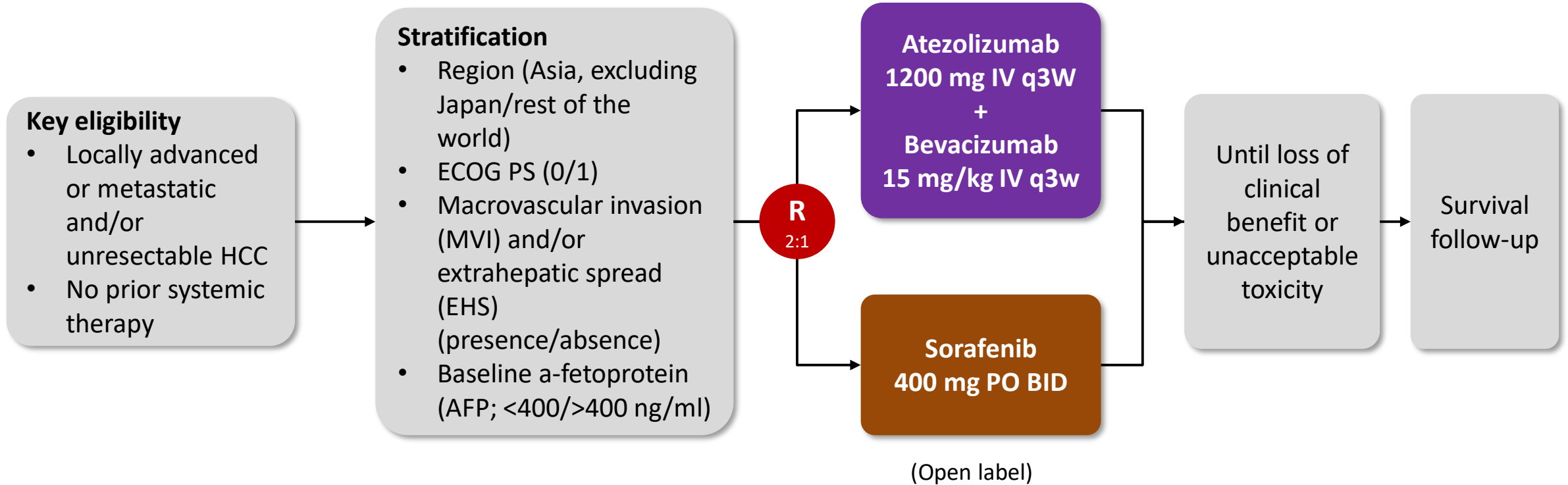




CHC avancé : le traitement de référence

Schéma de l'étude de Phase III IMbrave 150



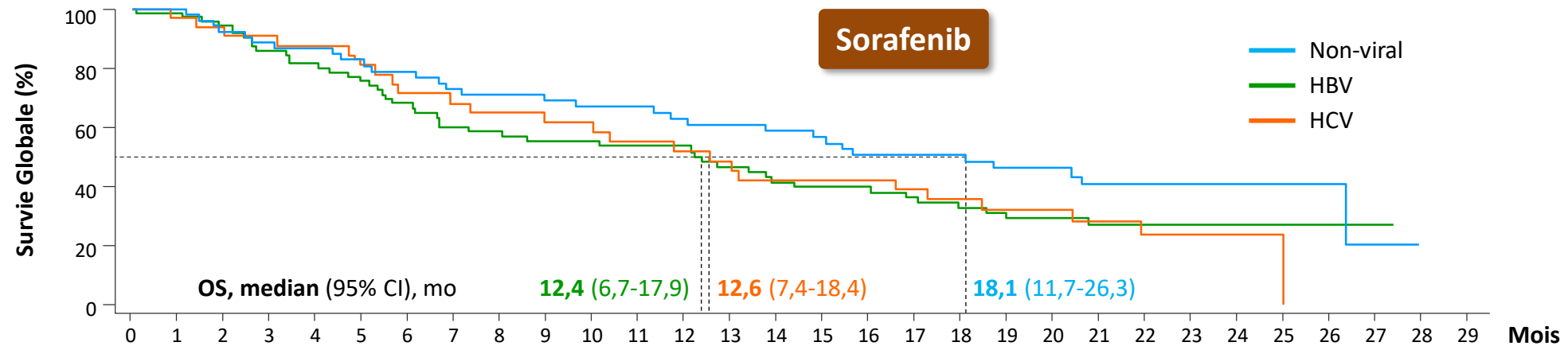
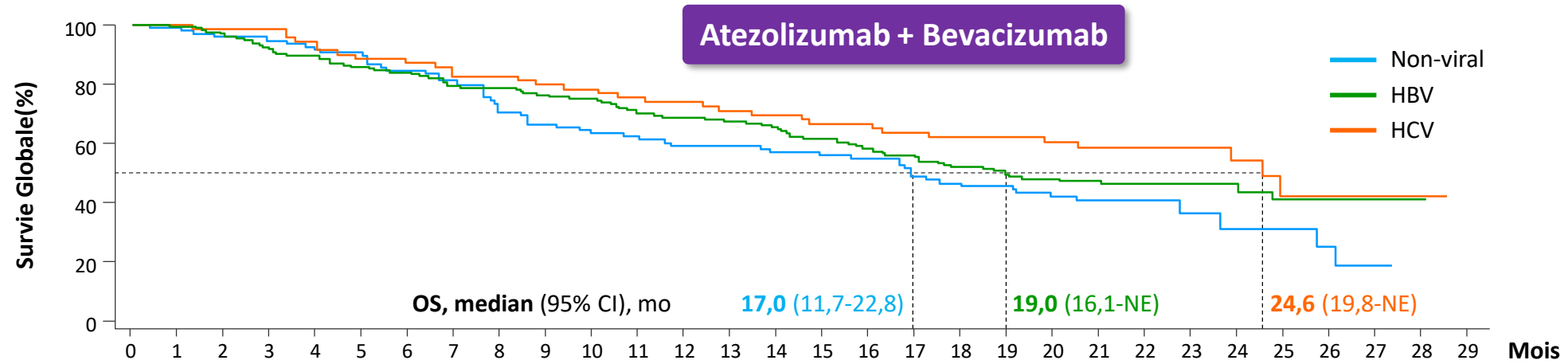
Co-primary endpoints

- OS
- IRF-assessed PFS per RECIST 1.1

Key secondary efficacy endpoints

- IRF-assessed PFS per HCC mRECIST
- IRF-assessed ORR per RECIST 1.1 and HCC mRECIST
- IRF-assessed DOR per RECIST 1.1 and HCC mRECIST

Courbes de survie globale



Effets indésirables grade 3/4 survenant chez $\geq 5\%$, n (%)

| | Non-viral population | | HBV population | | HCV population | |
|--------------------------------|-----------------------|---------------------|------------------------|---------------------|-----------------------|---------------------|
| | Atezo + Bev (n=98) | Sorafenib (n=52) | Atezo + Bev (n=162) | Sorafenib (n=71) | Atezo + Bev (n=69) | Sorafenib (n=33) |
| Hypertension | 18 (18) | 5 (10) | 24 (15) | 9 (13) | 14 (20) | 5 (15) |
| Augmentation Bilirubine | 7 (7) | 3 (6) | 5 (3) | 3 (4) | 1 (1) | 4 (12) |
| Diarrhée | 5 (5) | 5 (10) | 4 (2) | 3 (4) | 2 (3) | 0 |
| Ascite | 5 (5) | 3 (6) | 6 (4) | 0 | 1 (1) | 0 |
| Augmentation ASAT | 4 (4) | 5 (10) | 11 (7) | 3 (4) | 11 (16) | 1 (3) |
| Syndrome main-pied | 0 | 2 (4) | 0 | 5 (7) | 0 | 6 (18) |