

SUPPLEMENTAL FIGURE 1. Flow chart defining tissue positivity and negativity in the ⁶⁸Ga-DOTATOC retrospective trial.

SUPPLEMENTAL TABLE 1
Standardized Data Collection Criteria for non-Proprietary PET Radiopharmaceuticals

Standardized Criteria	Reason	Examples
Radiopharmaceutical end-product specification	Assure same drug quality across trials	pH range, radionuclidic purity, radiopharmaceutical purity
Adverse Event Data Collection	Assure safety standards consistently reported	Same reporting lexicon, same follow-up periods
Dose Range Acquisition Parameters	Assuring same dose Assuring same/similar imaging protocols	3-5 mCi, with target 4 mCi Scan commenced 55-70 minutes post injection
Interpretation criteria	Assure consistent interpretation across trials	Dual blinded reads, Threshold SUV _{max} , Minimum size criteria
Reference/Truth Standard	Assure true positive, true negative, false positive, and false negative results are reported based on same criteria.	Comparison with histopathology or if no pathology, then conventional imaging.