



**ESMO HIGHLIGHTS**

19-23 DE OCTUBRE 2018, MUNICH

# ESTADIOS LOCALMENTE AVANZADOS

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Iniciativa científica de:



Grupo Español de Cáncer de Pulmón  
Spanish Lung Cancer Group

- Situación clínica compleja, que requiere de valoración en un comité de tumores multidisciplinar.
- Representa aproximadamente un tercio de los pacientes con cáncer de pulmón.
- Hay que distinguir entre pacientes operables y/o resecables.

## STAGE III – TREATMENT STRATEGIES

Stage III  
inoperable



Definitive CTx/RTx



Consolidation ITx

Stage III  
operable



Induction CTx



Surgery ( Postoperative RTx )



Induction CTx/RTx



Surgery



Surgery



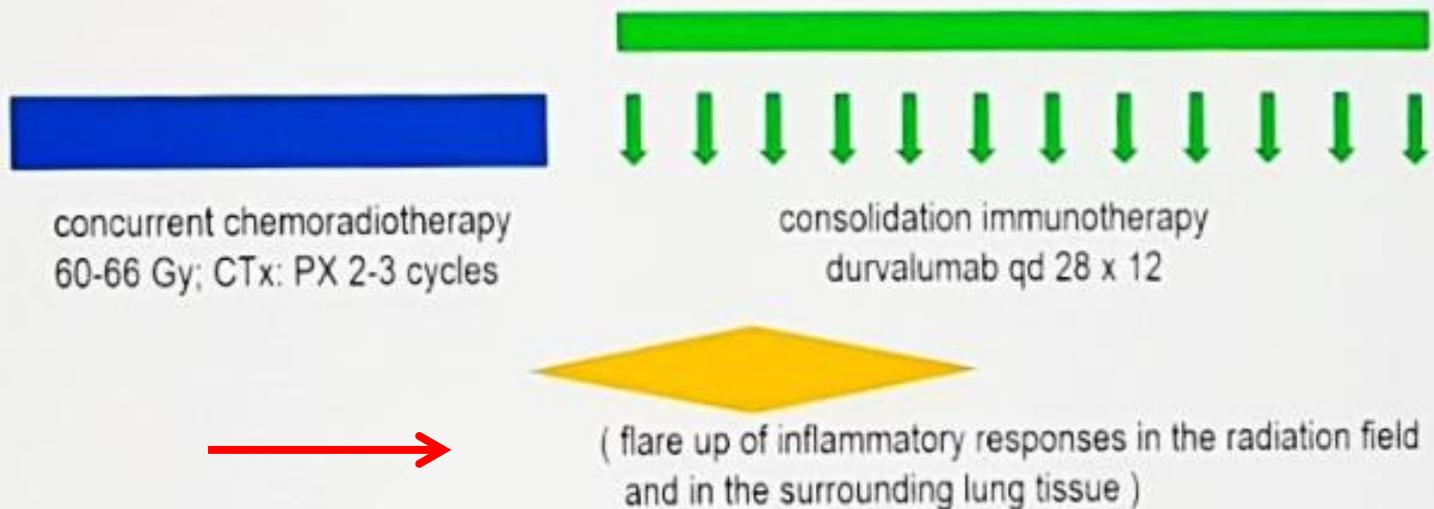
Adjuvant CTx



( Consolidation RTx )

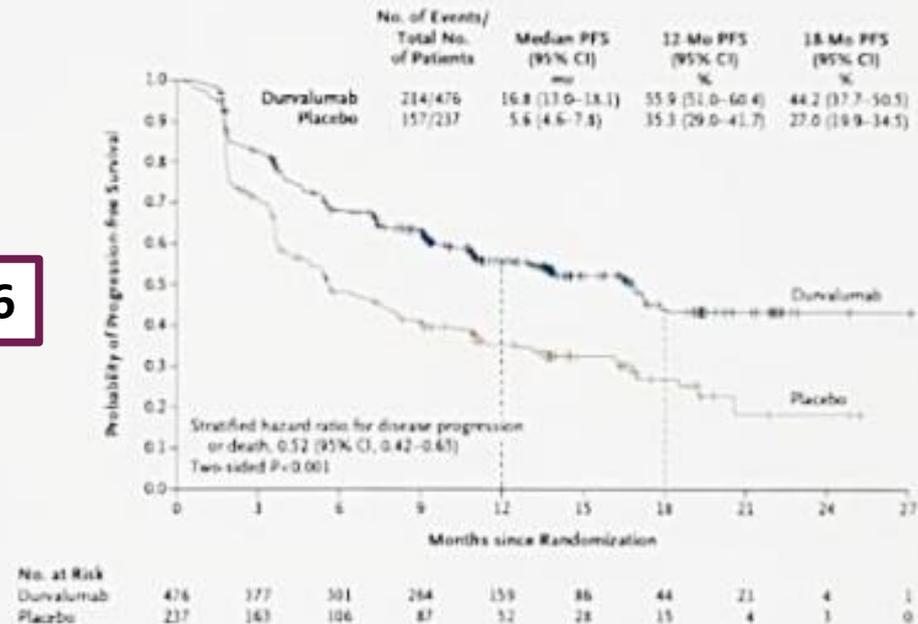
## IMMUNOTHERAPY IN STAGE III

- PACIFIC design



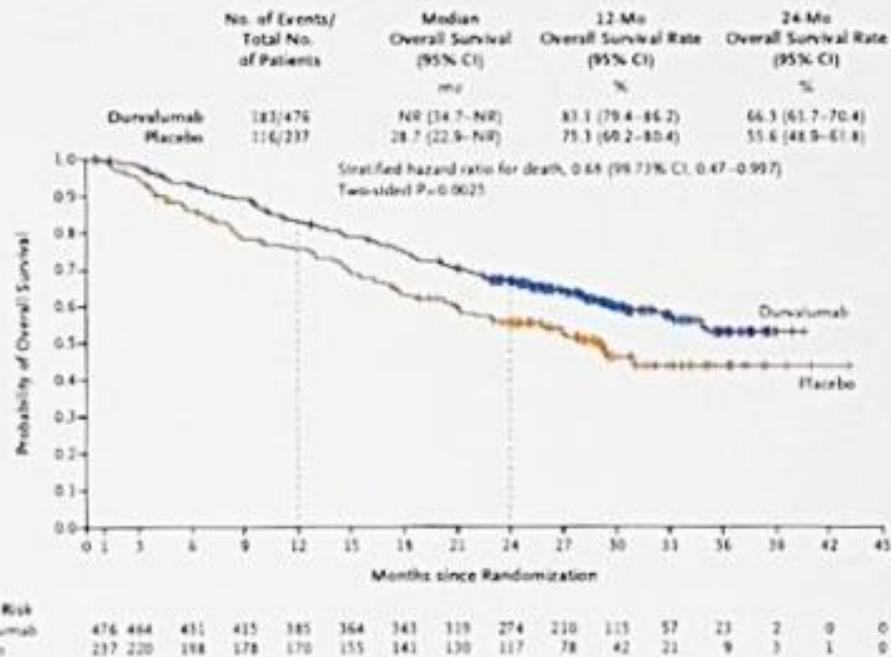
## PACIFIC – PROGRESSION-FREE SURVIVAL

**PFS= 16.8 VS 5.6**



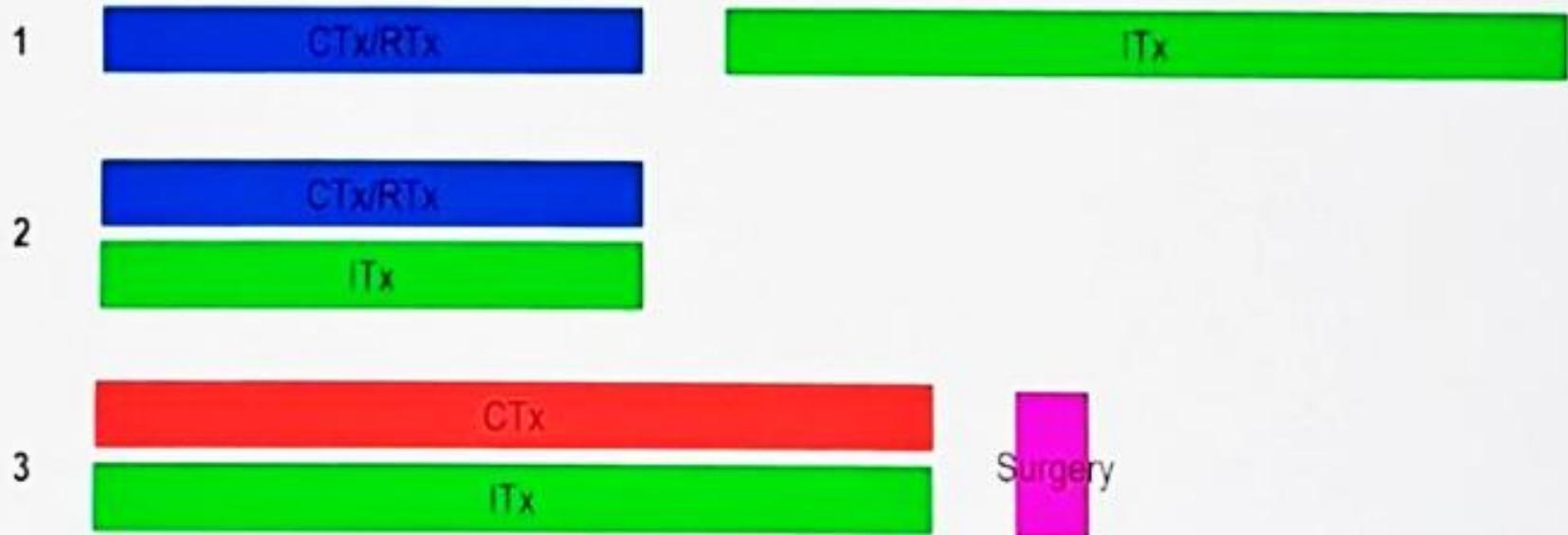
## PACIFIC - OVERALL SURVIVAL

OS= NR VS 28.7  
OS 2ª= 66.3% VS 55.6%  
HR=0.65  
PD-L1 >1%



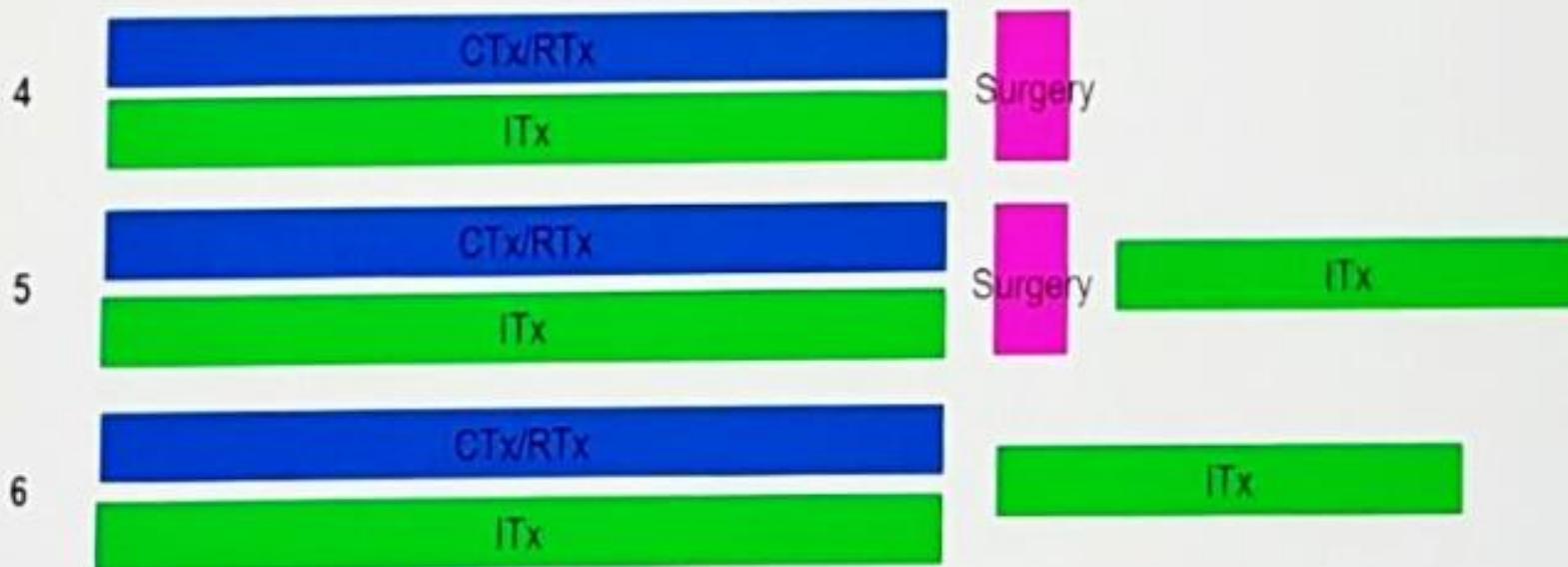
## IMMUNOTHERAPY IN STAGE III

- Possible permutations - 1



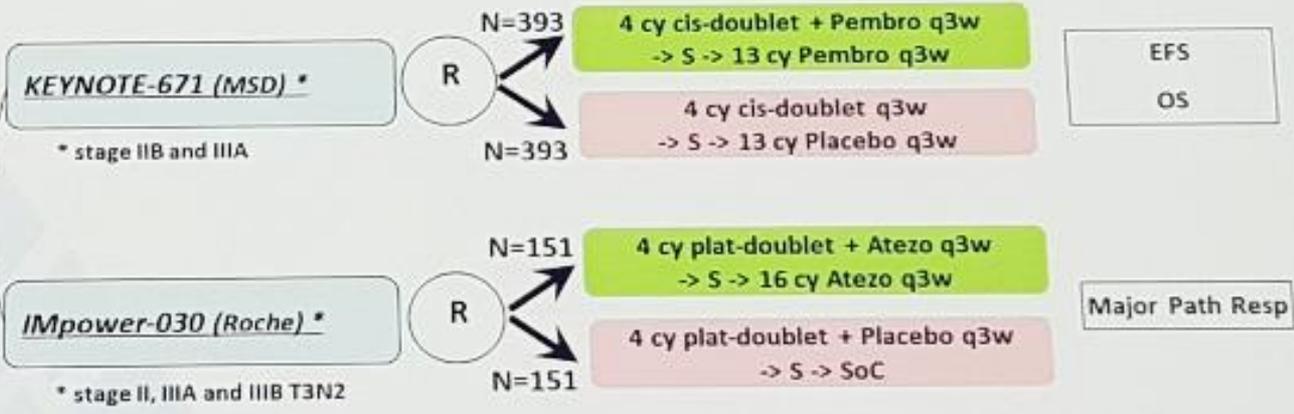
## IMMUNOTHERAPY IN STAGE III

- Possible permutations - 2



## Early stage NSCLC > neoadjuvant IO ph3 studies

- Selection of II-III
- PS 0-1
- Fit for surgery
- Any PDL1



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 Leuven Lung Cancer Group  
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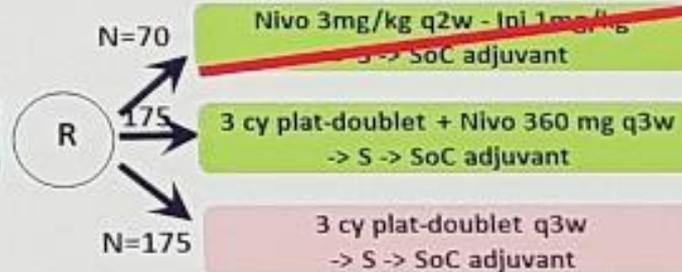
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## Early stage NSCLC

> neoadjuvant IO ph3 study

- Stage IB(>4cm)  
II and IIIA
- PS 0-1
- Fit for surgery
- EGFRwt ALK-
- Any PDL1

*Checkmate-816 (BMS)*



Path CR  
EFS



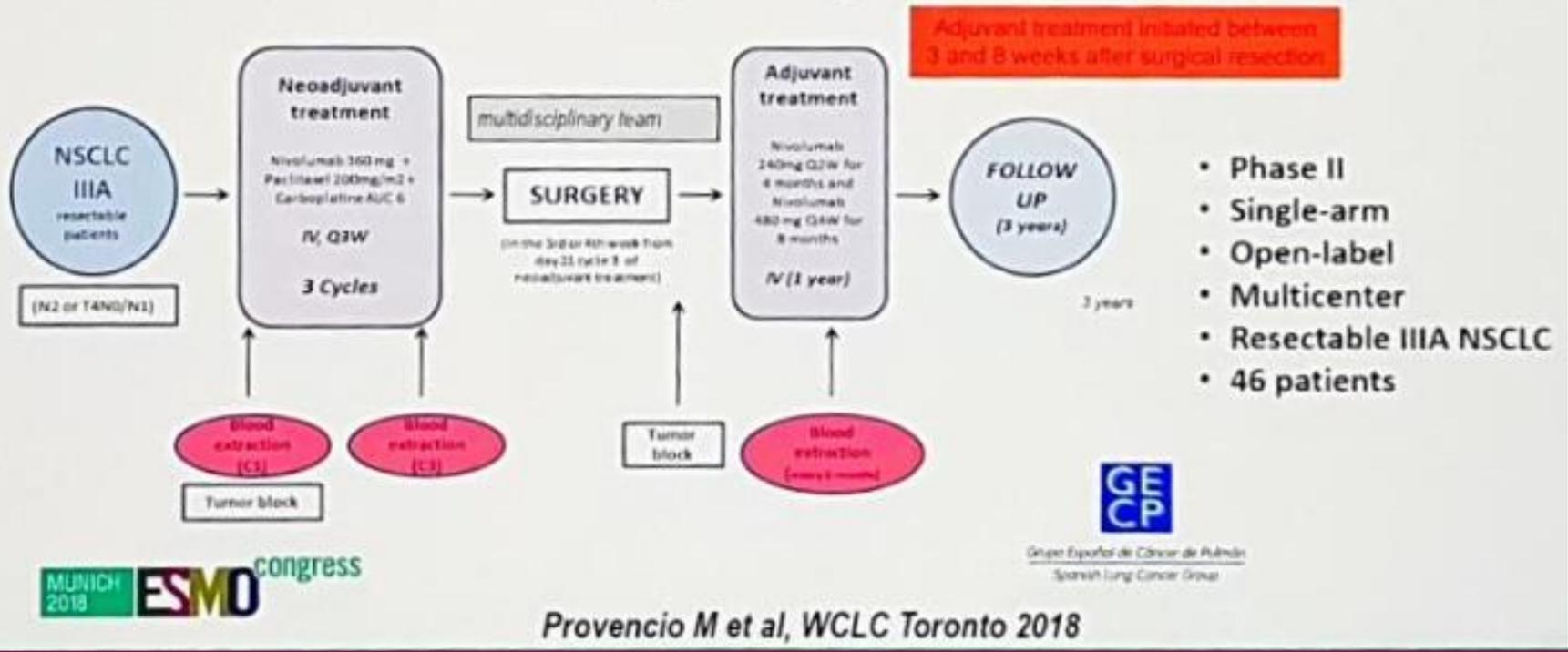
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## NADIM: Study design & Flow-chart



## Neoadjuvant treatment

	N	Median	Range
Cycles	45	3.0	(1.0-3.0)

CYCLES	N	%
1	3	5
3	43	95
<b>Total</b>	<b>46</b>	<b>100.0</b>

*All patients received three neoadjuvant cycles except for the three patients still being treated.*

## Clinical response

	N	%
Complete response (CR)	3	10.0
Partial response (PR)	18	60.0
Stable disease (SD)	9	30.0
<b>Total</b>	<b>30</b>	<b>100.0</b>

*No progressive disease has been observed.*



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## Ongoing Targeted Clinical Trials for NSCLC in Neoadjuvant setting

ClinicalTrials.gov ID	Phase	Drug	Setting	Endpoint	No. of Pts
NCT02347839	2	Gefitinib → surgery → gefitinib	Neoadjuvant, Stage III NSCLC, EGFRm	Resectability rate	37
NCT02804776	2	Gefitinib	Induction (4 weeks) for resectable NSCLC, EGFRm	Biomarker evaluation	40
<b>NCT03433469</b>	<b>2</b>	<b>Osimertinib</b>	<b>Neoadjuvant, Stage I–IIIA, resectable NSCLC, EGFRm</b>	<b>MPR</b>	<b>27</b>
NCT01407822 (EMERGING)	2, 3	Erlotinib versus gemcitabine / cisplatin	Neoadjuvant, Stage IIIA NSCLC, EGFRm	ORR	70
NCT03349203	2	Icotinib	Adjuvant / neoadjuvant, Stage IIIB NSCLC, EGFRm	ORR	60
NCT02820116	2	Icotinib	Neoadjuvant, Stage III NSCLC, EGFRm	Complete resection rate	67
NCT03088930	2	Crizotinib	Neoadjuvant, Stage I–IIIA, resectable NSCLC, with an activating alteration in ALK, ROS1, or MET	ORR	18
UMIN000017906 (SAKULA)	2	Ceritinib	Neoadjuvant, Stage II–IIIA, resectable NSCLC, with an activating alteration in ALK	MPR	7/19

