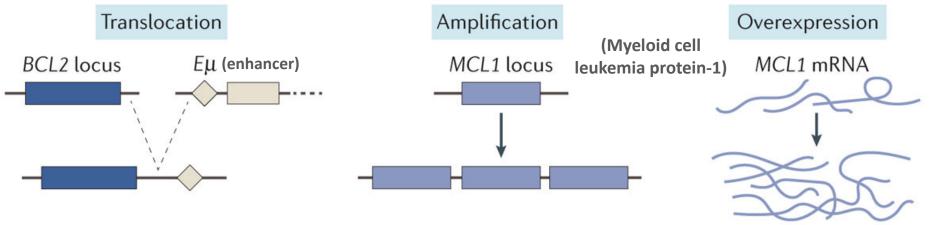
XIAP

Increased expression of IAPs

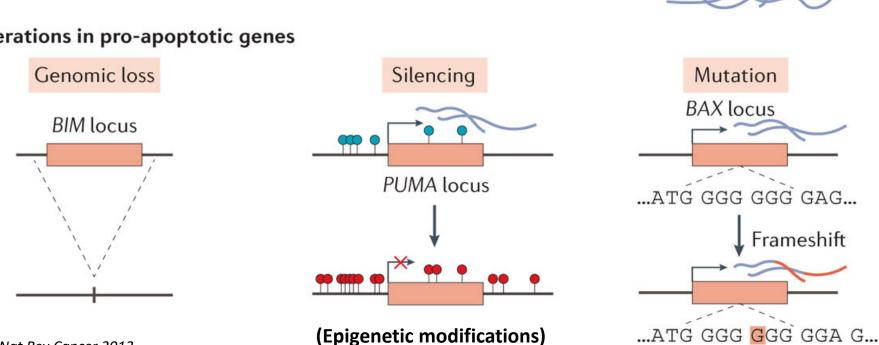
18

Deregolazione dei membri della famiglia di Bcl2

a Alterations in anti-apoptotic genes



Alterations in pro-apoptotic genes

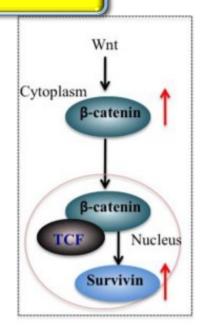


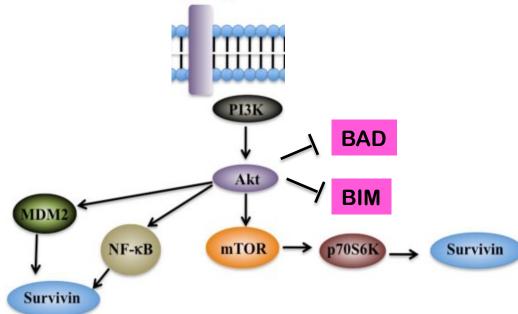
Deregolazione della survivina

Survivin aspase-9 Effector Caspases (-3 and -7) Apoptosis

È indotta in molti tumori da stress e pathways oncogeniche







Terapie mirate per l'induzione di apoptosi

TRAIL receptor agonists

Bcl-2 targeted chemotherapeutics

BH3 mimetics

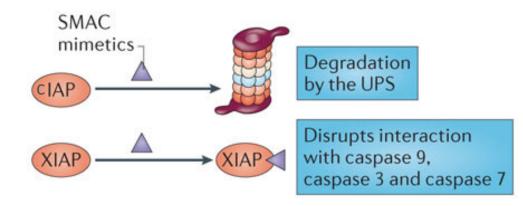
BCL-2 or BCL-X_L

BH3 mimetics

MOMP

IAP antagonist molecules

SMAC mimetics



Holohan Nat Rev Cancer 2013

Farmaci contro Bcl2

Non-Hodgkin lymphoma

Chronic lymphocytic leukaemia

B cell non-Hodgkin lymphoma

Diffuse large B cell lymphoma

Diffuse large B cell lymphoma

Haematological malignancies including

Follicular lymphoma

Follicular lymphoma

Non-Hodgkin lymphoma

myelodysplasia

Non-Hodgkin lymphoma

Clinical trial

Phase III

Phase I/II

Phase I

Phase I

Phase I

Phase I

Phase I

Phase I

Phase III

Phase I/II

Phase I/II

Phase II

Phase II

Phase I

Phase II

Phase II

Phase II

		•	diffici com				
Table 1 BH3-mimetics undergoing clinical trials for cancer indications							
BH3-mimetic	Alternative name	Targets	Therapy	Indication			
ABT-199	Venetoclax	BCL-2	Single agent	Chronic lymphocytic leukaemia			
				Acute myeloid leukaemia			
				Diffuse large B cell lymphoma			
				Follicular lymphoma			
				Lymphoma			
				Mantle cell lymphoma			
				Multiple myeloma			

Combination*

Single agent

Single agent

Data compiled from the Global Data database: http://healthcare.globaldata.com/ (accessed July 2015). *Combination with standard of care therapies.

None

None

Delbridge Nat Rev Cancer 2013

S-055746

PNT-2258

BCL-2

BCL-2

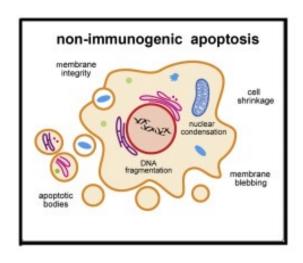
IAP targeted chemotherapeutics

Table 1. Clinical trials with Smac mimetics

Compound	Combination	Cancer type	Status	Phase I/II
LCL-161	None	Solid tumors	Completed	Phase I
LCL-161	None	Leukemia	Recruiting	Phase II
LCL-161	Paclitaxel	Solid tumors	Completed	Phase I
LCL-161	Paclitaxel	Solid tumors	Recruiting	Phase I
LCL-161	Paclitaxel	Breast cancer	Completed	Phase II
LCL-161	Gemcitabine	Pancreatic cancer	Recruiting	Phase I
LCL-161	Cyclophosphamide	Multiple myeloma	Recruiting	Phase II
GDC-0152	None	Solid cancers	Completed	Phase I
CUCD-427	None	Lymphoma	Recruiting	Phase I
Birinapant	None	Solid tumors	Completed	Phase I/II
Birinapant	None	Solid tumors, lymphoma	Completed	Phase I
Birinapant	None	AML	Completed	Phase I/II
Birinapant	None	Ovarian, peritoneal and fallopian tube cancer	Completed	Phase II
Birinapant	Gemcitabine	Solid tumors	Terminated	Phase I
Birinapant	5-Aza	MDS	Active, not recruiting	Phase I/II
Birinapant	5-Aza	MDS, CMML	Recruiting	Phase II
Birinapant	Conatumumab	Ovarian, peritoneal and fallopian tube cancer	Recruiting	Phase I
AT-406	Daunorubicin, cytarabine	AML	Terminated	Phase I
Debio1143	None	Solid tumors, lymphoma	Completed	Phase I
Debio1143	Paclitaxel, carboplatin	Solid tumors	Recruiting	Phase I
Debio1143	Cisplatin, radiotherapy	Head and neck carcinoma	Recruiting	Phase I/II
HGS1029	None	Solid tumors	Completed	Phase I
HGS1029	None	Lymphoid malignancies	Terminated	Phase I

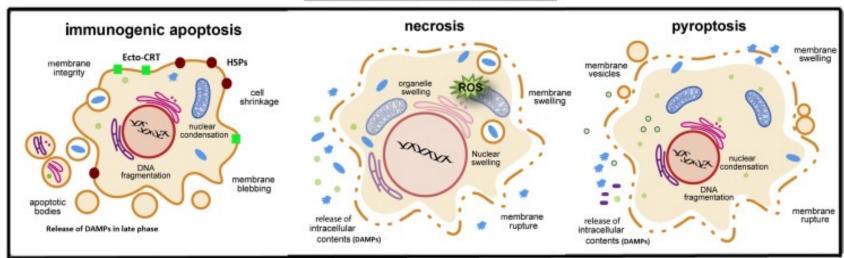
Abbreviations: 5-Aza, 5-Azacitidine; AML; acute myelogenous leukemia; CMML, chronic myelomonocytic leukemia; MDS, myelodysplastic syndrome.

Induzione di morte cellulare tolerogenica e immunogenica:



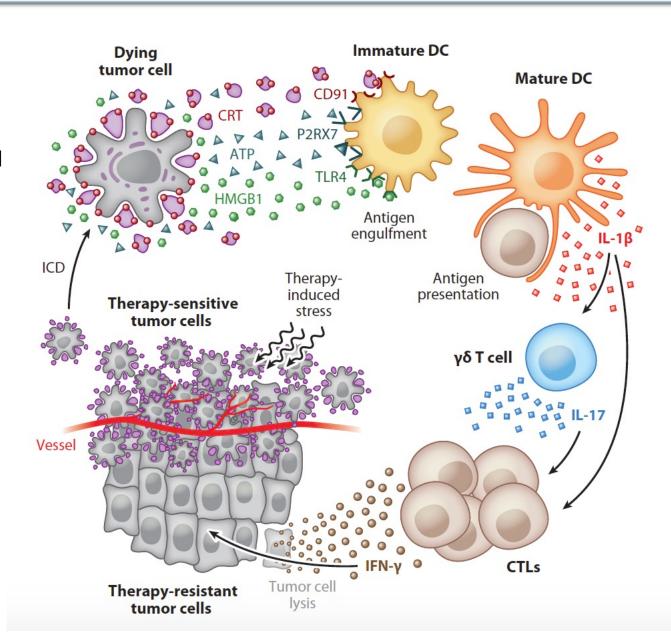
Non-immunogenic cell death mode

Immunogenic cell death modes



Induzione di morte cellulare tolerogenica e immunogenica:

Calreticulin, HSPs, Release of DAMP (damage-associated molecular patterns: ATP, HMGB1...)



Induzione di morte cellulare immunogenica e non immunogenica: implicazioni terapeutiche

