

Molekulární chaperony a jejich úloha v patogenezi lidských chorob

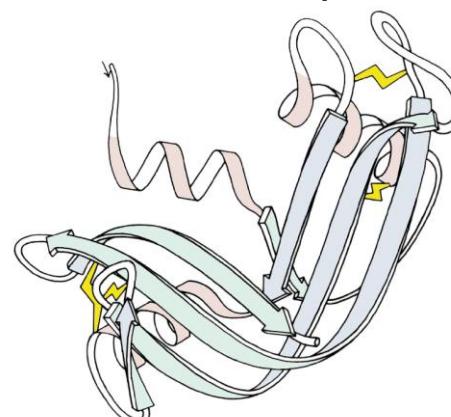
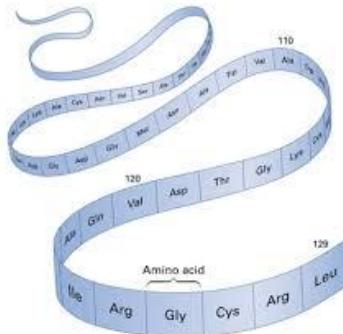
Petr Muller

RECAMO

Regional Centre
for Applied Molecular
Oncology



At the environmental conditions (temperature, solvent concentration and composition, etc.) at which folding occurs, the native structure is a unique, stable and kinetically accessible minimum of the free energy

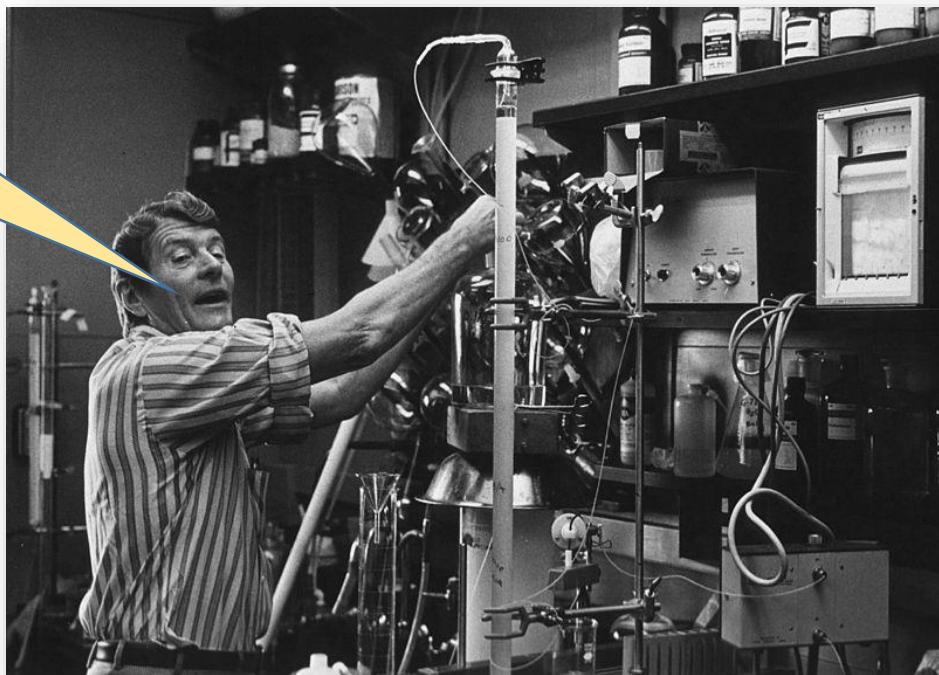


Ribonuclease A

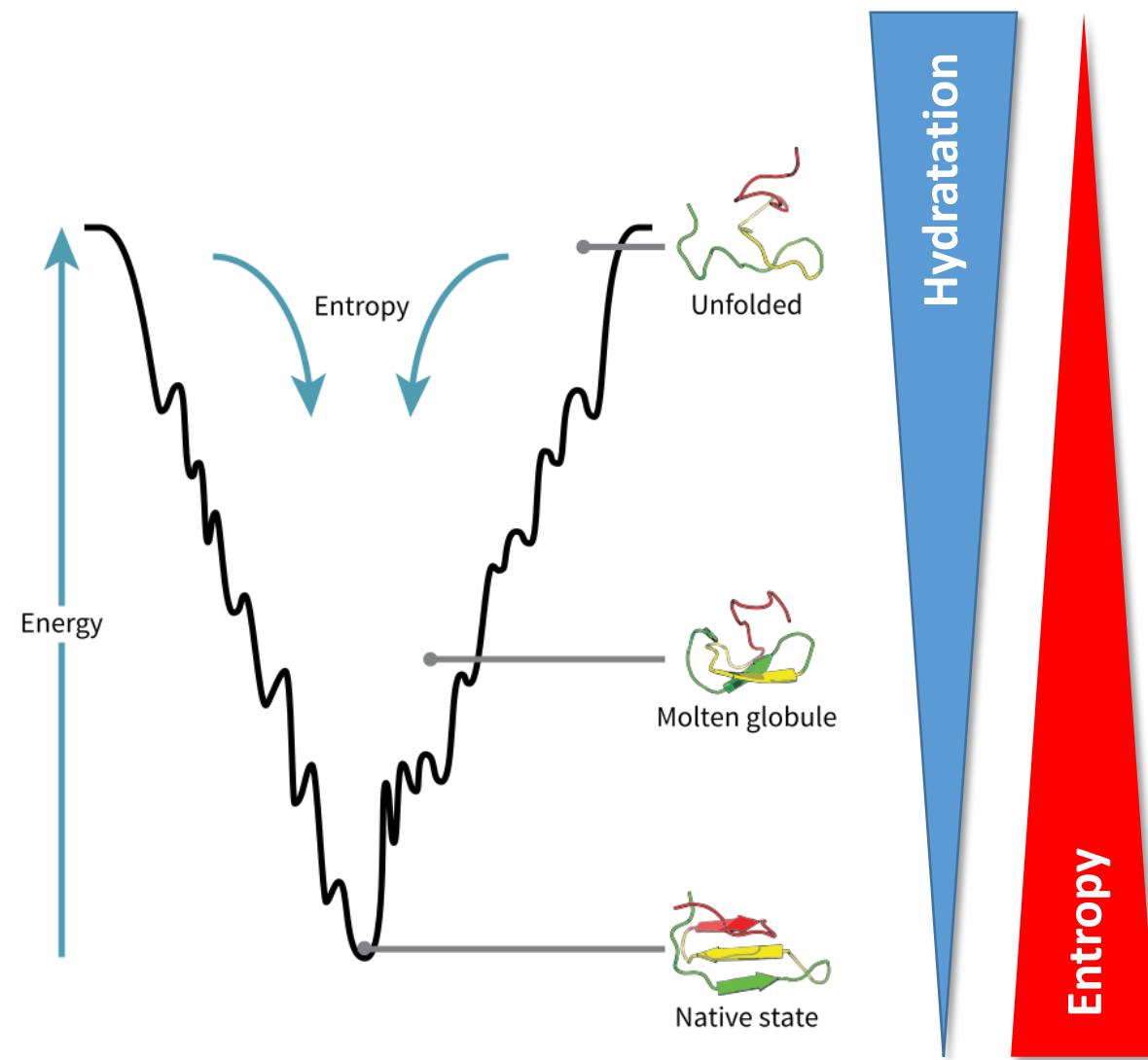
The native structure is determined only by the protein's amino acid sequence

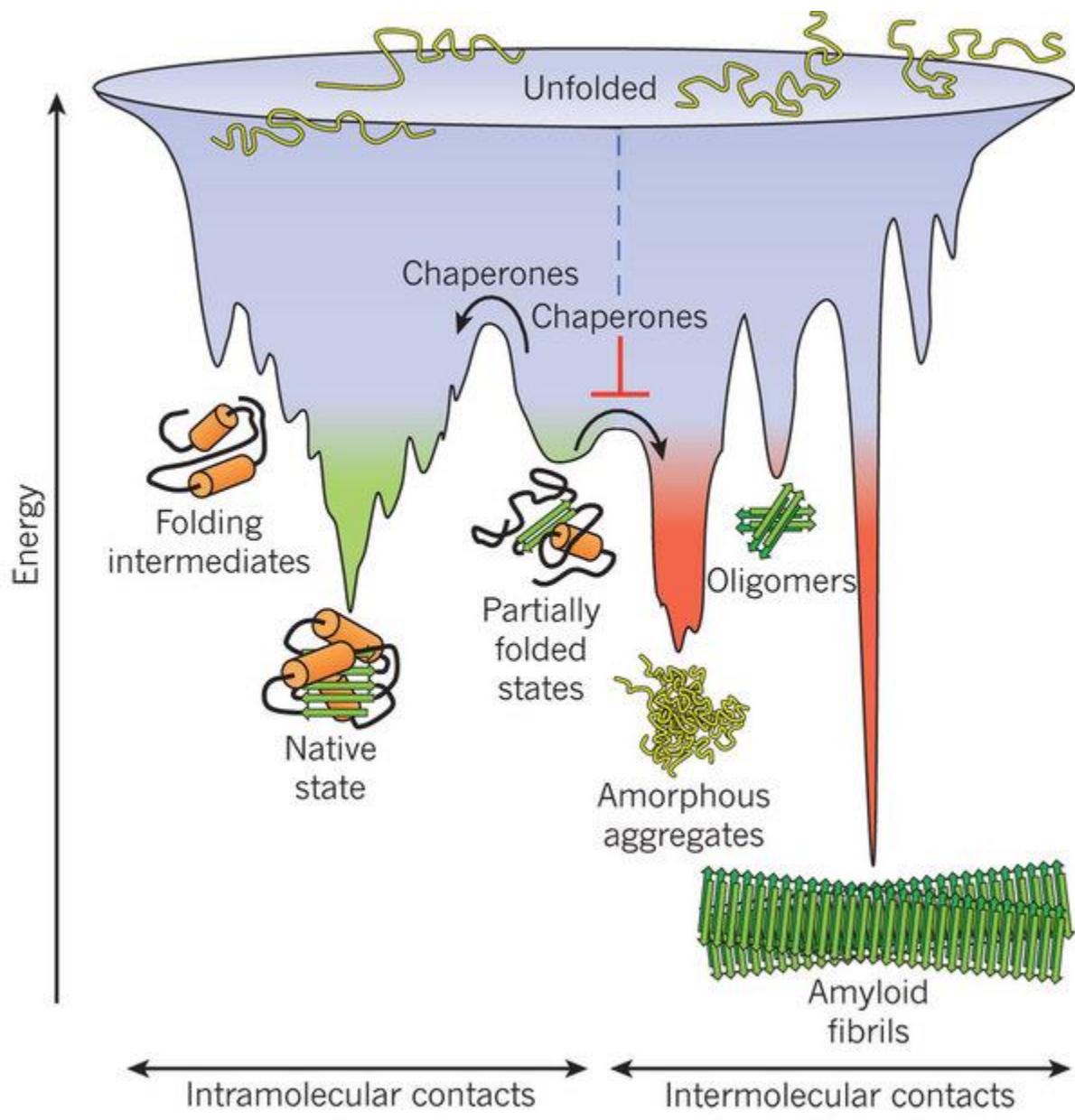
Christian Boehmer Anfinsen, Jr.
(March 26, 1916 – May 14, 1995)

Nobel Prize in Chemistry (1972)



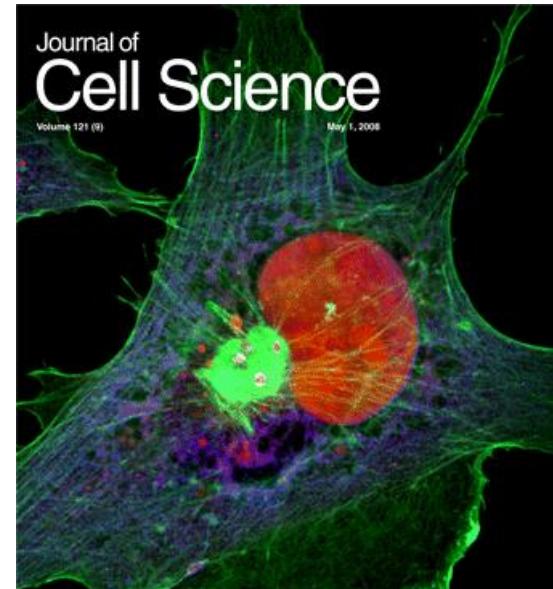
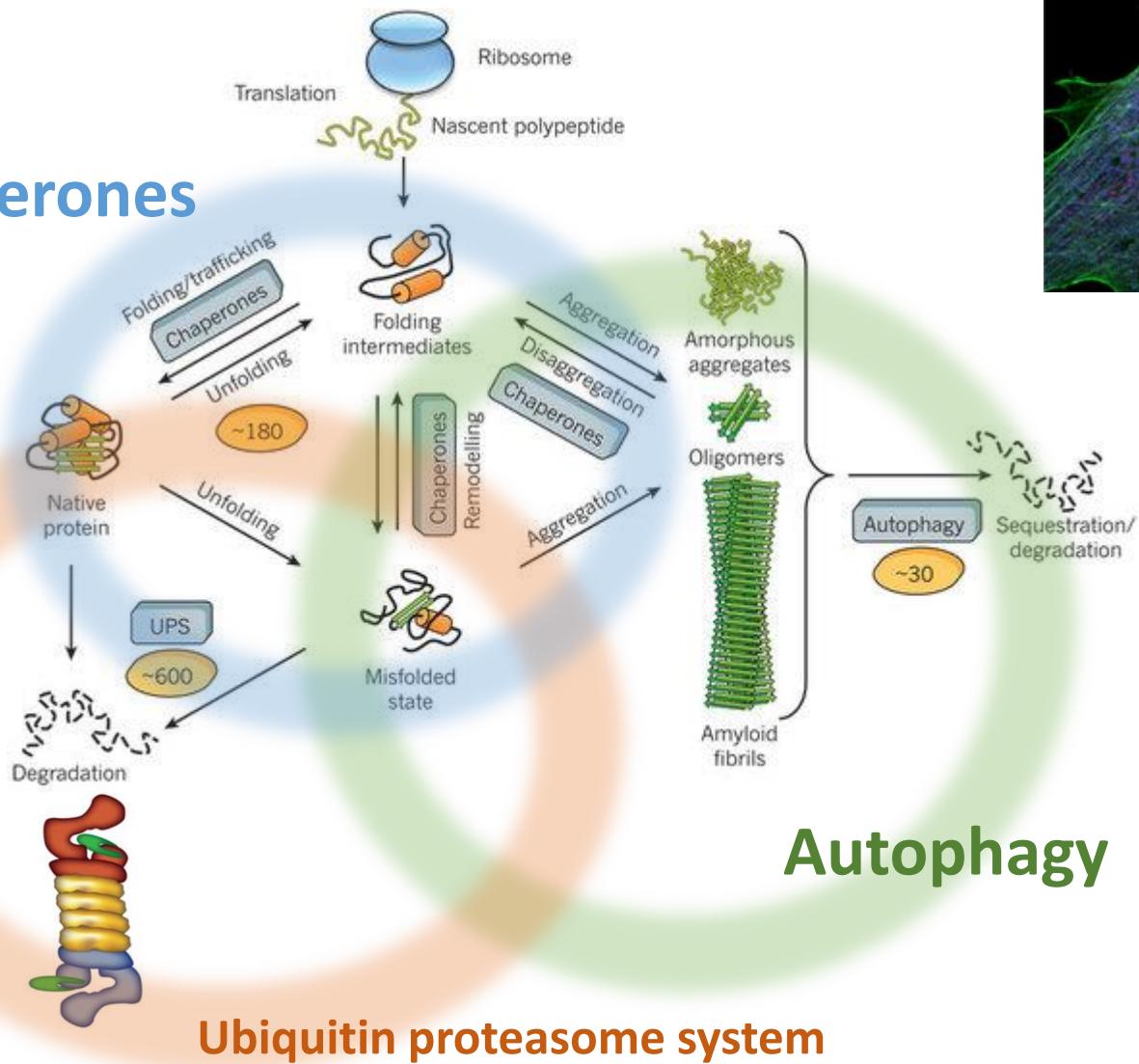
Folding is entropy driven process





Protein homeostasis / proteostasis

Chaperones

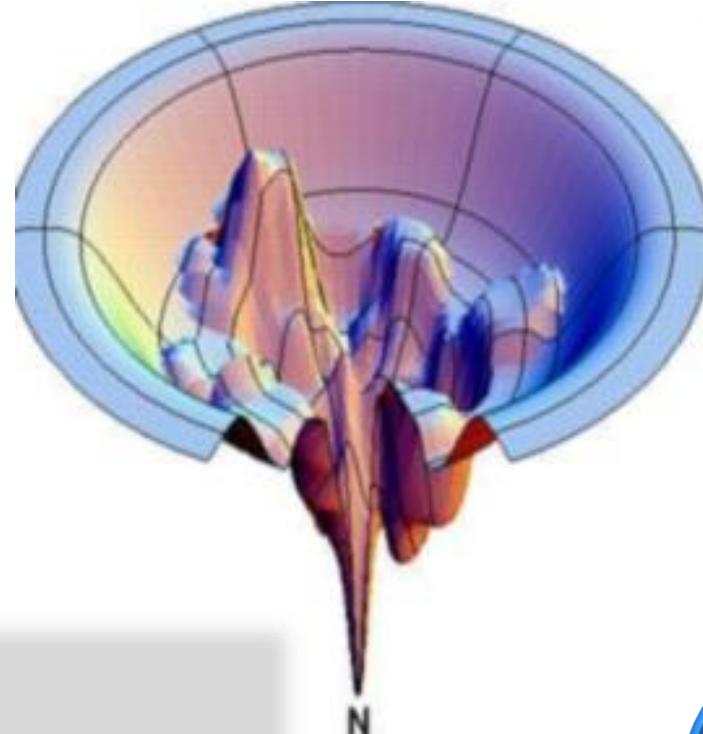
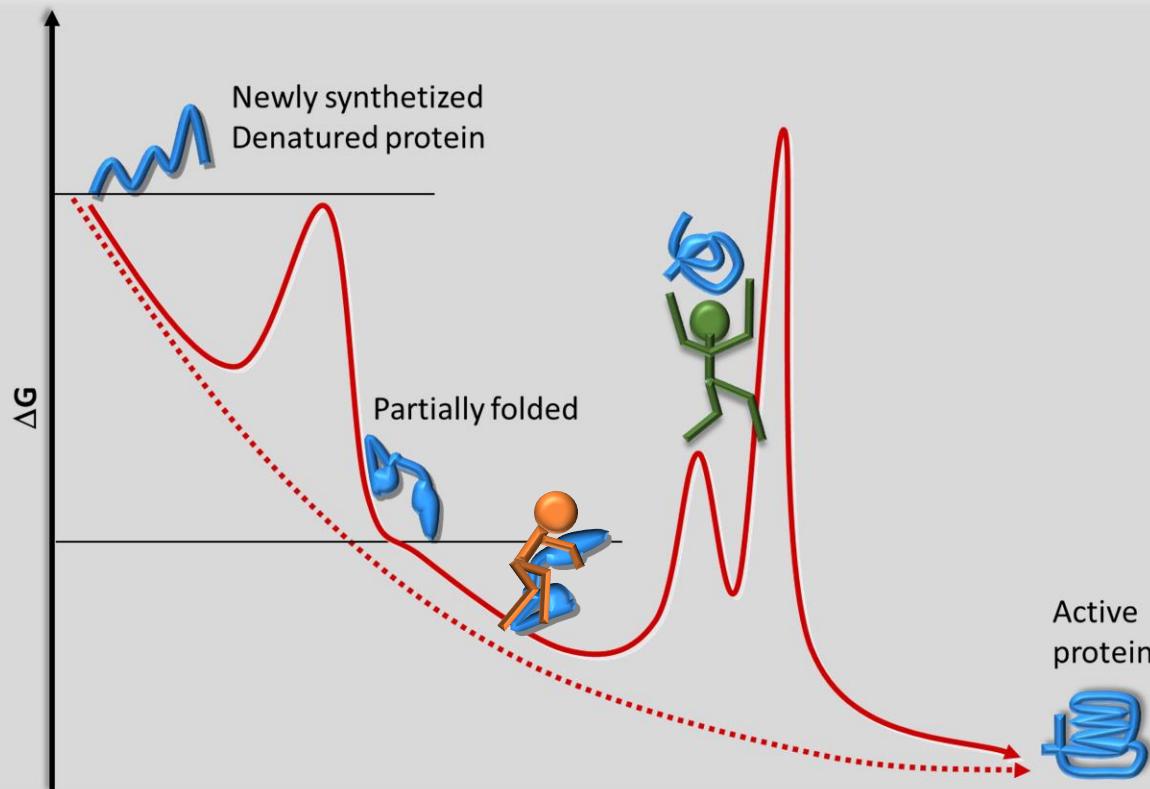


Stress proteins / Chaperones

Holdases bind folding intermediates to prevent their aggregation



Crystalins, p23, Hsp40...

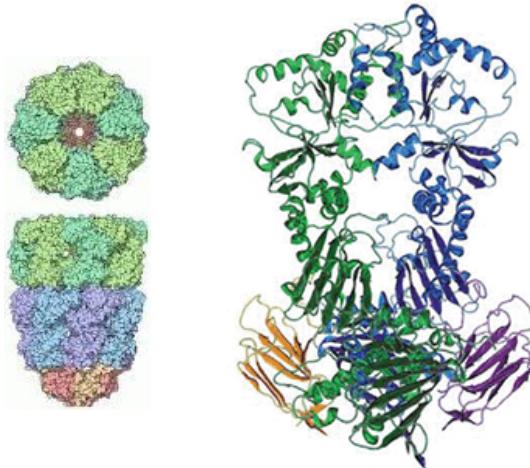


Foldases are chaperones that accompany other proteins to help them to overcome the energy barriers during folding to native conformation (ATP dependent)

Hsp70, Hsp90, GroEL...

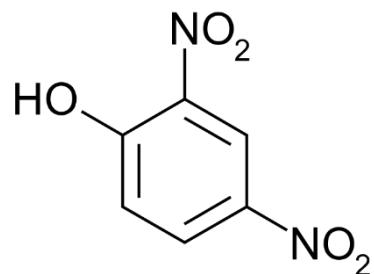
Stresové proteiny

Proteiny indukované tepelným stresem, převážně chaperony

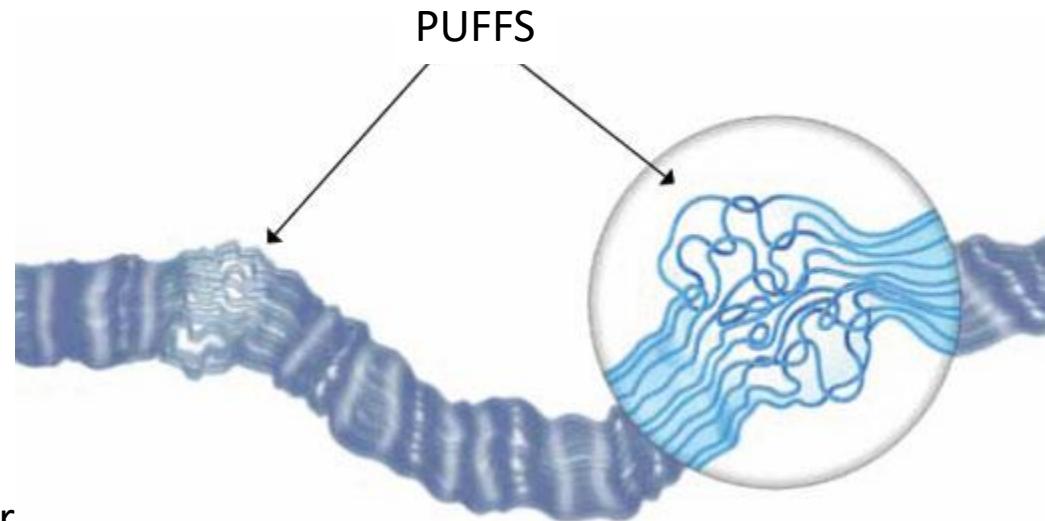


Hsp90
Hsp70, DnaK
Hsp60, GroEL
Hsp40 DnaJ
Hsp27, Crystalins,
Hsp10, chaperonins, GroES

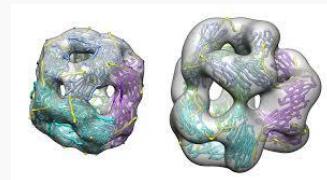
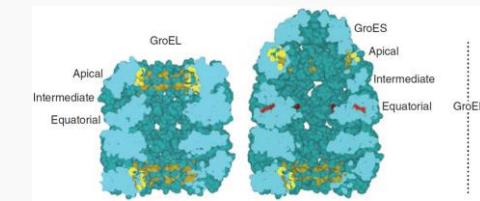
Teploplota



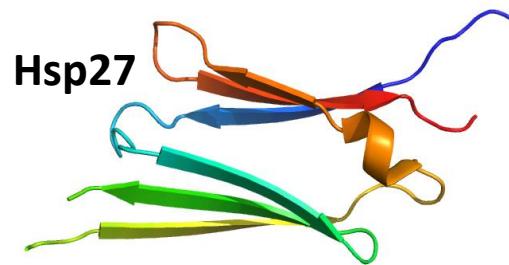
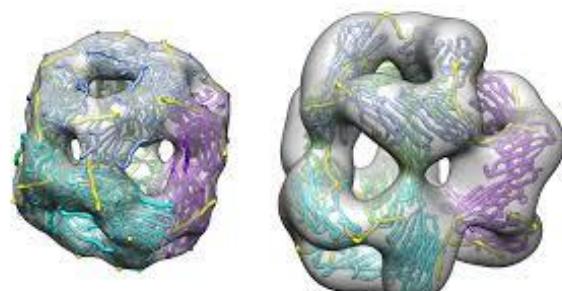
Dinitrophenol-mitochondrialí uncoupler



1962 by Ferruccio Ritossa

<u>Approximate molecular weight(kDa)</u>	<u>Prokaryotic</u> proteins	<u>Eukaryotic</u> proteins	Function
<u>10 kDa</u>	GroES	Hsp10	
20-30 kDa	GrpE	The HspB group of Hsp. Eleven members in mammals including Hsp27 , HSPB6 or HspB1 ^[28]	
<u>40 kDa</u>	DnaJ	Hsp40	Co-factor of Hsp70
<u>60 kDa</u>	GroEL, 60kDa antigen	Hsp60 	Involved in protein folding after its post-translational import to the mitochondrion/chloroplast
<u>70 kDa</u>	DnaK	The HspA group of Hsp including Hsp71, Hsp70 , Hsp72 , Grp78 (BiP), Hsx70 found only in primates	Protein folding and unfolding, provides thermotolerance to cell on exposure to heat stress. Also prevents protein folding during post-translational import into the mitochondria/chloroplast.
<u>90 kDa</u>	HtpG, C62.5	The HspC group of Hsp including Hsp90, Grp94	Maintenance of steroid receptors and transcription factors
100 kDa	ClpB, ClpA, ClpX	Hsp104, Hsp110	Tolerance of extreme temperature

HspB group/ small chaperones



Crystallins

Small Hsps

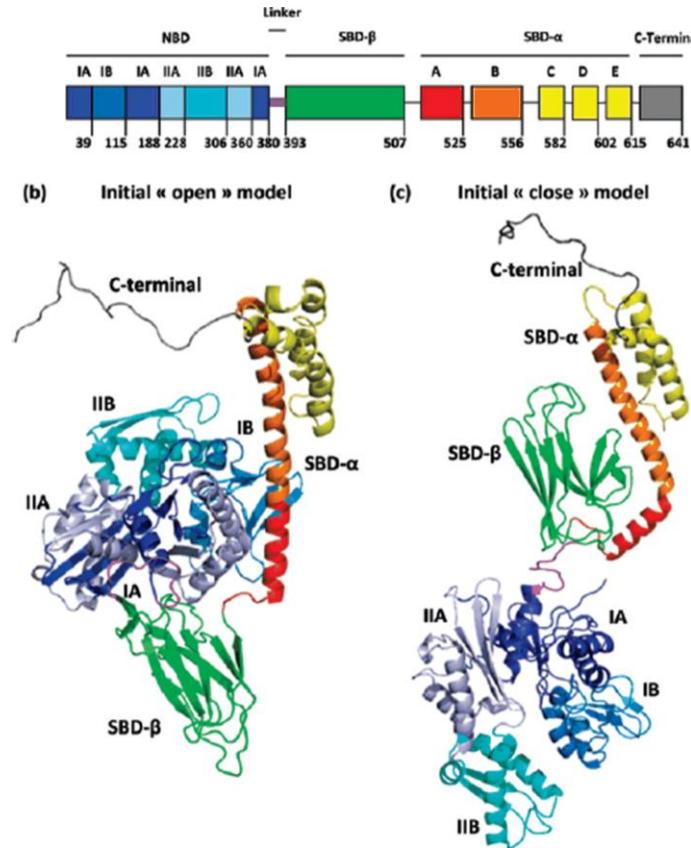
Ubiquitin-like

Prevent aggregation
Thermotolerance

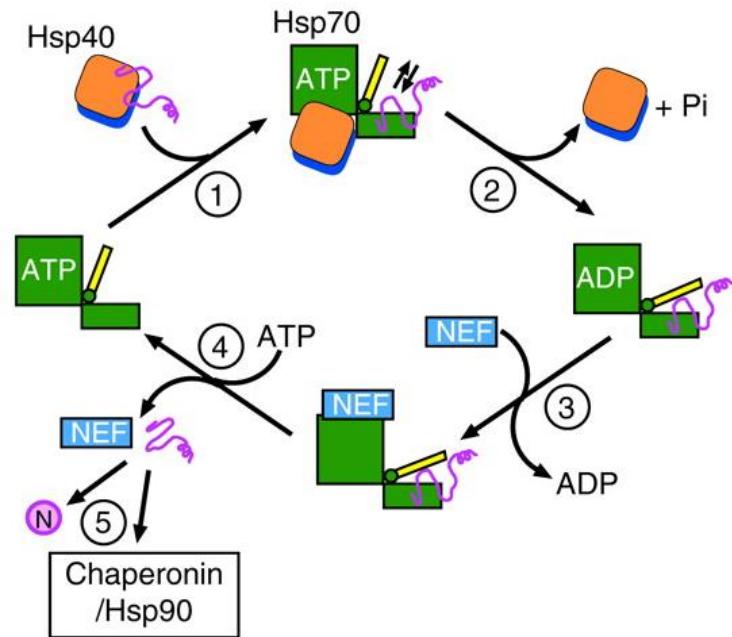
	Gene name	Protein name	Old names	Human gene ID	Mouse ortholog ID
1	<i>HSPB1</i>	HSPB1	CMT2F; HMN2B; HSP27; HSP28; HSP25; HS.76067; DKFZp586P1322	3315	15507
2	<i>HSPB2</i>	HSPB2	MKBP; HSP27; Hs.78846; LOH11CR1K; MGC133245	3316	69253
3	<i>HSPB3</i>	HSPB3	HSPL27	8988	56534
4	<i>HSPB4^a</i>	HSPB4	crystallin alpha A; CRYAA, CRYA1	1409	12954
5	<i>HSPB5^a</i>	HSPB5	crystallin alpha B, CRYAB; CRYA2	1410	12955
6	<i>HSPB6</i>	HSPB6	HSP20; FLJ32389	126393	243912
7	<i>HSPB7</i>	HSPB7	cVHSP; FLJ32733; DKFZp779D0968	27129	29818
8	<i>HSPB8</i>	HSPB8	H11; HMN2; CMT2L; DHMN2; E2IG1; HMN2A; HSP22	26353	80888
9	<i>HSPB9</i>	HSPB9	FLJ27437	94086	75482
10	<i>HSPB10^a</i>	HSPB10	ODF1; ODF; RT7; ODF2; ODFP; SODF; ODF27; ODFPG; ODFPGA; ODFPGB; MGC129928; MGC129929	4956	18285
11	<i>HSPB11</i>	HSPB11	HSP16.2; Clorf41; PP25	51668	72938

Hsp70 (DnaK, Grp78,..) chaperone machinery

ATP



ADP



BAG

NEF-Nucleotide exchange factor

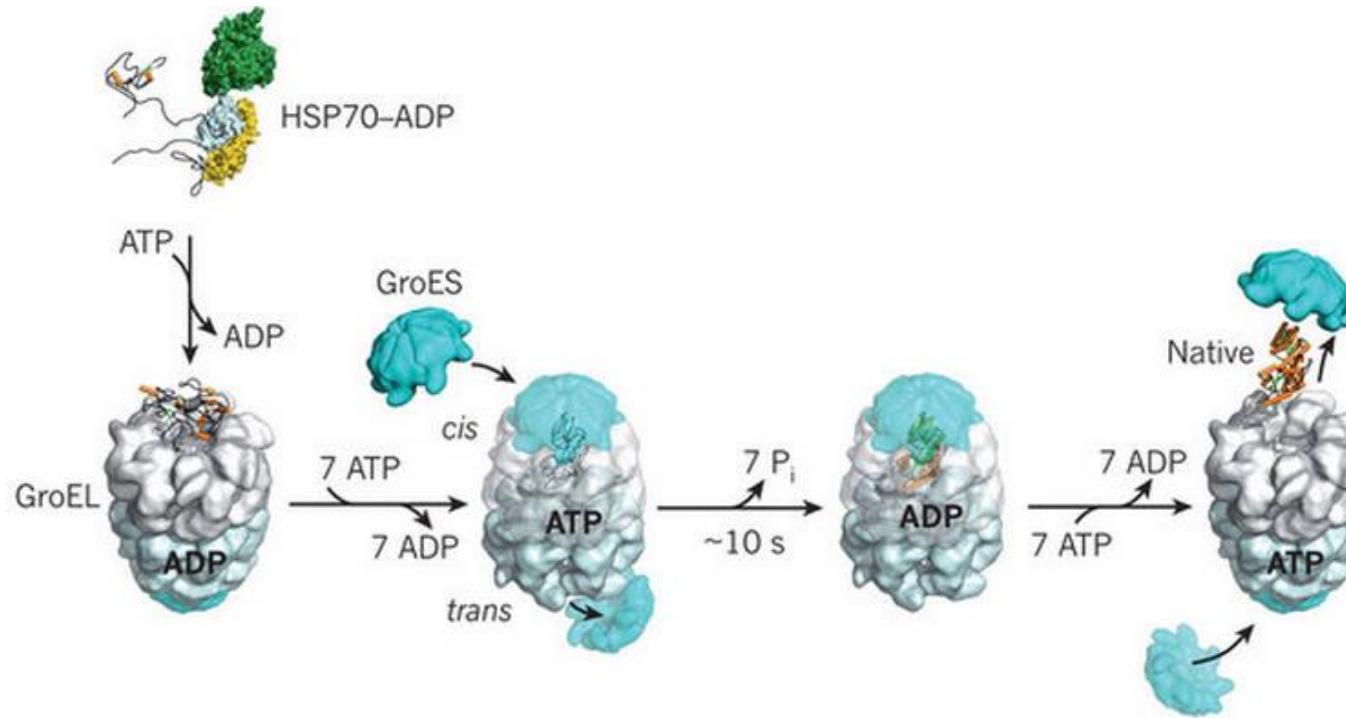
Hsp40

DnaJ

J-proteins

Chaperonins

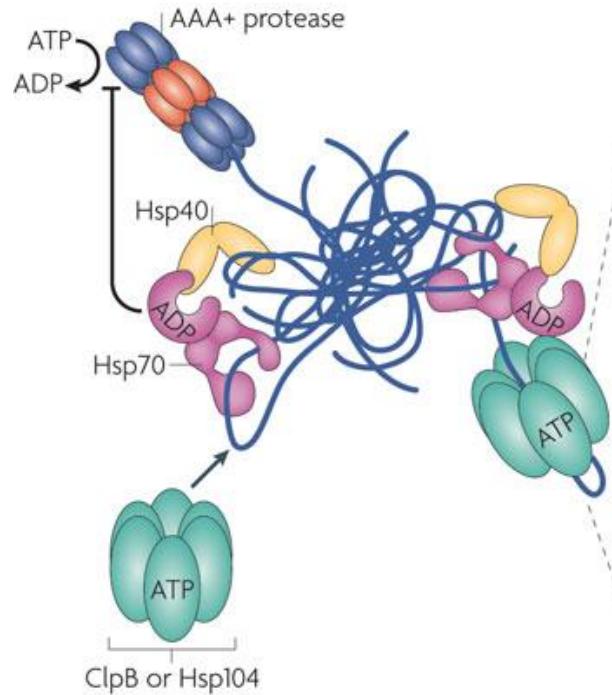
(GroEL-GroES, Hsp60, CCT-TRiC)



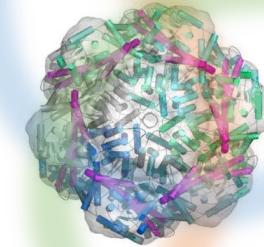
Folding of cytoskeletal proteins (tubulin)
Protein transport

Hsp104 (ClpB, ClpX,...)

Thermotolerance
Aggregate refolding
Prion folding (yeast Psi⁺/-)



Proteasome



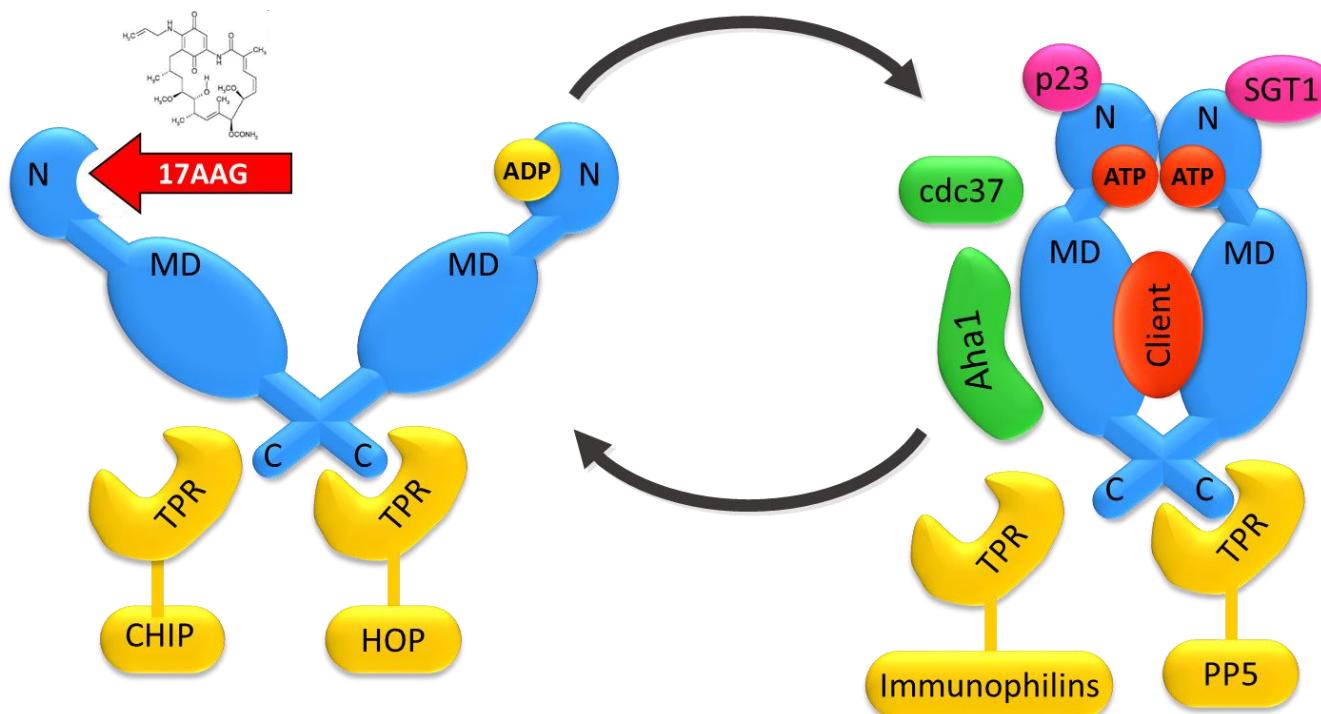
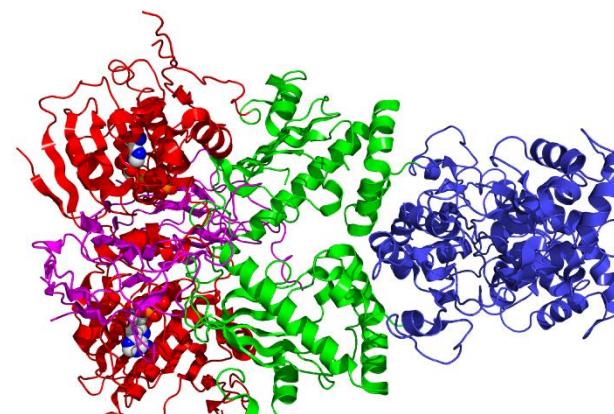
AAA+ proteases

AAA+ ATPases

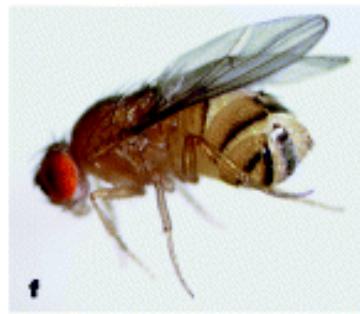
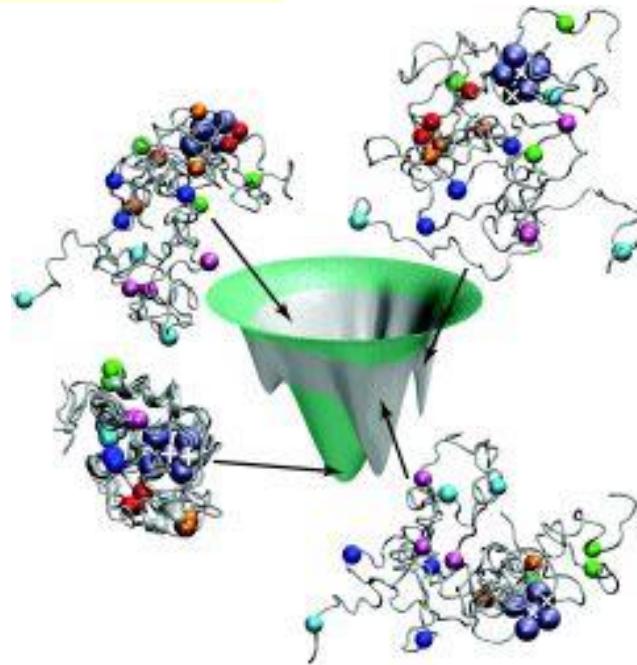
Converts ATP to
“mechanical” energy
(molecular motors)

Hsp90 chaperone machinery

- Conserved from prokaryotes to mammals
- ATPase activity (like gyrase)
- Mitochondrial, ER, cytoplasmic
- Redundant isoforms



Stress proteins/ Chaperones/Hsp90



Hsp90 as a capacitor for morphological evolution

Suzanne L. Rutherford*† & Susan Lindquist*

*Howard Hughes Medical Institute, University of Chicago, 5841 South Maryland Avenue MC1028, Chicago, Illinois 60637, USA

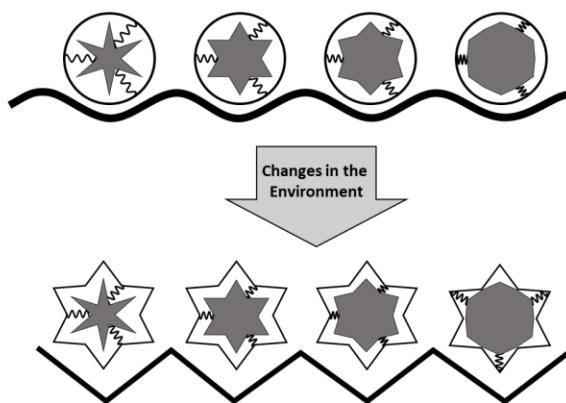
NATURE | VOL 396 | 26 NOVEMBER 1998 | www.nature.com

CHAPERONES AND EVOLUTION

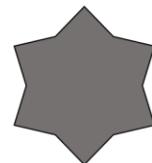
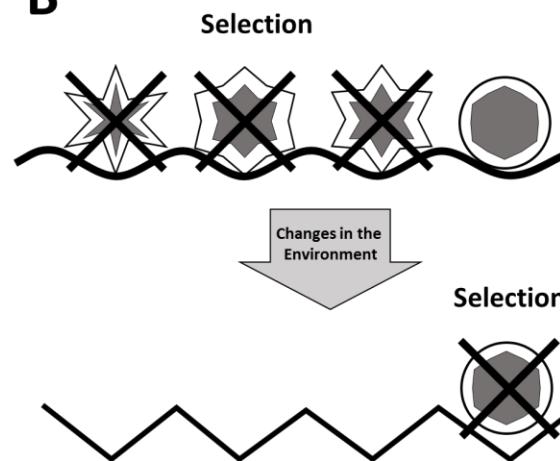
FILIP TRCKA, BORIVOJ VOJTESEK, PETR MULLER

Regional Centre for Applied Molecular Oncology, Masaryk Memorial Cancer Institute,
Zluty kopec 7, 656 53 Brno

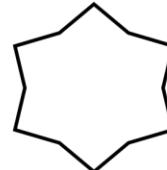
A



B



Genotype

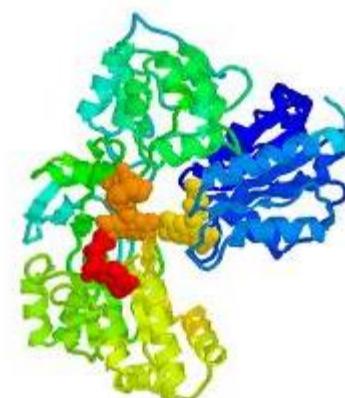
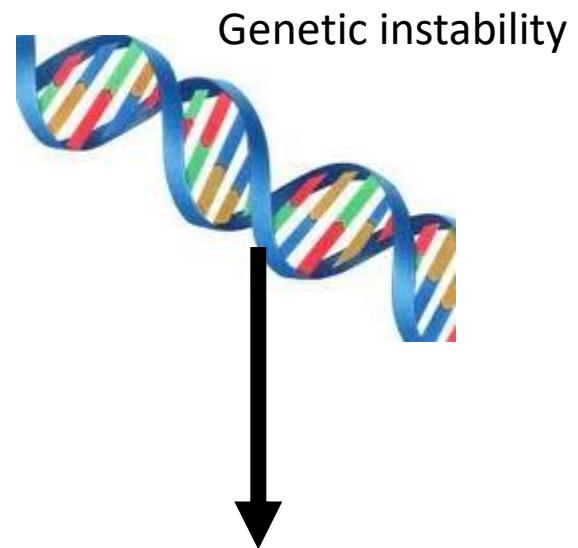


Phenotype



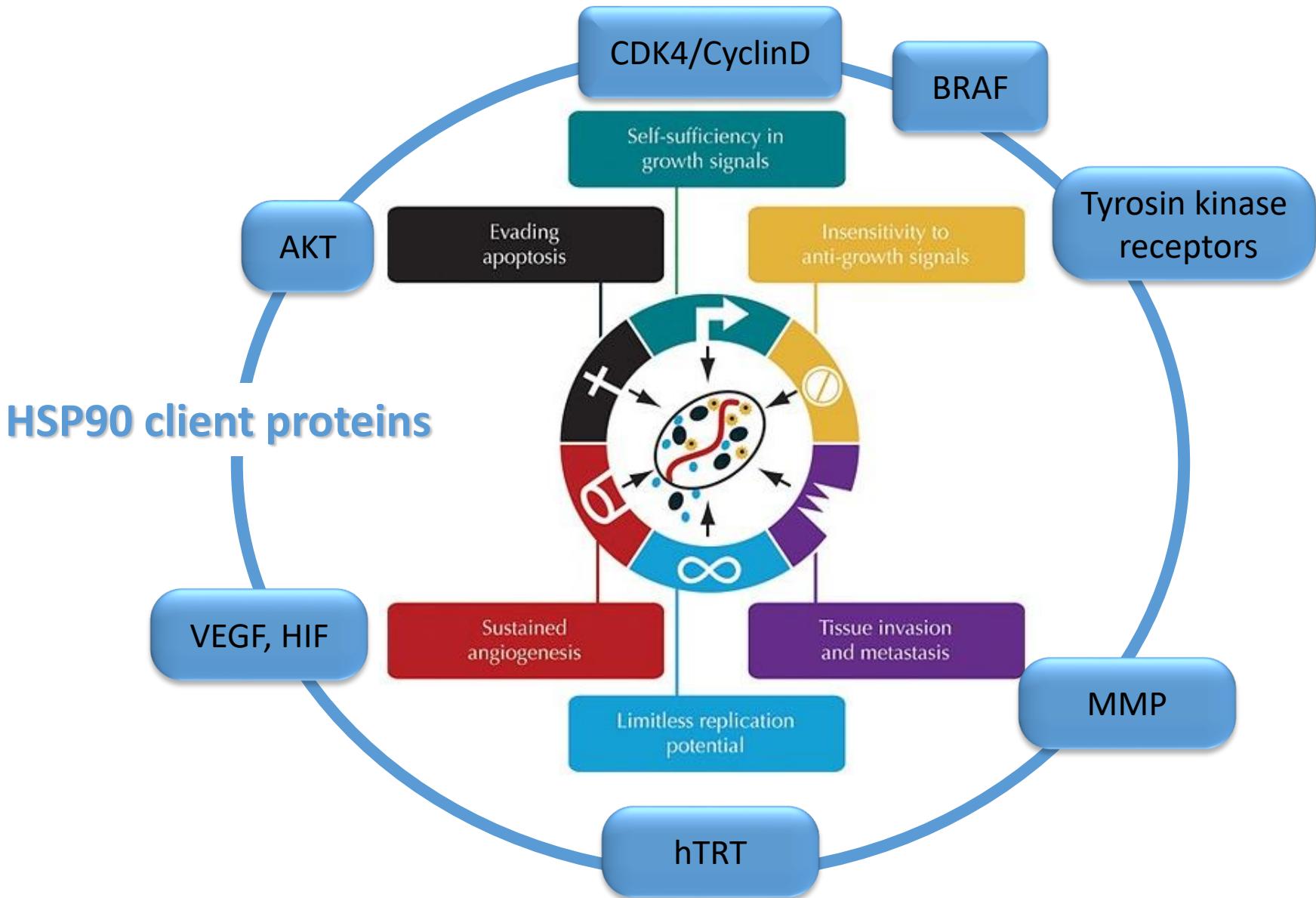
The activity of
molecular chaperones

The tumor cells demand high quality and amount of protein



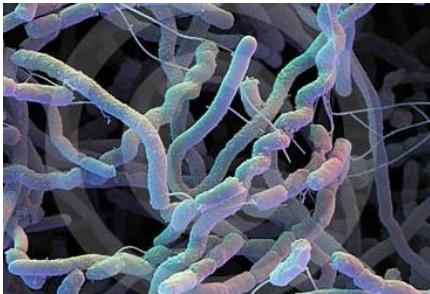
Hanahan D, Weinberg RA.: Cell. 2000 Jan 7;100(1):57-70.

Activity of Hsp90 is essential for expression of cancer phenotype

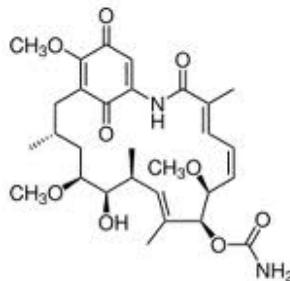


Specific inhibitors Hsp90

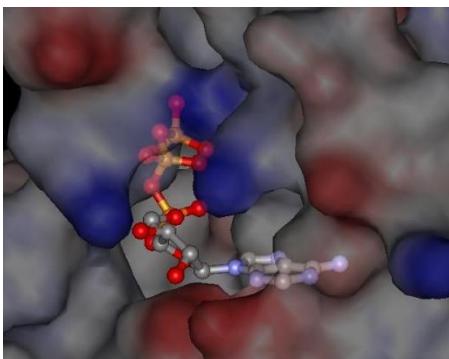
Clinical trials with Geldanamycin(2000)



Isolation of Geldanamycin (1970)



Geldanamycin binds ATP cavity of Hsp90 (1997)

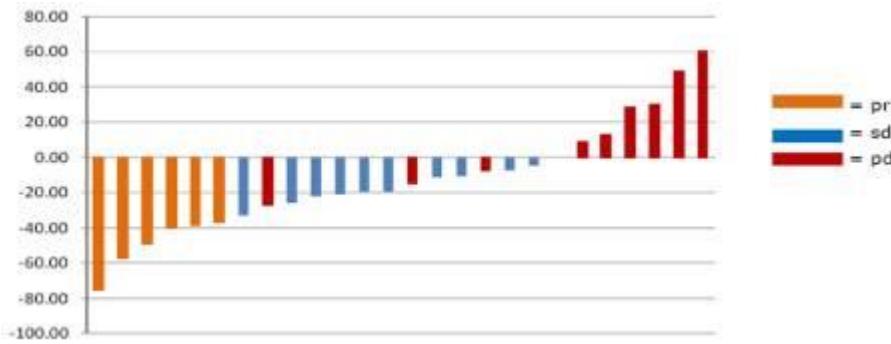
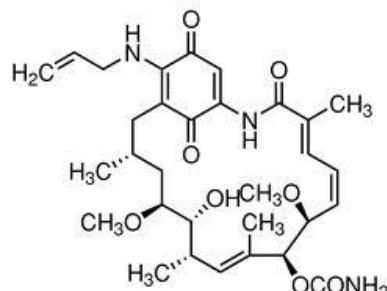


	inhibitor	No of studies	phase	Company
1	tanespimycin (17AAAG)	36	III	Bristol-Myers Squibb, Kosan
2	retaspimycin (IPI-504)	11	II/III*	Infinity Pharmaceuticals
3	alvespimycin (17DMAG)	7	II	Bristol-Myers Squibb, Kosan
4	STA-9090	14	II	Synta Pharmaceuticals Corp.
5	AUY922	11	II	Novartis Pharmaceuticals
6	CNF2024 (BIIIB021)	7	II	Biogen Idec
7	SNX-5422	4	I	Pfizer, Serenex, Inc.
8	AT13387	3	I	Astex Therapeutics
9	KW-2478	2	I/II	Kyowa Hakko Kirin Pharma, Inc.
10	IPI-493	2	I	Infinity Pharmaceuticals
11	HSP990	2	I	Novartis Pharmaceuticals
12	MPC-3100	1	I	Myrexis Inc.
13	Debio 0932	1	I	Debiopharm S.A.
15	BIIIB028	1	I	Biogen Idec

Hsp90 is unique therapeutic target for anti-cancer therapy



more than 17 different molecules in clinical trials



Variable response
need for predictive markers



Different assembly of Hsp90 machinery ?

- posttranslational modifications
- expression pattern of co-chaperones

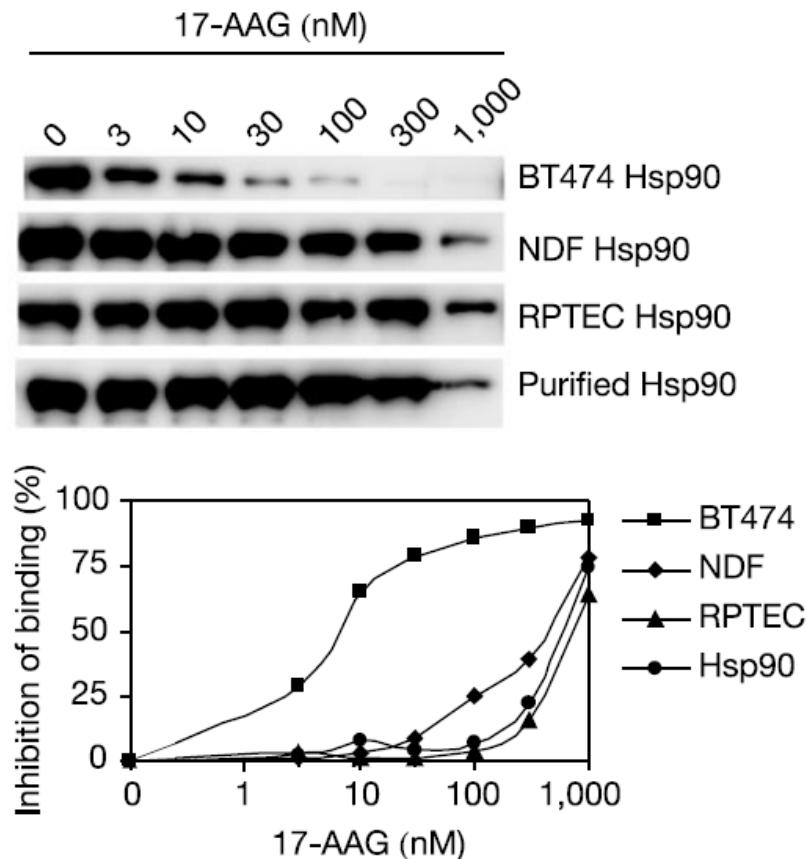
Client spectrum ?

What does kill the cells:

- apoptosis,
aggregation,

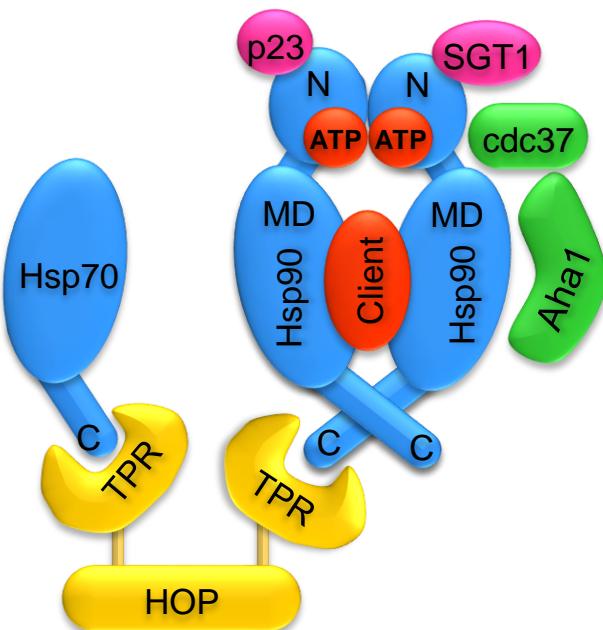
A high-affinity conformation of Hsp90 confers tumour selectivity on Hsp90 inhibitors

Adeela Kamal, Lia Thao, John Sensintaffar, Lin Zhang, Marcus F. Boehm,
Lawrence C. Fritz & Francis J. Burrows



Multichaperone complex

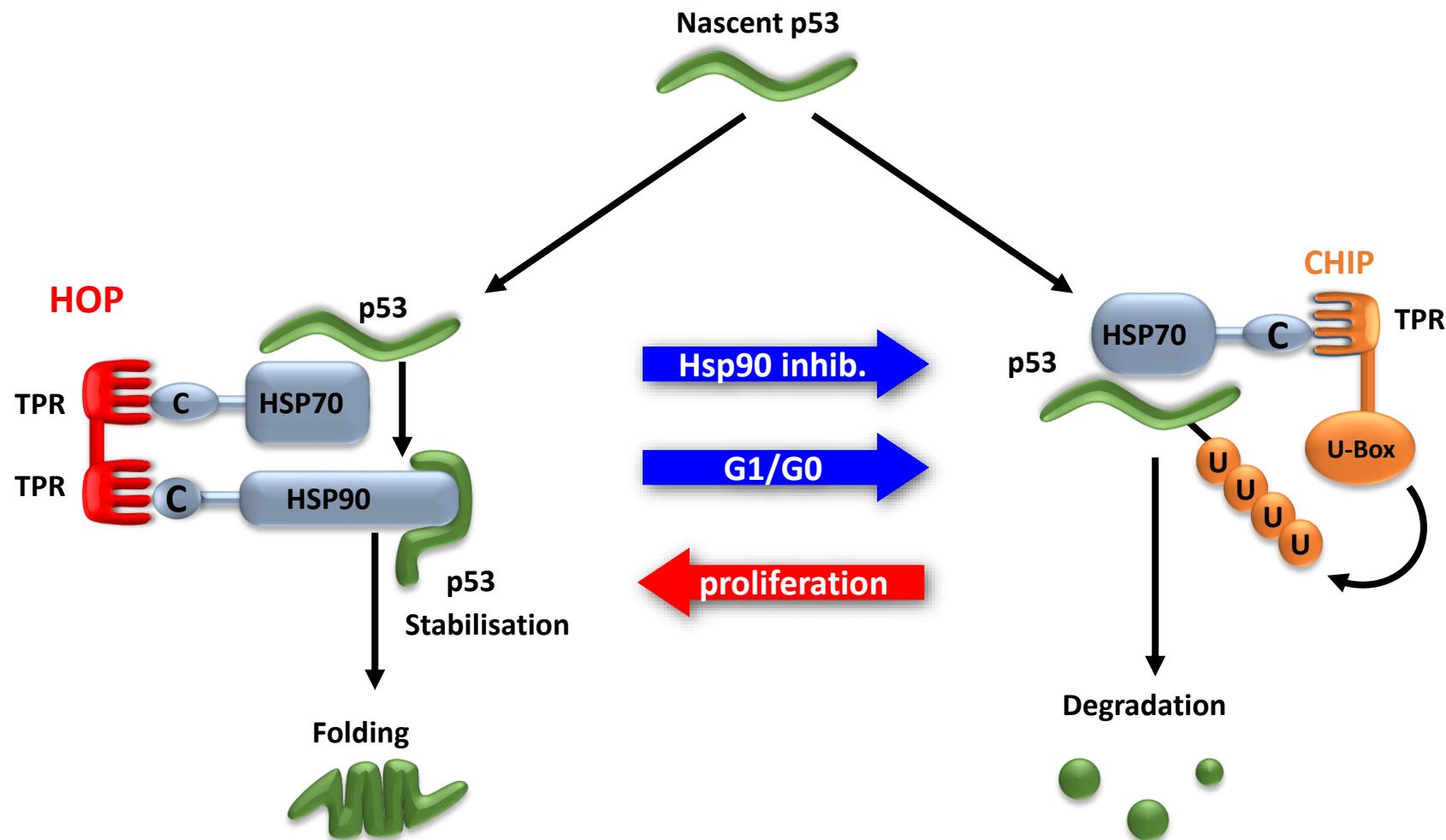
- Hsp90+Hsp70
- cochaperones



ORIGINAL ARTICLE

Chaperone-dependent stabilization and degradation of p53 mutants

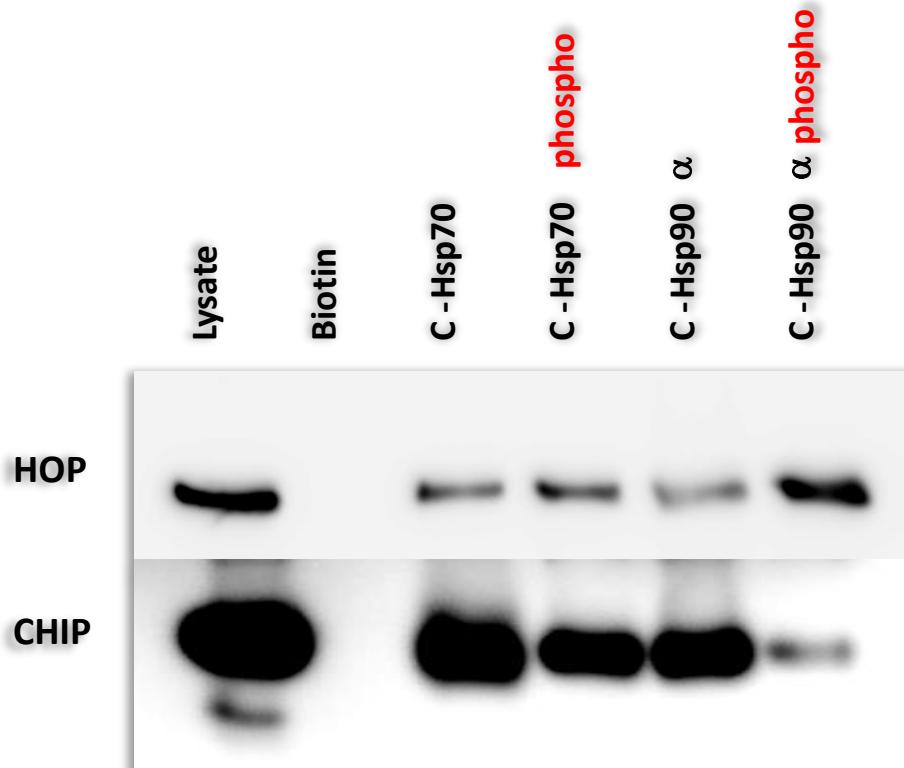
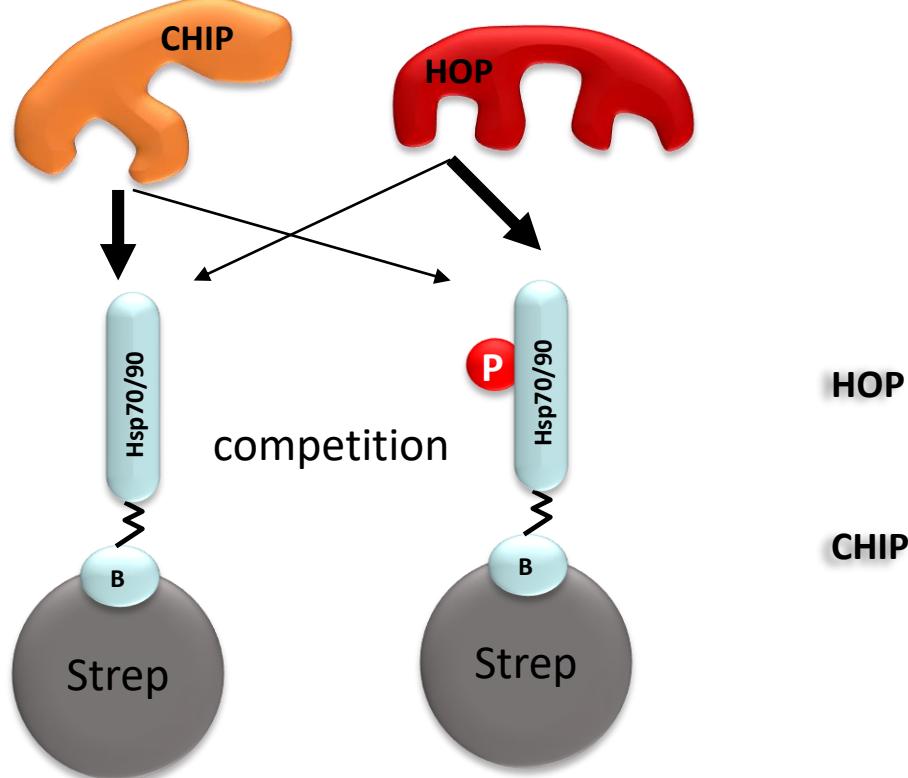
P Muller^{1,2}, R Hrstka¹, D Coomber², DP Lane² and B Vojtesek¹



What is the mechanism regulating folding/degradation balance?

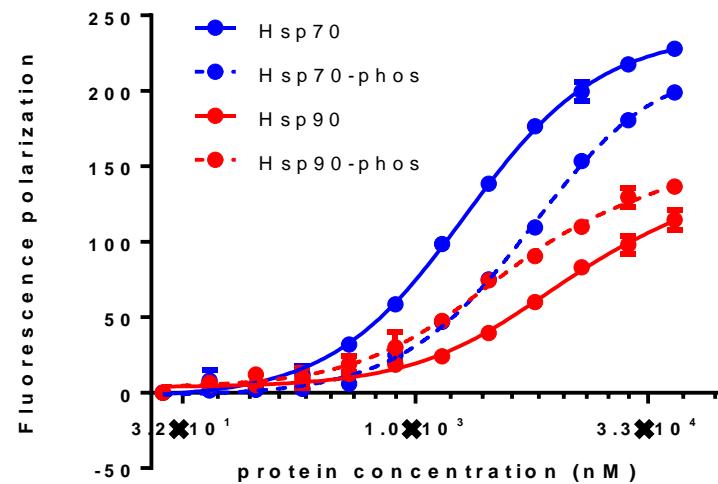
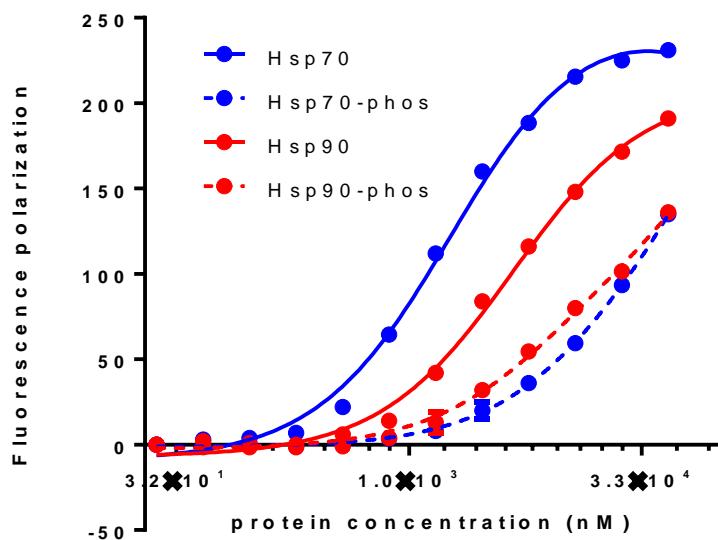
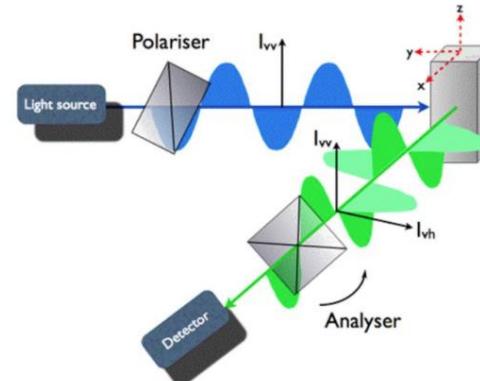
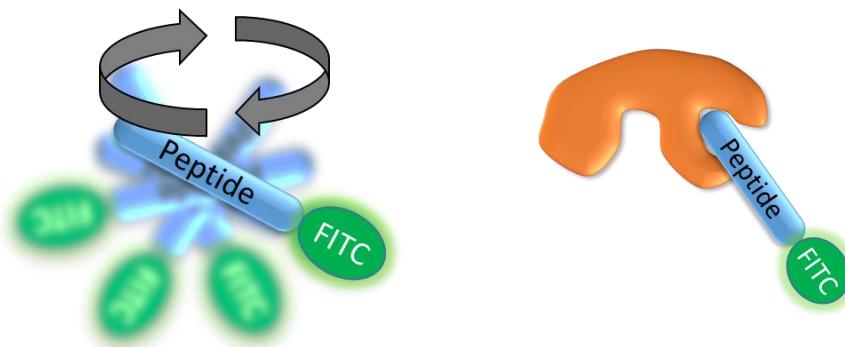
Cell lysate pulldown of HOP and CHIP

- Biotinylated phospho/non phospho peptides of Hsp70/Hsp90



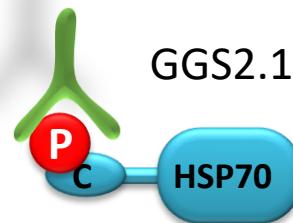
Detekce interakce proteinu s peptidem pomocí fluorescenční polarizace

Fluorescence lifetime – „doba mezi přijetím excitačního fotonu a vyzářením fotonu emitovaného“

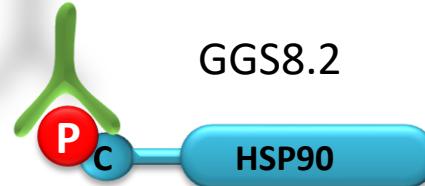


Detection of phosphorylated Hsp70 and Hsp90

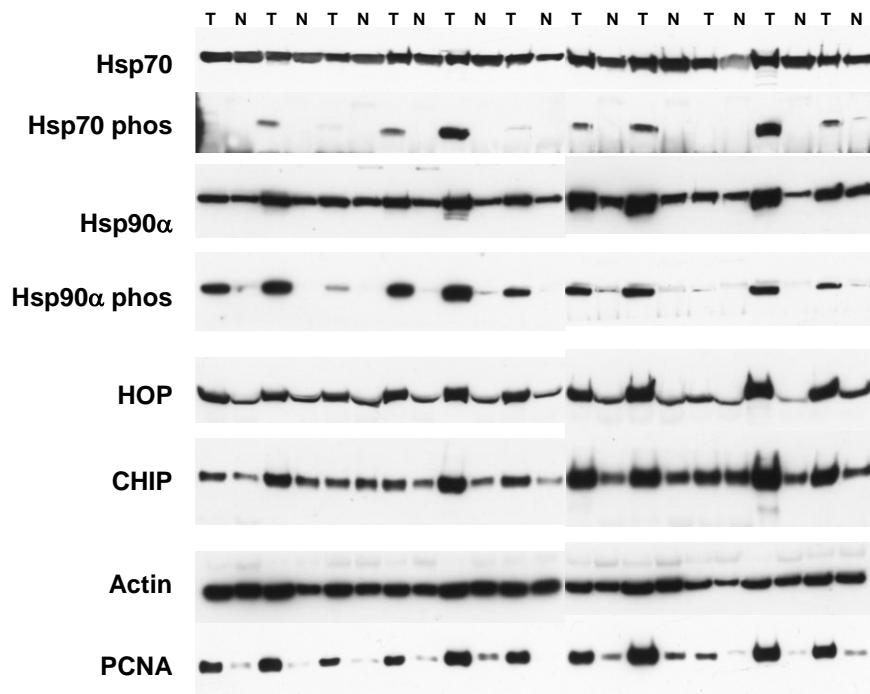
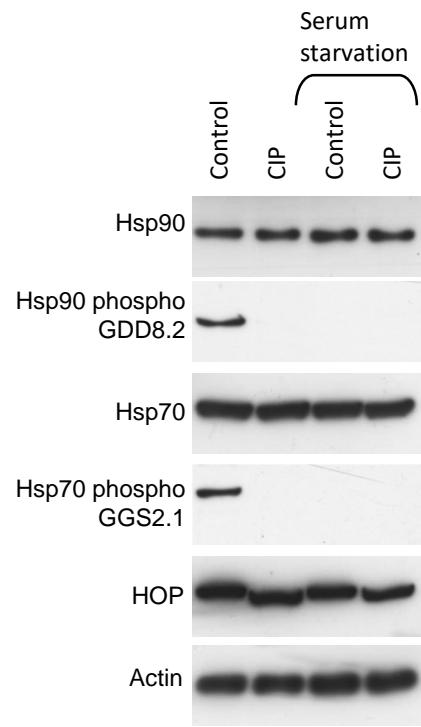
phospho-specific monoclonal antibodies



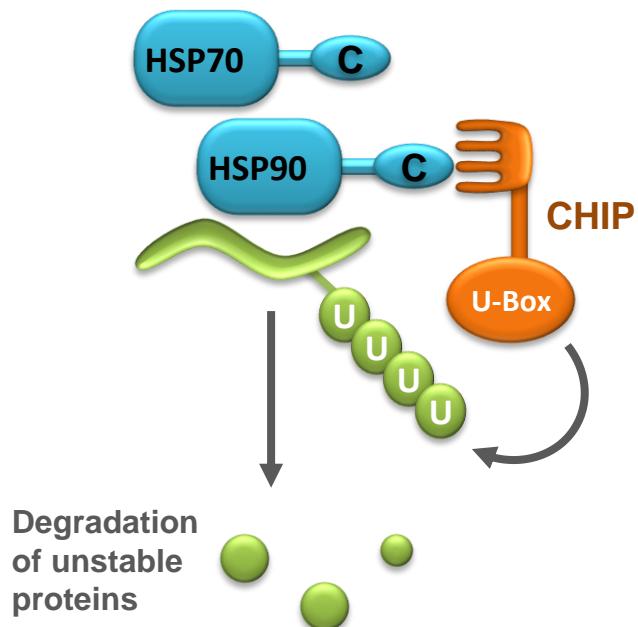
GGS2.1



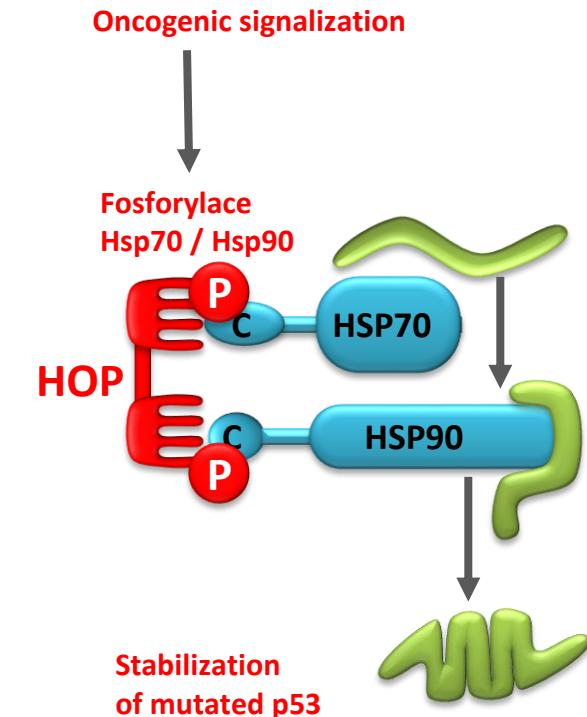
GGS8.2



Normal differentiated cell



Cancer cell

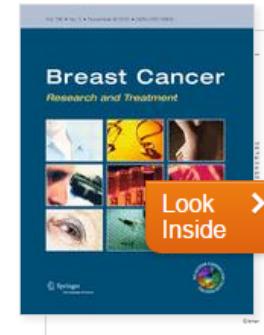


Normal differentiated cell	Cancer cell
C-terminus Hsp70/90 non phosphorylated	Phosphorylated Hsp90 Hsp70
Hsp bind preferentially CHIP	Hsps bind preferentially HOP
Designed to degrade unfolded protein	High folding capacity of Hsp90
Higher expression of CHIP	Increased level of HOP
Lower sensitivity to Hsp90 inhibitors	High sensitivity to Hsp90 inhibitors

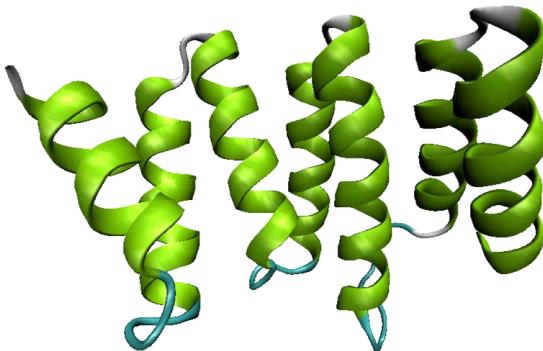
Date: 04 Oct 2012

TOMM34 expression in early invasive breast cancer: a biomarker associated with poor outcome

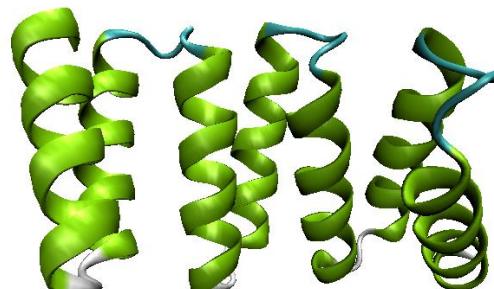
Mohammed A. Alekandarany, Ola H. Negm, Emad A. Rakha, Mohamed A. H. Ahmed, Christopher C. Nolan, Graham R. Ball, Carlos Caldas, Andrew R. Green, Patrick J. Tighe, Ian O. Ellis



TPR1



TPR2



Positively charged clamp

Positively charged clamp sequence alignment:

Tomm34-TPR1 Q15785 9-118	Helix 1A	Helix 1B
VEEL R AAG G NESFRNGQYAAEASALYGRALRVLQAQG-----		
ARV L KEEG N ELVKKGNHKKAI E KYSSESLLC-----		
AQE L KE Q G N RLFVGRKY P EAACYGRAITR-----		
VNE L KE G NKA S VGN I DDALQCYS E AIKL-----		
ALK E KEL G NDAYKKKD F DTALKHYDKAKE L -----		
STI V KERGT T YFKEGKY K QAL L QYKKIVSWLEYESSFSNE E AQ		
AEEL K TQ A N DYFKAKDY E NAIK F YSQ A IEL-----		

Positively charged clamp sequence alignment:

Tomm34-TPR1 Q15785 9-118	Helix 2A	Helix 2B
SSDP E EESVLYSNRAACHLKDGNCRDCIKDCTSALALVPFS-----		
- S N-LE S ATYSNRA C YLVLKQY T EA V KDC T EA L KLDGKN-----		
- N P-LV A V Y YTNRA C YL K M Q QHE Q ALAD C RR A LE L D G QS-----		
- D P-HN H VLYSNRS A AYAK K GD Y Q K AYED G C K TV D L K PDW-----		
- D P-TNMTY T TNQAA V Y F E K GD Y N K CRE L C E KA I E V GRE N R E		
KA Q A-LR L ASHLN L AM C HL K L Q A F S A AI E S C N K ALE L DSNN-----		
- N P-SNA I YYGNRS L AYL R TE C Y G Y A LG D AT R AI E LD K Y-----		

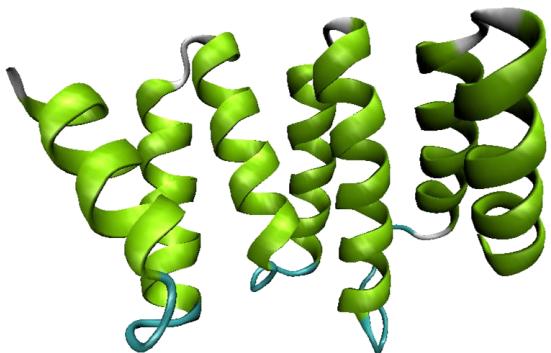
Positively charged clamp sequence alignment:

Tomm34-TPR1 Q15785 9-118	Helix 3A	Helix 3B
- I KP L LR R AS A YE A LE K YPM A Y V D K T V L Q IDD N V-----		
- V K A F Y R R QA H K A L K D Y K S S F A D I S N L Q I E PR N -----		
- V K A H F FL G QC Q L E M E S Y D E A I A N L Q R A Y S L A K E Q -----		
- G K G Y S R K AA A L E F L N R F E E A K R T Y E E G L K H E A N N -----		
D Y R Q I A K Y A R I G N S Y F K E E K Y K D A I H F Y N K S L A E H R T P -----		
- E K G L F R R G E A H L A V N D F E L A R A D F Q K V L Q L P NN -----		
- I K G Y Y R R A S N M A L G K F R A A L R D Y E T V V K V K P H -----		

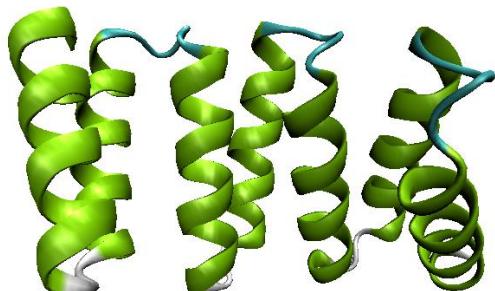
TOMM34 protein – co-chaperone

Tetratricopeptide repeat (TPR) domain

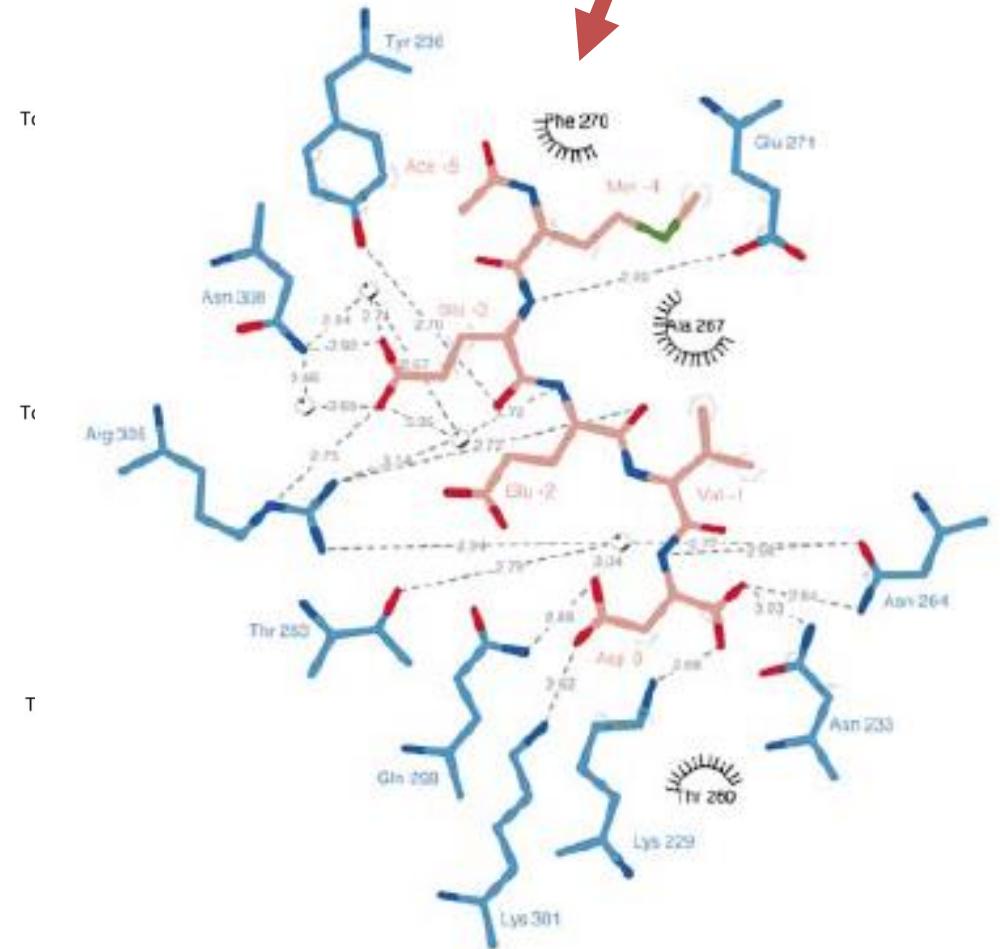
TPR1



TPR2



C-terminus Hsp70/Hsp90
EEVD

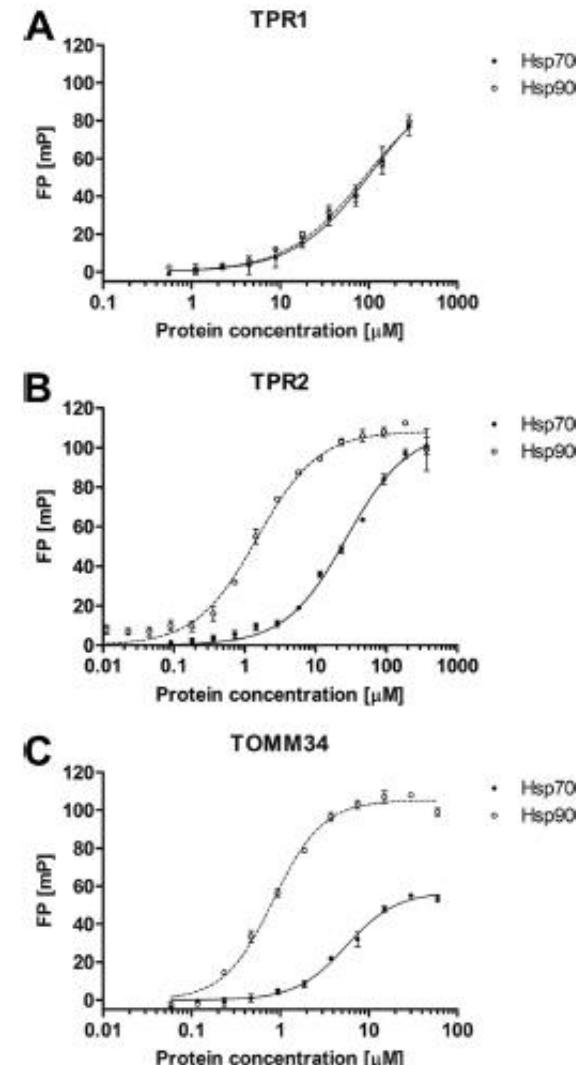
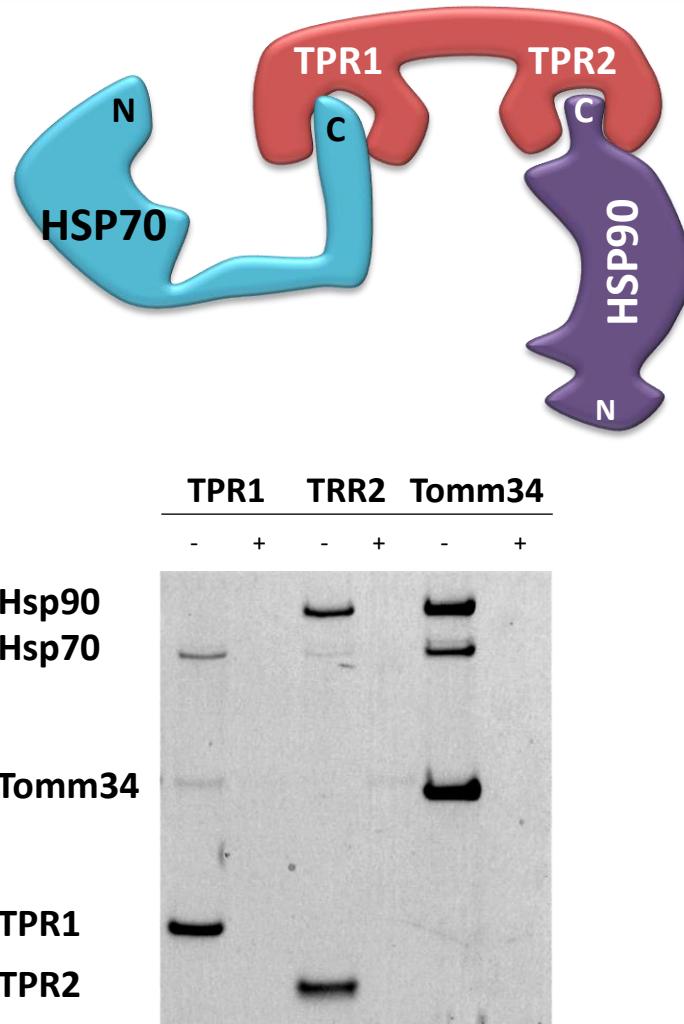


De novo modeled structure of **TOMM34** domains

The Assembly and Intermolecular Properties of the Hsp70-Tomm34-Hsp90 Molecular Chaperone Complex*

Received for publication, October 11, 2013, and in revised form, February 19, 2014. Published, JBC Papers in Press, February 24, 2014, DOI 10.1074/jbc.M113.526046

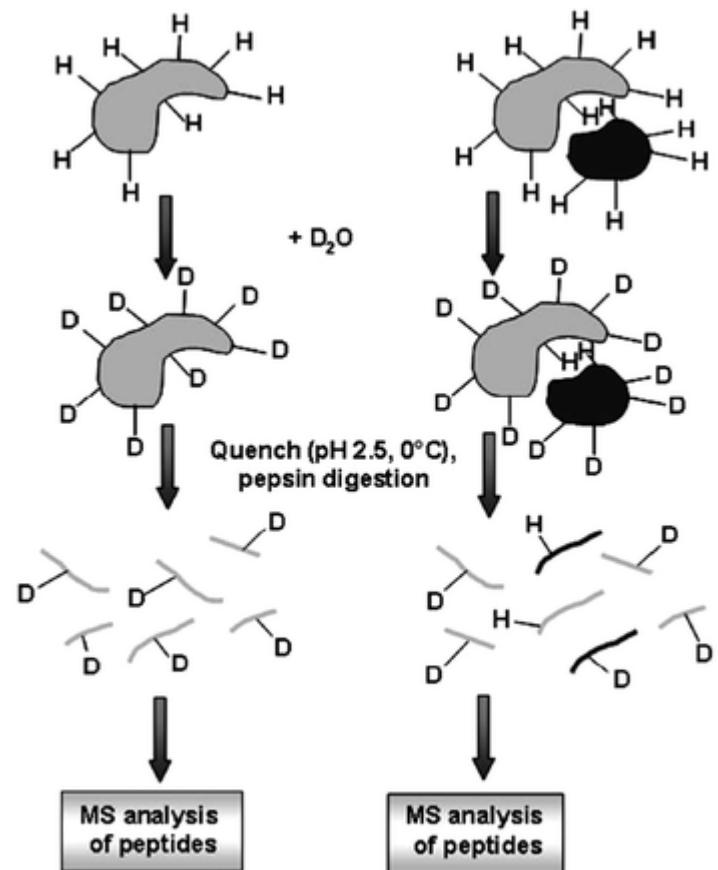
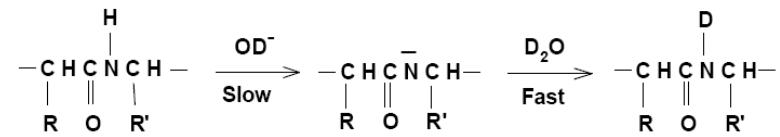
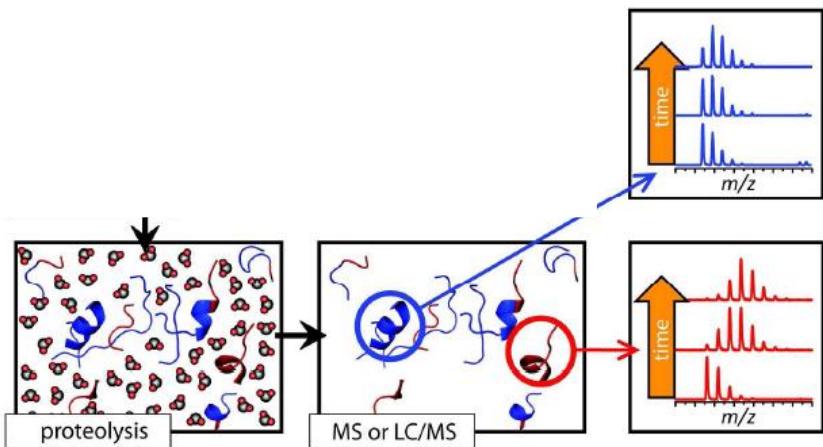
Filip Trcka[‡], Michal Durech[‡], Petr Man^{§¶}, Lenka Hernychova[‡], Petr Muller^{‡,1,2}, and Borivoj Vojtesek^{‡,1,3}



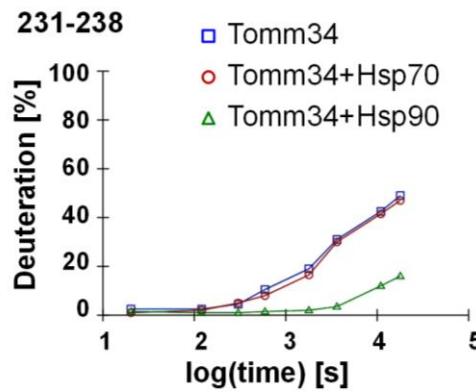
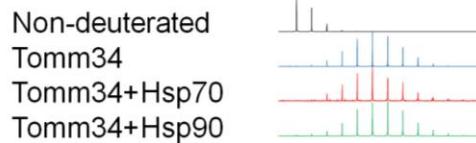
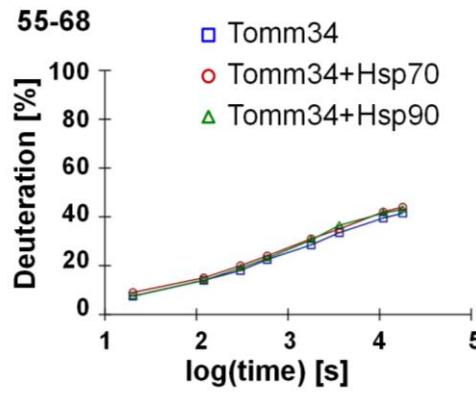
HDX – basic basics

Exchangeable hydrogens:

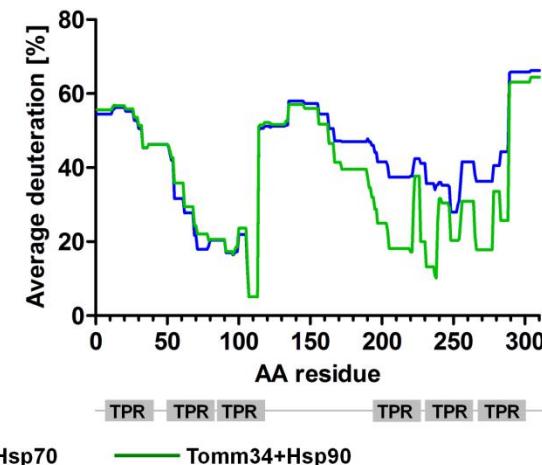
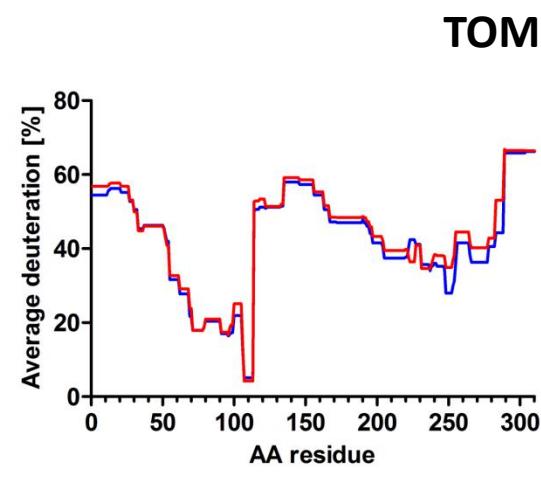
- 1) side chains containing $-OH$, $-SH$, $-NH_2$, $-COOH$ and $-CONH_2$ groups and hydrogens from the amino and carboxy termini
- 2) carbon-bound aliphatic and aromatic hydrogens
- 3) hydrogens arising from the amide linkages between amino acids of the protein polypeptide chain



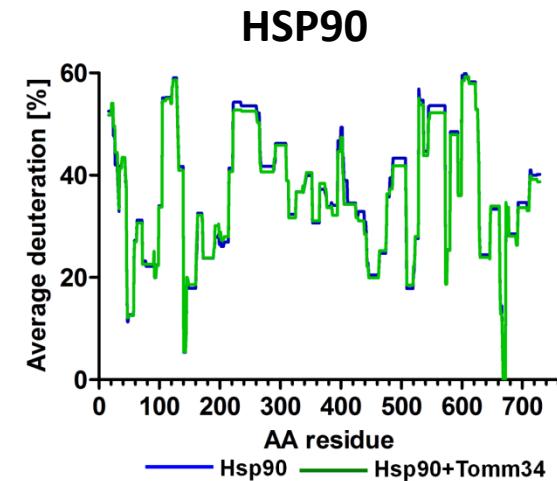
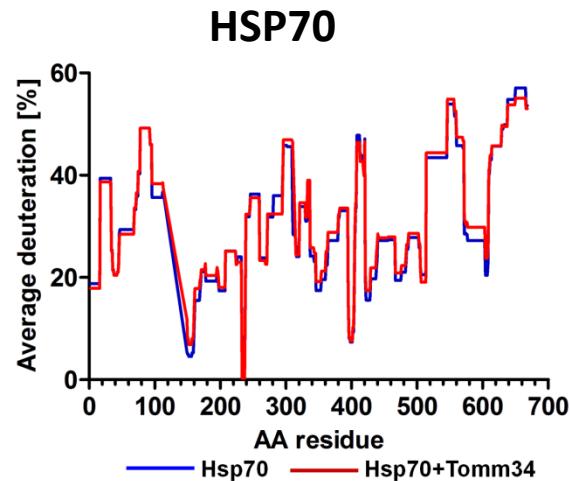
HDX – HSP70/90-TOMM34 interaction without ATP



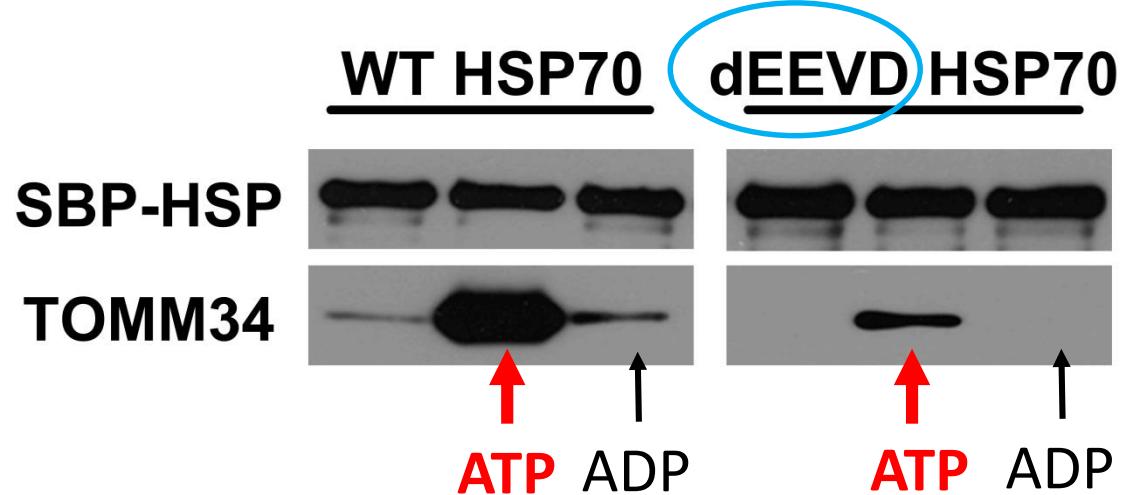
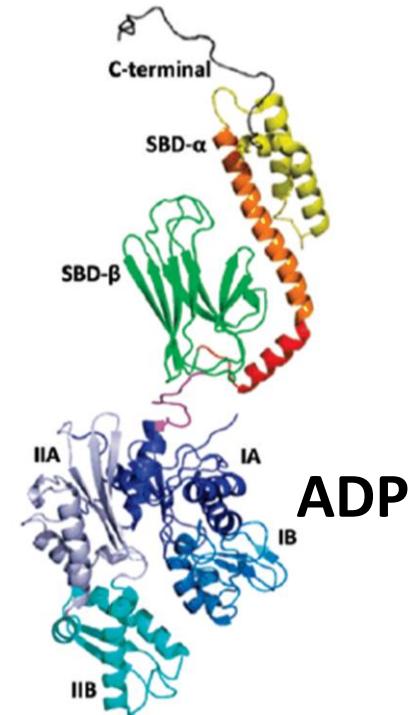
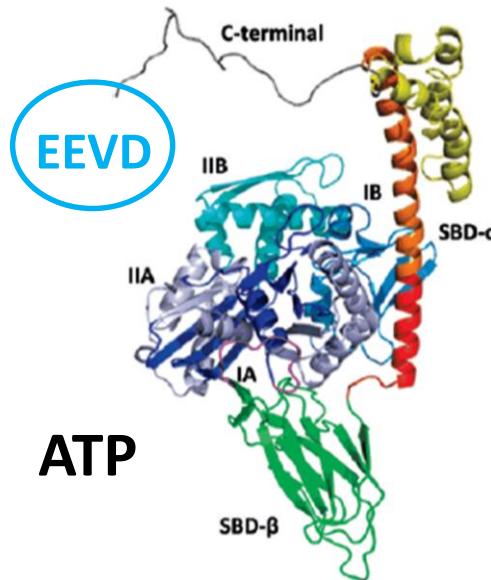
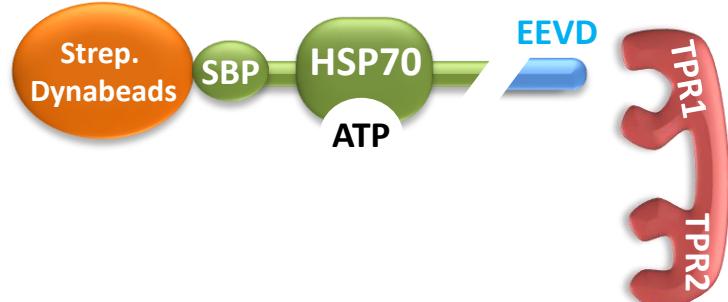
A



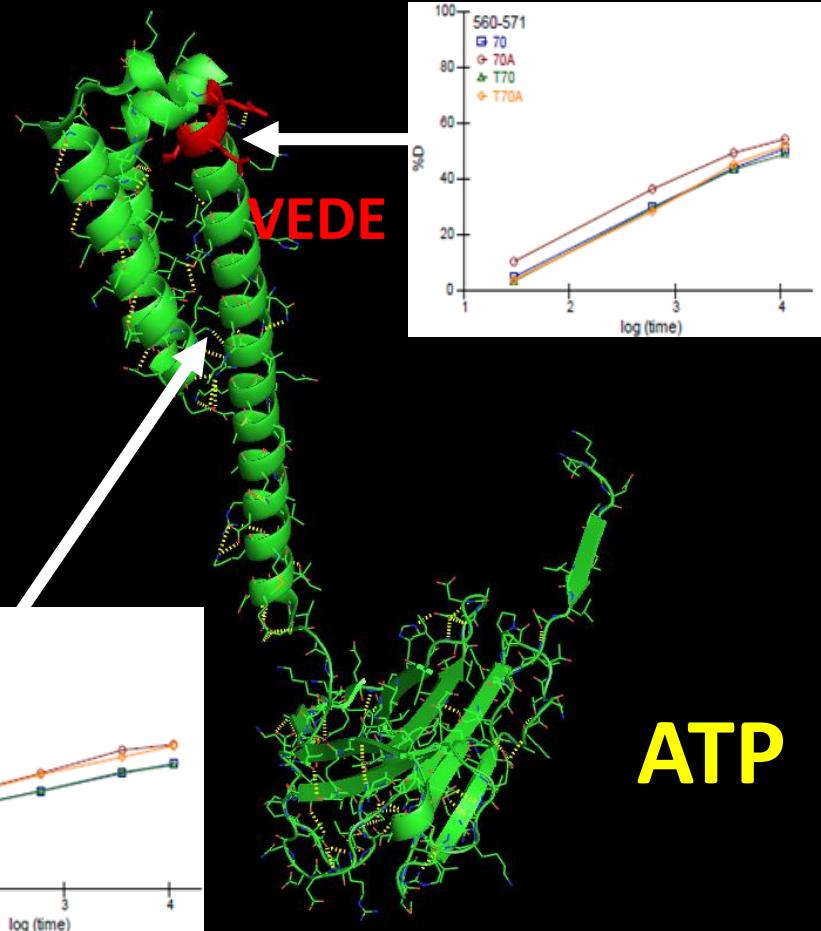
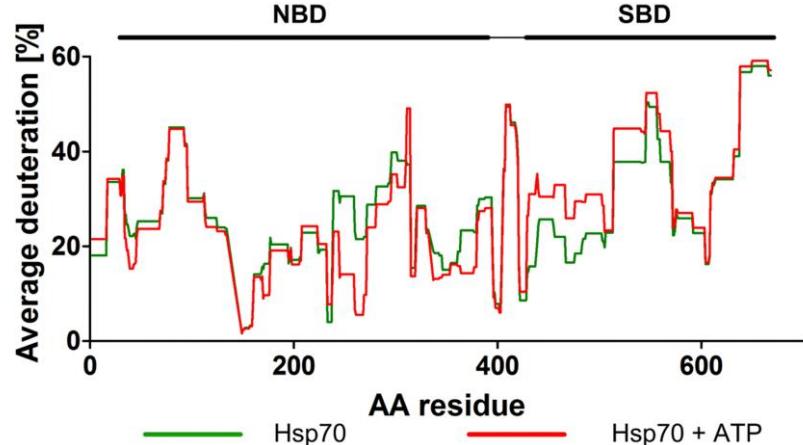
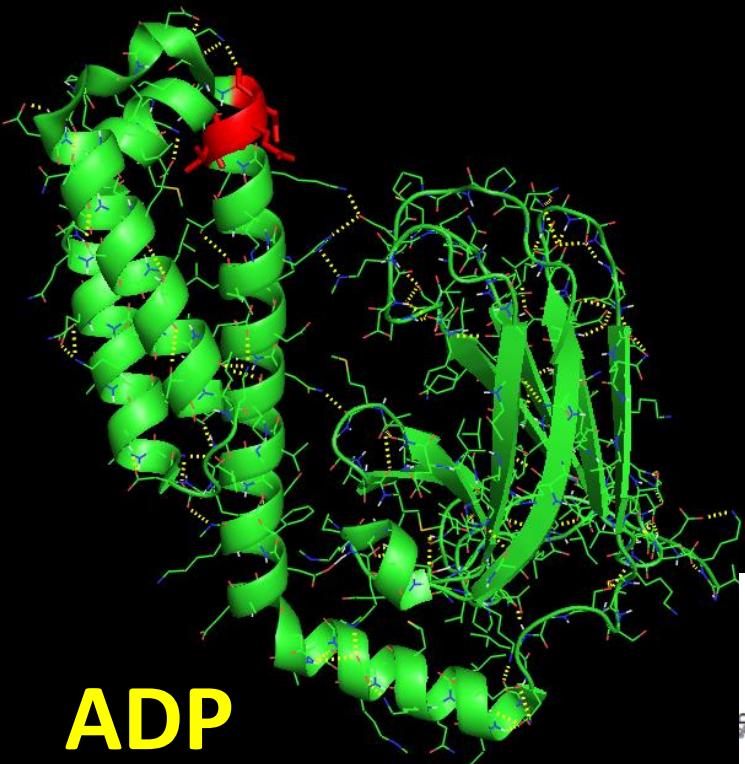
C



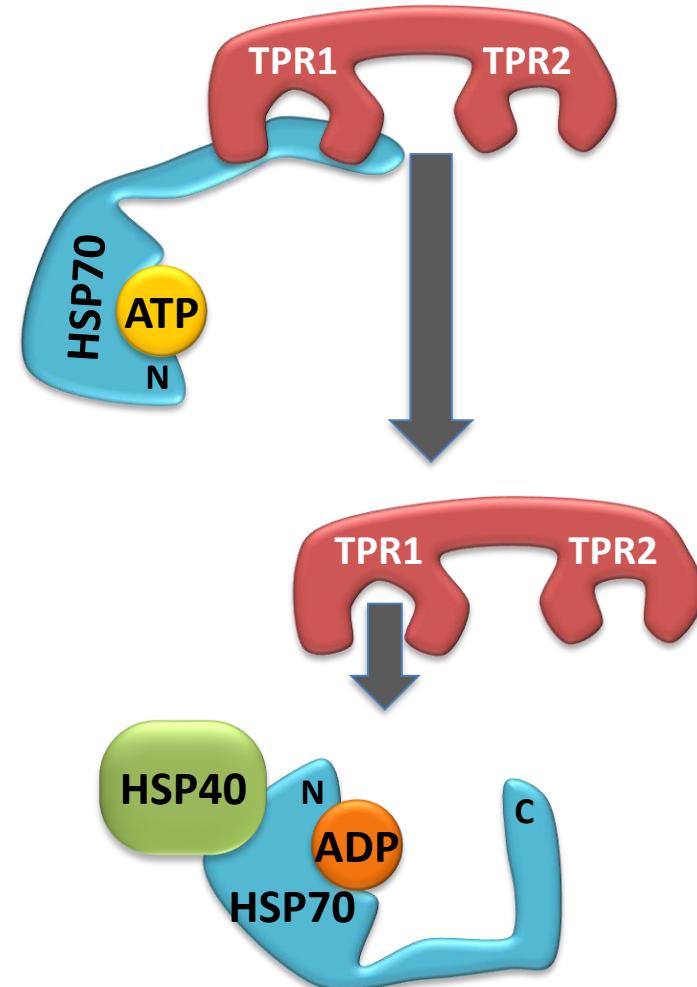
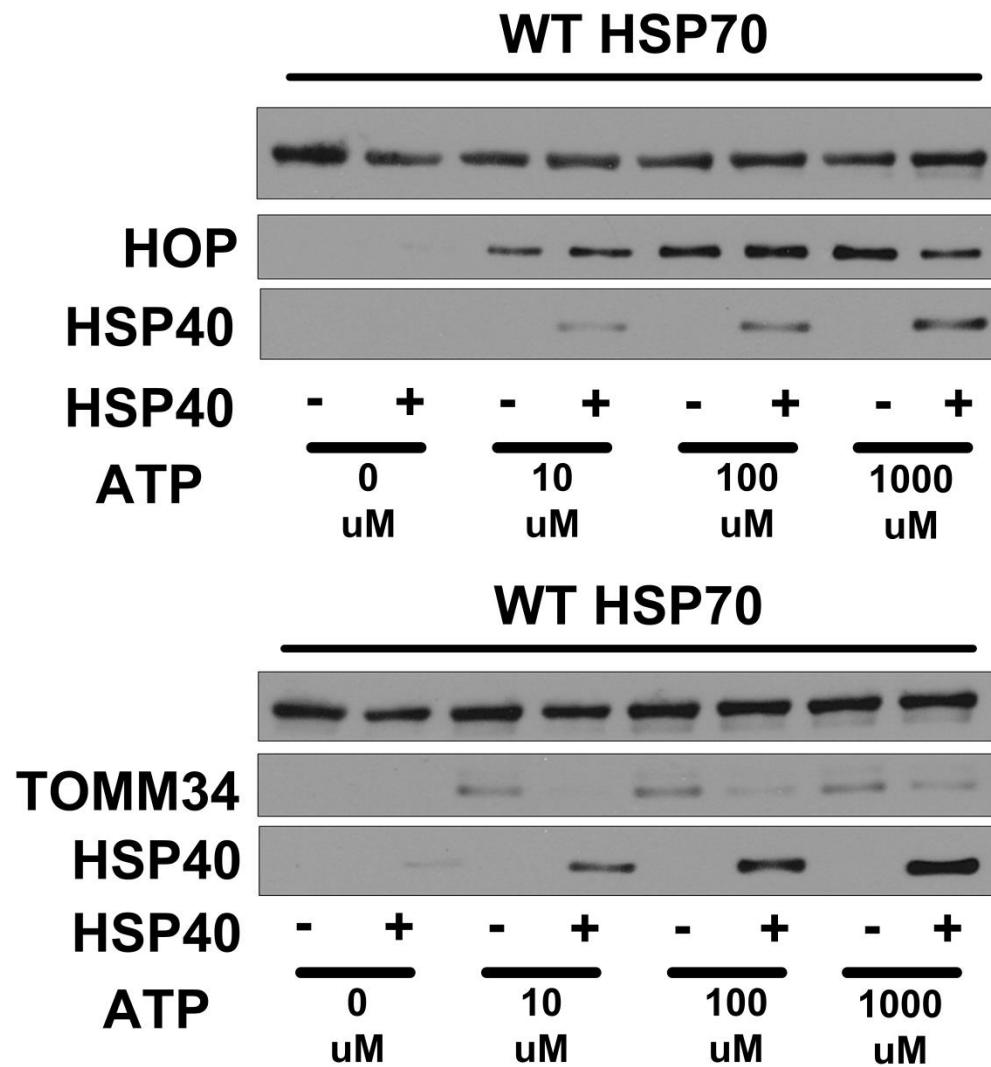
The effect of ATP on HSP70 – Tomm34 interaction



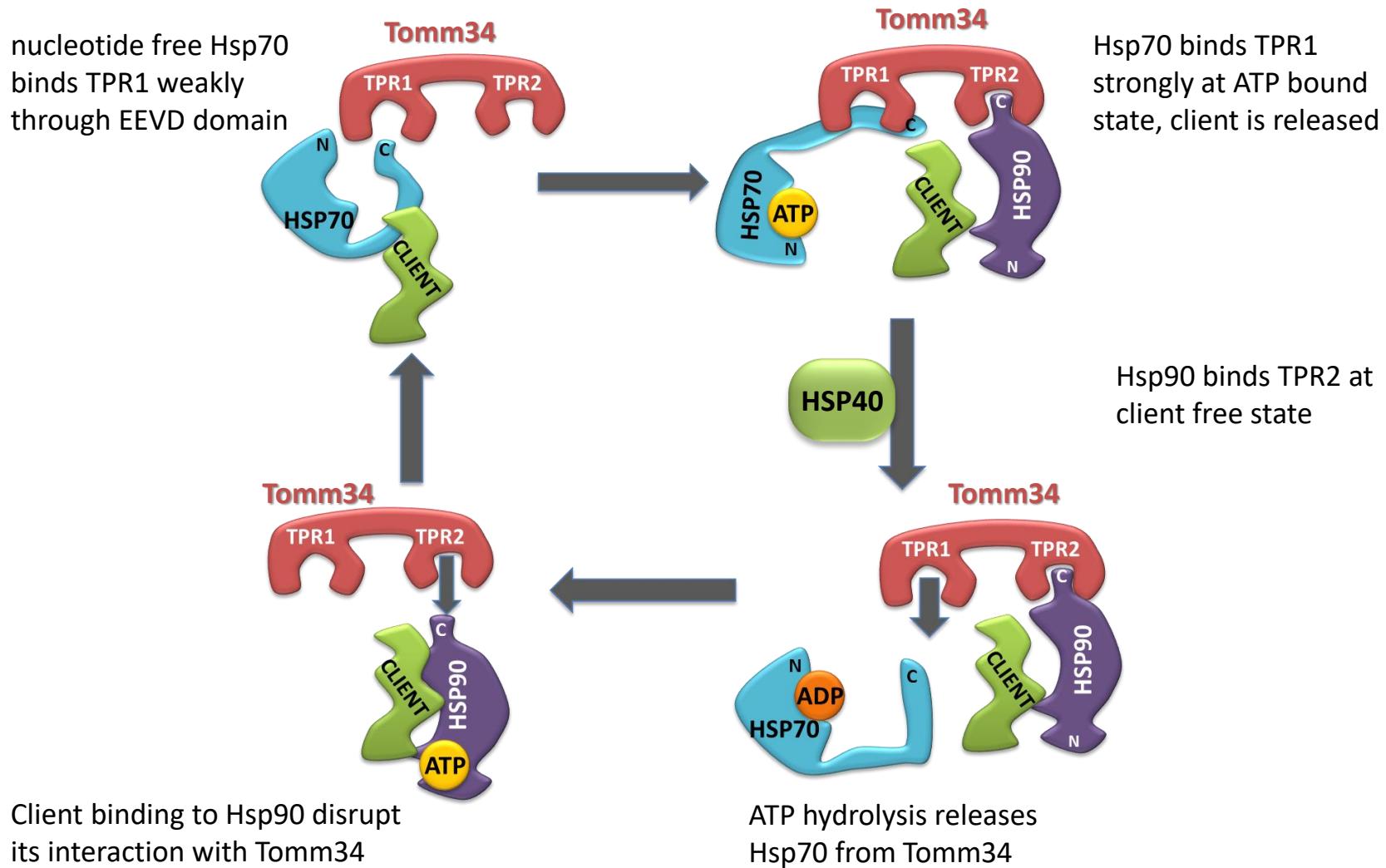
HDX – HSP70-TOMM34 interaction with ATP



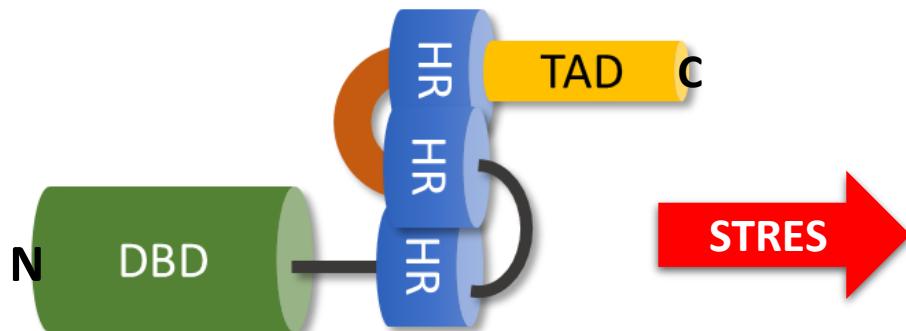
HSP70-TOMM34 interaction with ATP, the role of HSP40



Cooperation of Hsp70/Hsp90 folding by Tomm34



Regulace genové exprese chaperonů

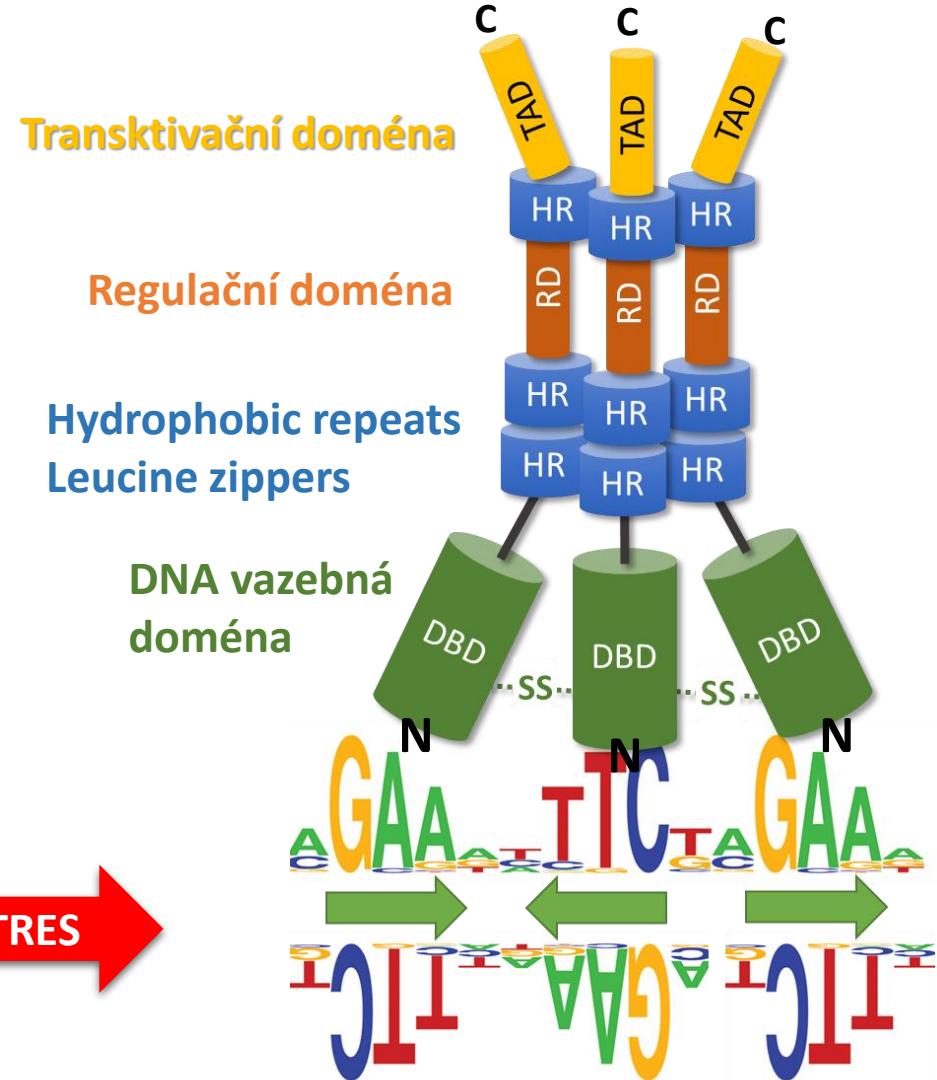


Transktivační doména

Regulační doména

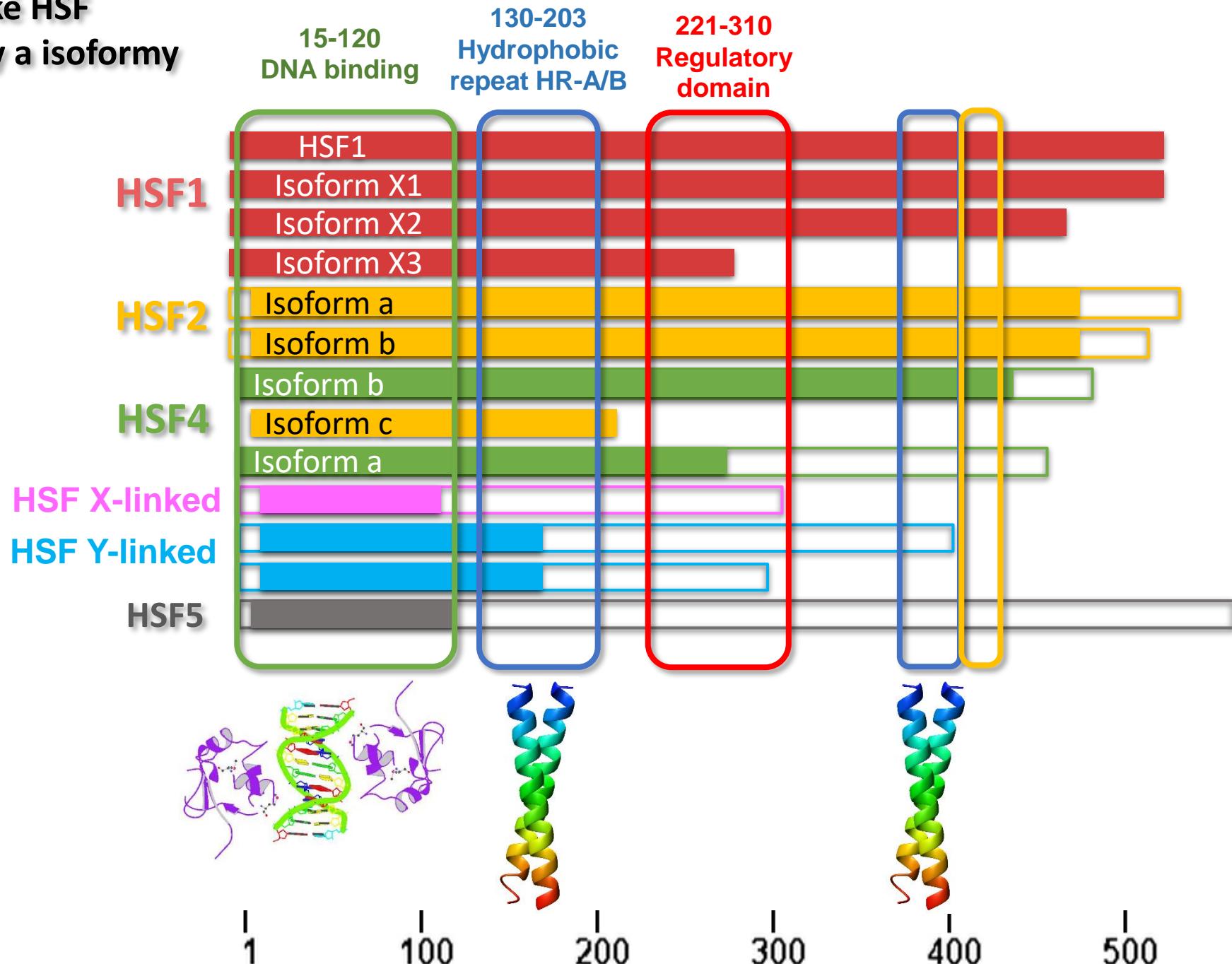
Hydrophobic repeats
Leucine zippers

DNA vazebná
doména



Lidské HSF

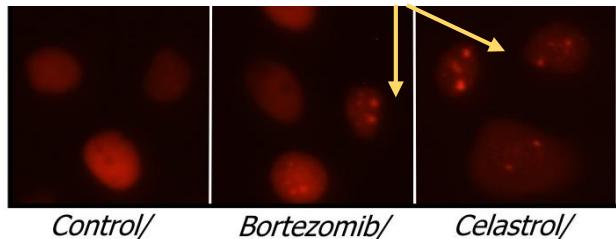
Geny a isoformy



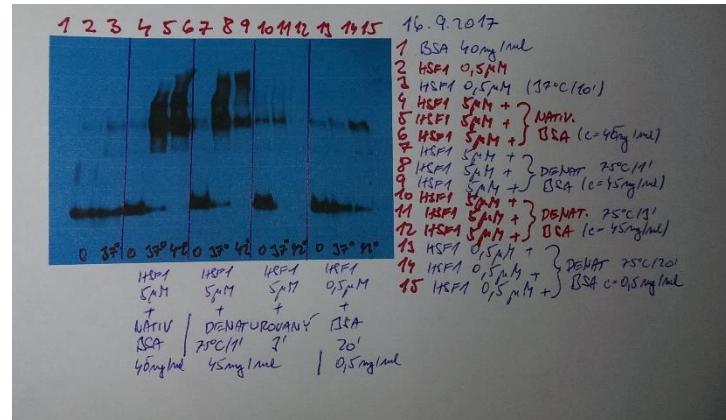
Analysis of HSF1 activation

HSF1-mCherry in A375 and H1299

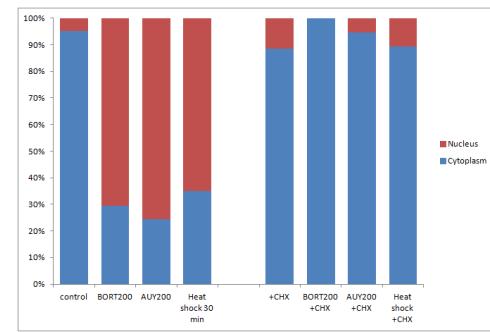
Nuclear stress bodies



Native gel, detection of trimers



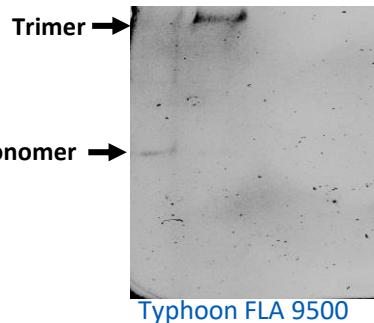
Cell fractionation



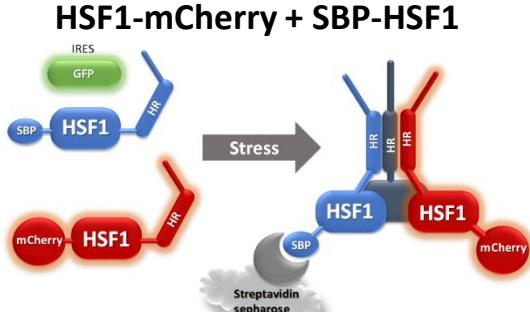
WB

- Crosslinkink
- Fractionation
- phosphorylation

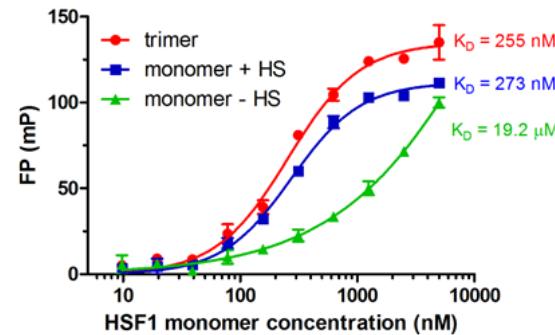
Native gel - mCherry



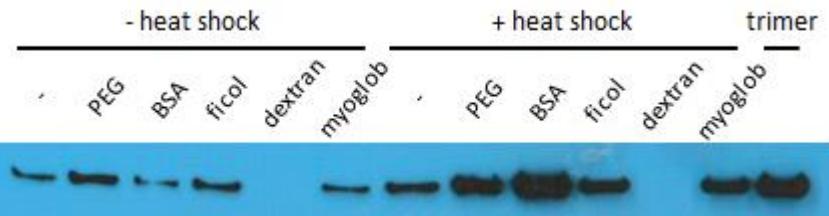
HSF1-mCherry + SBP-HSF1



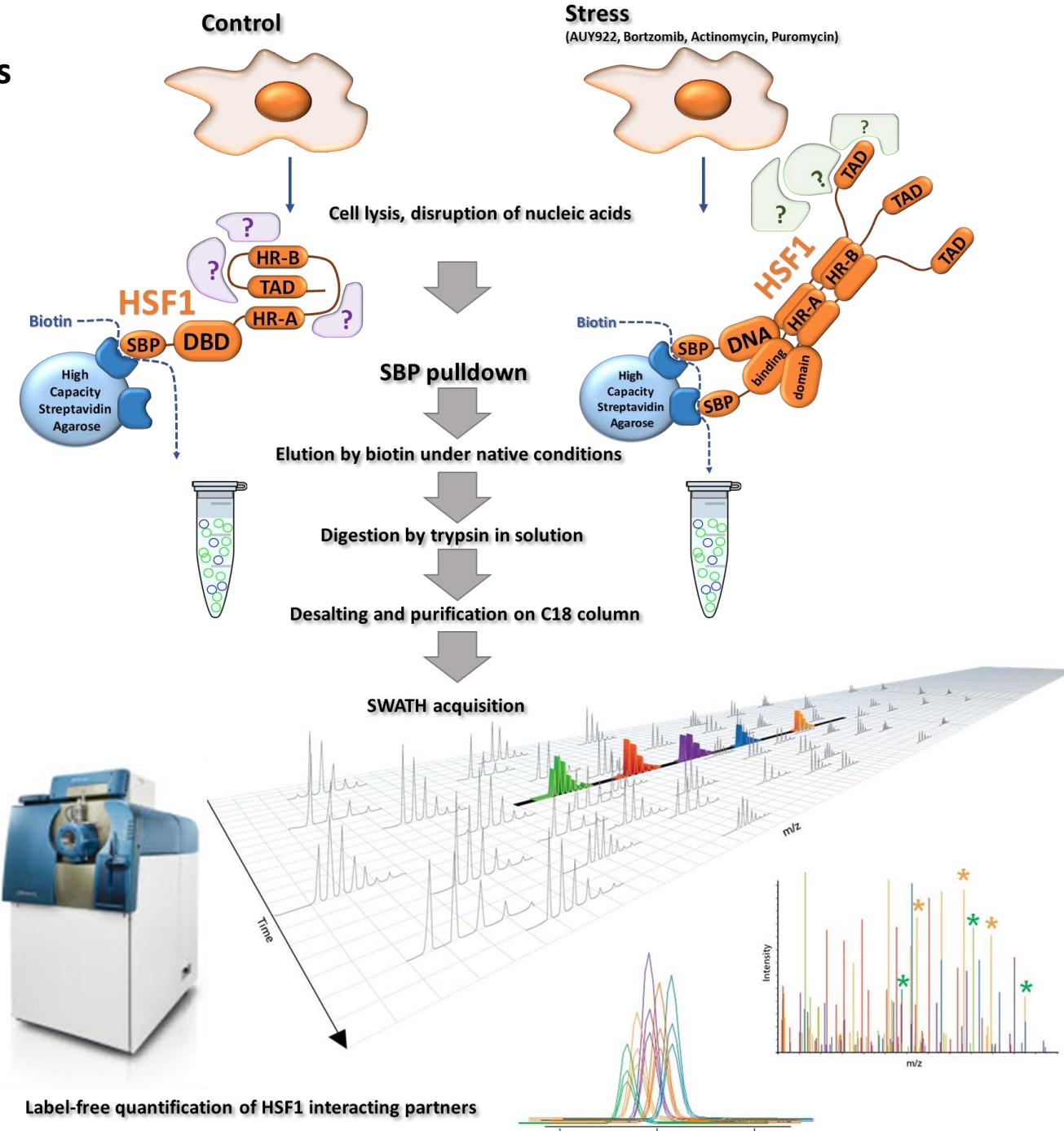
FAM-HSE binding to HSF1



Fluorescence polarization

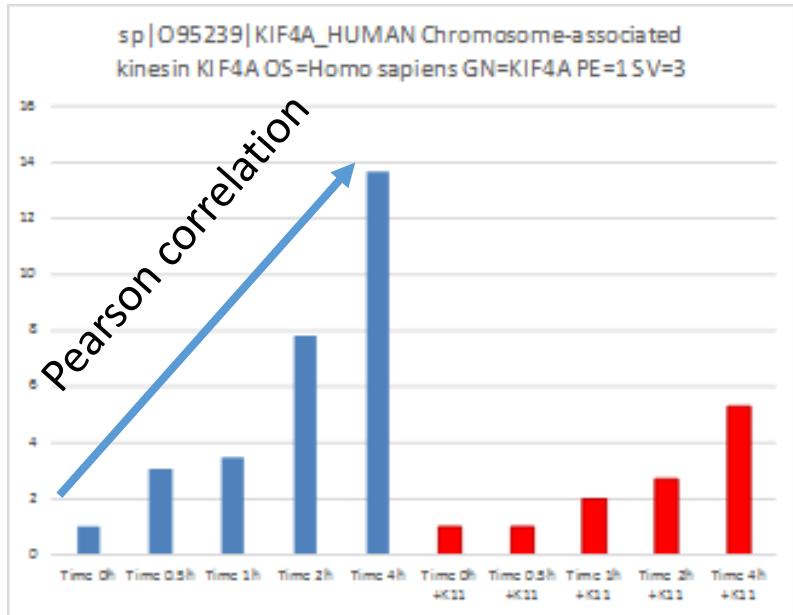


Analysis of HSF1 interaction partners Using functional proteomics



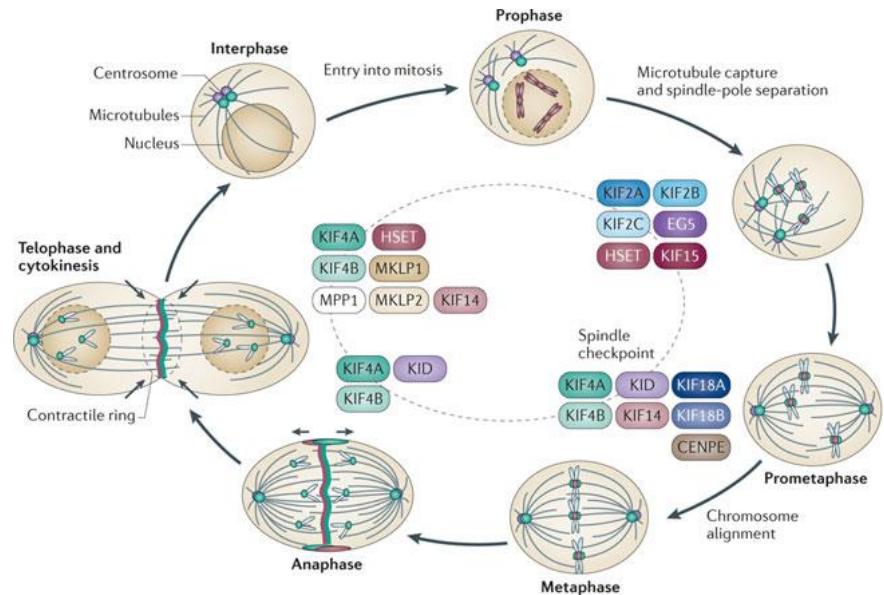
Time dependent interaction

KIF4A



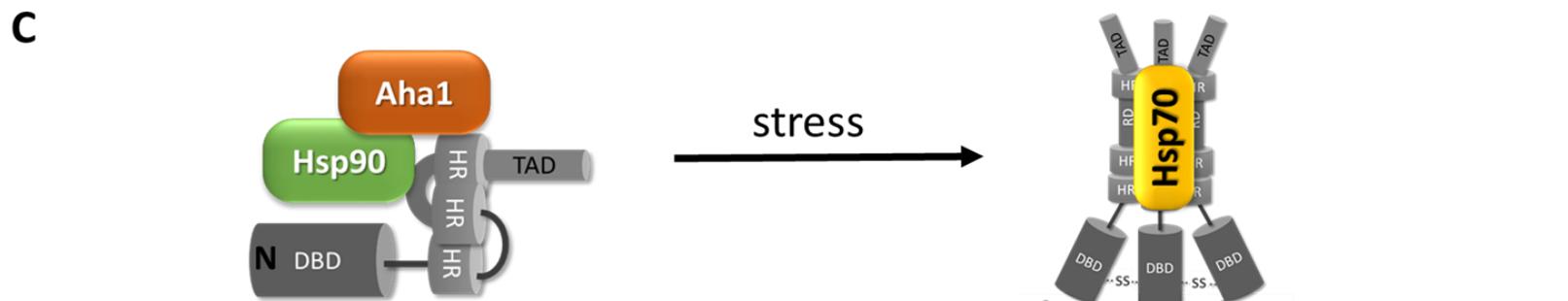
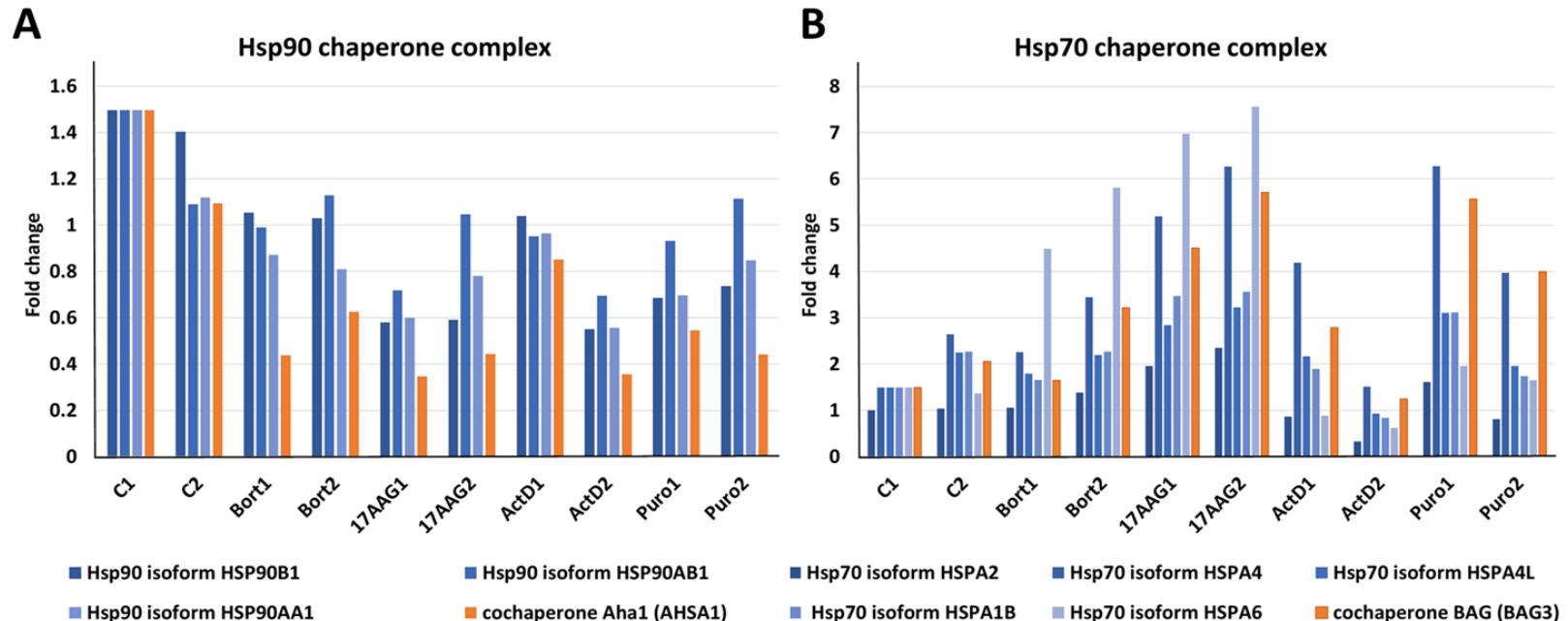
Bortezomib

Bortezomib
+ KRIBB11

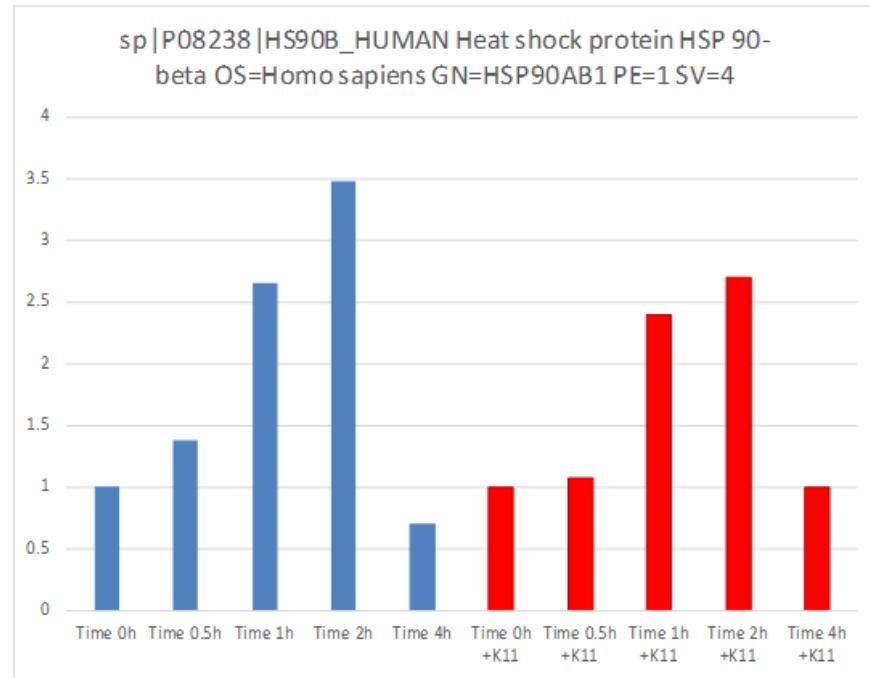
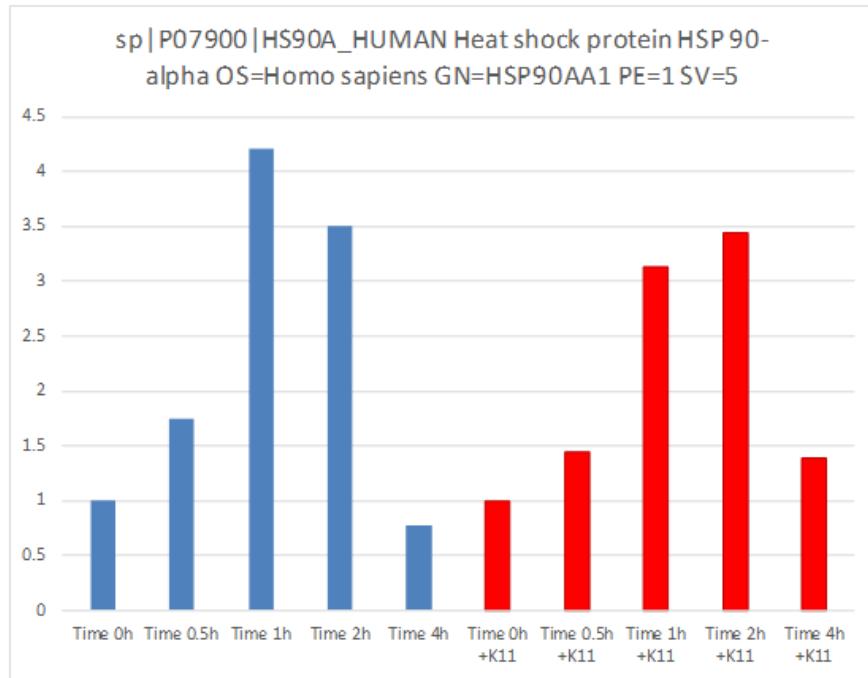


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Hsp90alpha/Hsp90beta

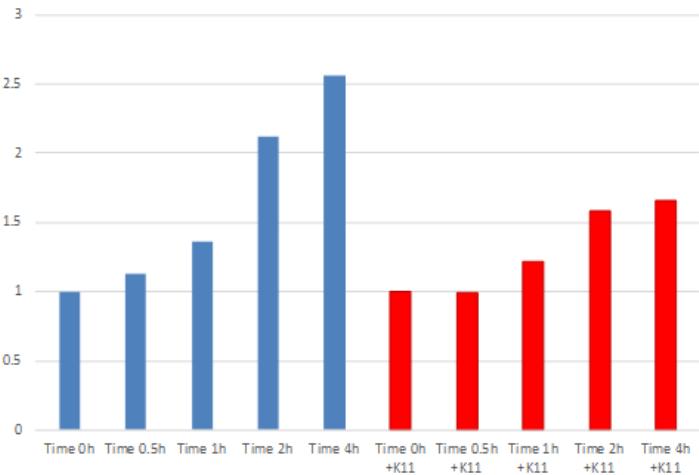


Interaction of HSF1 with Hsp90alpha/Hsp90beta after Bortezomib treatment



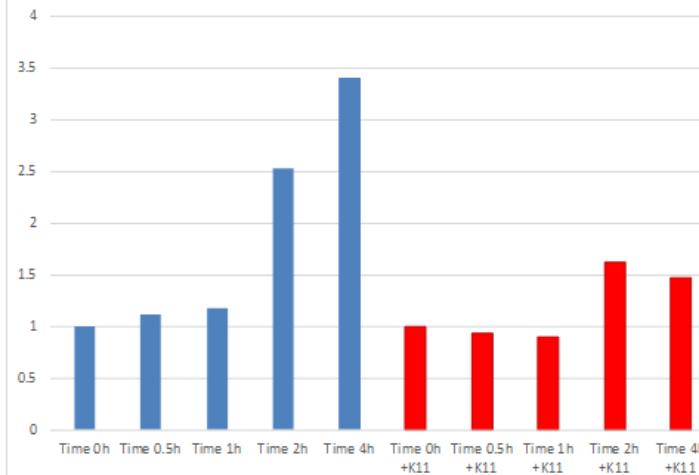
PRP8

sp|Q6P2Q9|PRP8_HUMAN Pre-mRNA-processing-splicing factor 8 OS=Homo sapiens GN=PRPF8 PE=1 SV=2



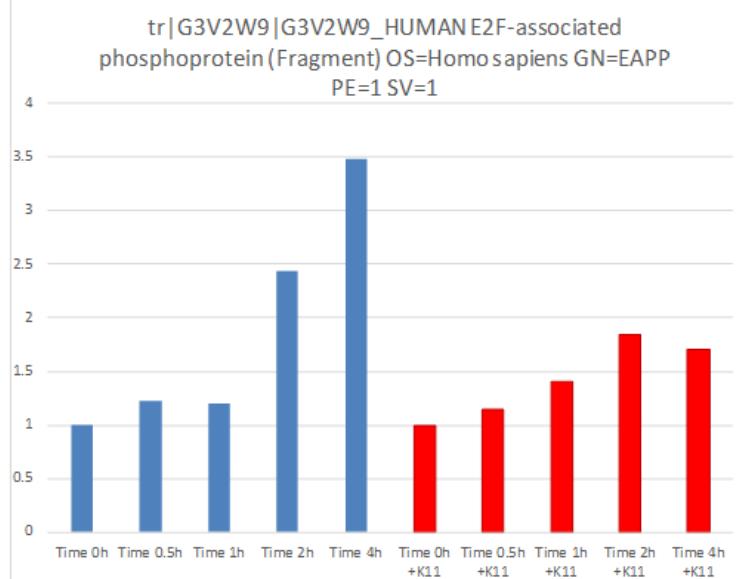
AAR2

sp|Q9Y312|AAR2_HUMAN Protein AAR2 homolog OS=Homo sapiens GN=AAR2 PE=1 SV=2



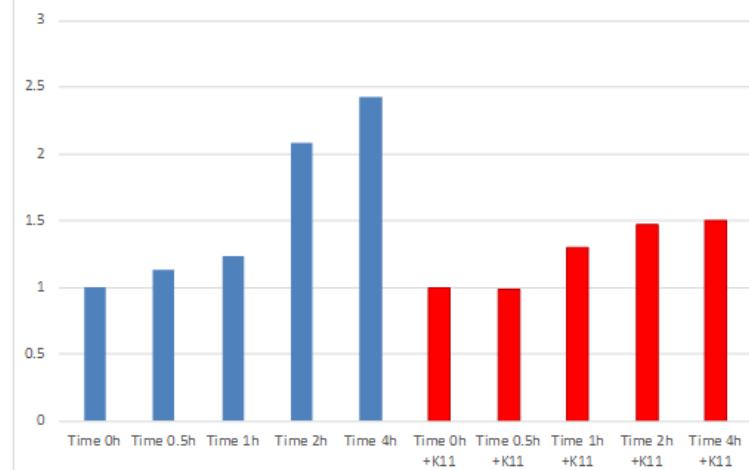
EAPP

tr|G3V2W9|G3V2W9_HUMAN E2F-associated phosphoprotein (Fragment) OS=Homo sapiens GN=EAPP PE=1 SV=1

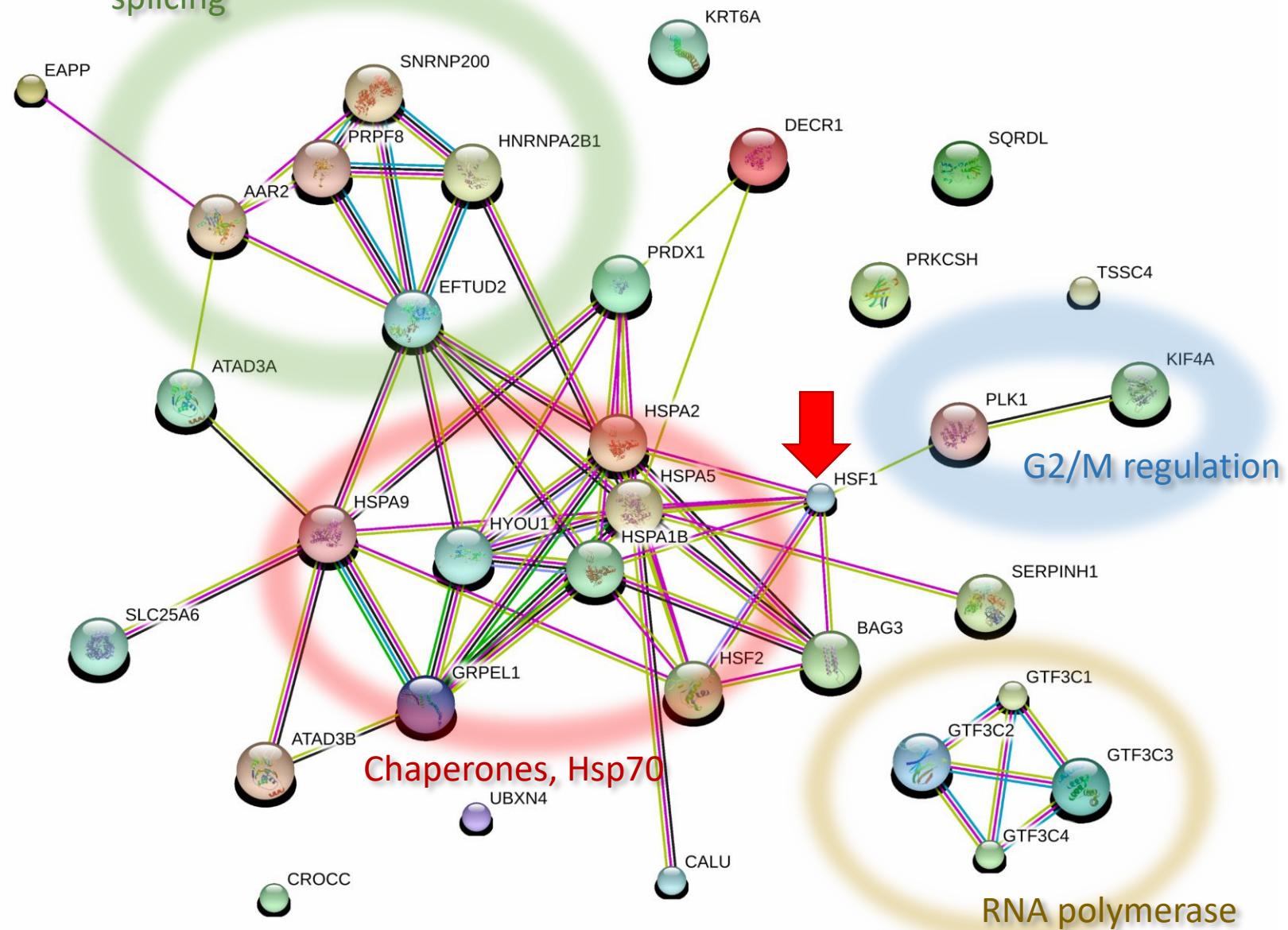


EFTUD2

sp|Q15029|U5S1_HUMAN 116 kDa U5 small nuclear ribonucleoprotein component OS=Homo sapiens GN=EFTUD2 PE=1 SV=1

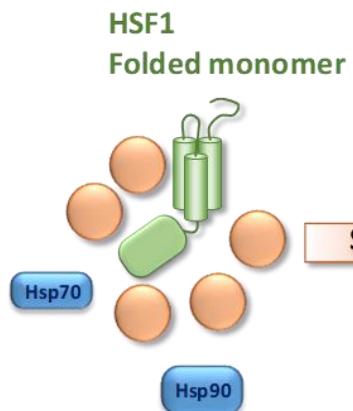


Pre mRNA processing splicing

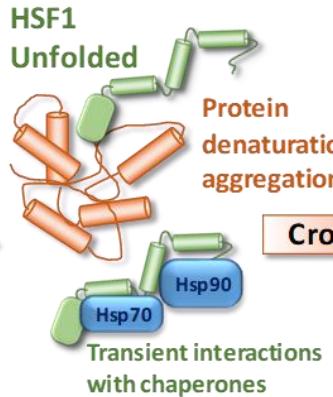


Mechanisms of HSF1 activation

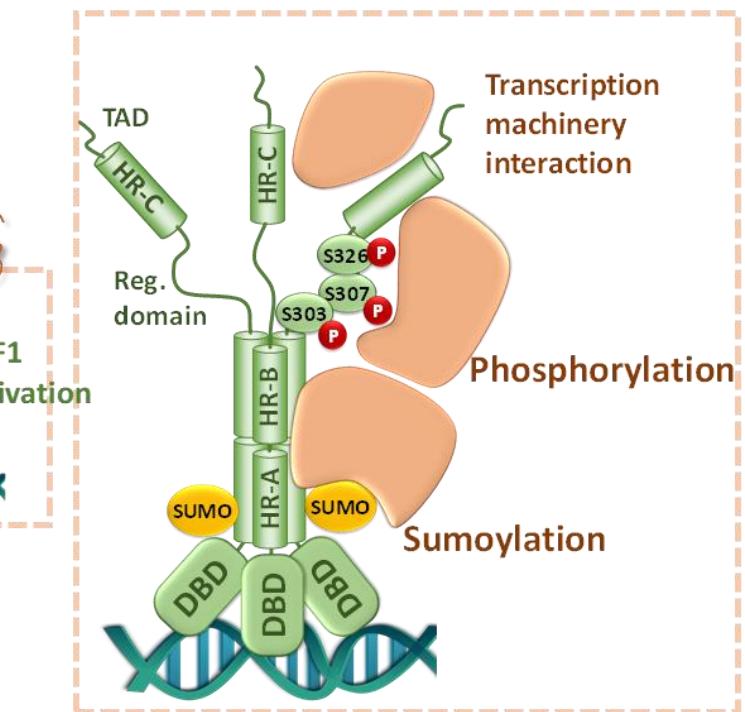
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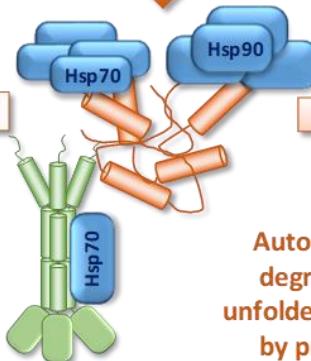
B



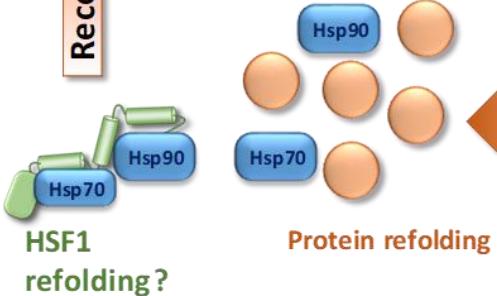
C



D



E



Autophagy and degradation of unfolded proteins by proteasome

This panel shows the pathway where unfolded proteins are targeted for degradation by the proteasome, leading to autophagy and protein degradation.

Děkuji za pozornost



- Bořivoj Vojtěšek
- Filip Trčka
- Eva Růčková
- Michal Ďurech
- Kateřina Křivánková



Edinburgh Cancer
Research Centre

Ted R. Hupp