

S3 Table. Feasibility of comprehensive molecular profiling in tumour tissue. The table shows the absolute numbers and percentage of successful comprehensive molecular profiling. Also, the frequency of missing requests for profiling (due to unspecified reasons) is shown, and reasons for profiling being unfeasible. The data correspond to the graphs in figure 3. Numbers indicate n(%).

Any histology	Stage I-III		Stage IV	
	Run-in	Protocol	Run-in	Protocol
Total number of patients	60	414	76	328
Molecular profiling status unknown	0	2 (0.5%)	0	1 (0.3%)
Completed molecular profiling	31 (51.7%)	249 (60.4%)	52 (68.4%)	259 (79.2%)
Incomplete molecular profiling	29 (48.3%)	163 (39.6%)	24 (31.6%)	68 (20.8%)
NGS and/or ALK test not requested	23 (38.3%)	70 (17.0%)	10 (13.2%)	11 (3.4%)
ROS1 and/or RET test not requested	0	28 (6.8%)	7 (9.2%)	17 (5.2%)
<i>EGFR</i> and <i>ALK</i> testing not feasible	6 (10.0%)	62 (15.0%)	5 (6.6%)	35 (10.7%)
Insufficient tumour tissue	5 (8.3%)	40 (9.7%)	5 (6.6%)	25 (7.6%)
No tumour tissue obtained	1 (1.7%)	11 (2.7%)	0	6 (1.8%)
No tumour tissue available at LEMA centre	0	10 (2.4%)	0	1 (0.3%)
Patient died before molecular analysis	0	1 (0.2%)	0	3 (0.9%)
Biomarker (ROS1/RET) analysis failed ^a	0	3 (0.7%)	2 (2.6%)	5 (1.5%)

Non-squamous NSCLC	Stage I-III		Stage IV	
	Run-in	Protocol	Run-in	Protocol
Total number of patients	37	295	67	286
Molecular profiling status unknown	0	1 (0.3%)	0	1 (0.3%)
Completed molecular profiling	20 (54.1%)	202 (68.7%)	51 (76.1%)	241 (84.6%)
Incomplete molecular profiling	17 (45.9%)	92 (31.3%)	16 (23.9%)	44 (15.4%)
NGS and/or ALK test not requested	11 (29.7%)	40 (13.6%)	4 (6.0%)	0 (0.4%)
ROS1 and/or RET test not requested	0	9 (3.1%)	6 (9.0%)	11 (3.9%)
<i>EGFR</i> and <i>ALK</i> testing not feasible	6 (16.2%)	40 (13.6%)	5 (7.5%)	28 (9.8%)
Insufficient tumour tissue	5 (13.5%)	28 (9.5%)	5 (7.5%)	19 (6.7%)
No tumour tissue obtained	1 (2.7%)	8 (2.7%)	0	6 (2.1%)
No tumour tissue available at LEMA centre	0	4 (1.4%)	0	0
Patient died before molecular analysis	0	0	0	3 (1.1%)
Biomarker (ROS1/RET) analysis failed ^a	0	3 (1.0%)	1 (1.5%)	4 (1.4%)

Squamous NSCLC	Stage I-III		Stage IV	
	Run-in	Protocol	Run-in	Protocol
Total number of patients	23	119	9	42
Molecular profiling status unknown	0	1 (0.8%)	0	0
Completed molecular profiling	11 (47.8%)	47 (39.8%)	1 (11.1%)	18 (42.9%)
Incomplete molecular profiling	12 (52.2%)	71 (60.2%)	8 (88.9%)	24 (57.1%)
NGS and/or ALK test not requested	12 (52.2%)	30 (25.4%)	6 (66.7%)	10 (23.8%)
ROS1 and/or RET test not requested	0	19 (16.1%)	1 (11.1%)	6 (14.3%)
<i>EGFR</i> and <i>ALK</i> testing not feasible	0	22 (18.6%)	0	7 (16.7%)
Insufficient tumour tissue	0	12 (10.2%)	0	6 (14.3%)
No tumour tissue obtained	0	3 (2.5%)	0	0
No tumour tissue available at LEMA centre	0	6 (5.1%)	0	1 (2.4%)
Patient died before molecular analysis	0	1 (0.8%)	0	0
Biomarker (ROS1/RET) analysis failed ^a	0	0	1 (11.1%)	1 (2.4%)

^a in some patients, ROS1 and/or RET tested using FISH failed due to technical reasons