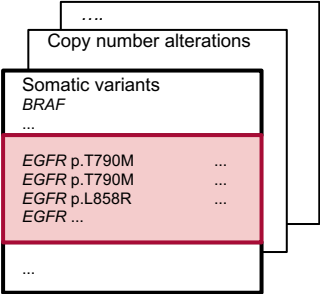


Matching a clinically relevant somatic variant to catalogued assertions

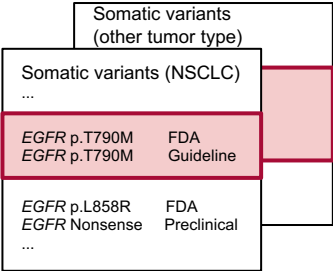
a NSCLC profile harboring
somatic variant **EGFR** p.T790M

Match to
Molecular Oncology Almanac
by data type and gene



b **NSCLC** profile harboring
somatic variant *EGFR* **p.T790M**

Match by ontology and additional
feature details



c NSCLC profile harboring
somatic variant *EGFR* p.T790M

Select strongest evidence,
additional items are returned as
equivalent matches

