virus_beacon_schema_v1_to_generic

	–VARIANT BASIC (basic beacon variant schema: from VCF)
	taxon_id (taxon of sequenced species)
	ref_assembly_id_version:
	start_nucleotide
	end_nucleotide
	ref_sequence
	alt_sequence
	variantType: SNV, indel, CNV, structural variant
	-VARIANT ANNOTATION
	variant_id: (external reference if it exists)
\checkmark	variant_type: e.g "del" (belongs in beacon variant basic)
	variant effect: e.g "missense variant" → molecularConsequence
	genomic_region: categorical, from virus genomic annotation in VIRUS: annotation (SARS-
	CoV2: 5UTR,ORF1ab, S, ORF3a, Intergenic, E,M, ORF6, ORF7a, ORF8, N, ORF10, 3UTR) →
	genomicRegionClass and featureId
\checkmark	functional_region: categorical, from functional annotation file VIRUS: annotation e.g
	"HVR", "RBD", "RNA modification site" or manual entering by user.
	-VARIANT IN SAMPLE
	biosample_id: (external ref) e.g "SRS6007144" → Variant in Sample biosampleId
	host_id: (external ref if it exists) → Variant in Sample individualId
	variant_frequency_dataset: from vcf → Variant in Sample variantFrequency
	info
	✓ study_info:
	✓ study_id: (study accession): e.g "SRP242226"
	✓ study_ref: (article PUMED ID or URL)
	experiment_info
	√ variant_file_id: (accession if external ref or internal if we run pipeline) > (to info?)
	✓ sequence_file_id: (run accession) e.g "SRR10903401"
	✓ exp_id (experiment accession): e.g "SRX7571571"
	exp. title: e.g. "Total RNA sequencing of BALE (human reads removed)"

exp_lib_strategy: ("RNA-Seq", "WGS", "AMPLICON", "Targeted-Capture") →
Variant in Sample libraryStrategy
exp_lib_source: ("METATRANSCRIPTOMIC", "METAGENOMIC", "GENOMIC",
"VIRAL RNA")
exp_lib_selection: ("RANDOM", "RT PCR", "RANDOM PCR", "unspecified", "PCR",
"cDNA")
<pre>exp_lib_layout: ("PAIRED" "SINGLE")</pre>
exp_platform: ("Illumina , "Nanopore") → Variant in Sample seqPlatform
exp_platform_model: ("Illumina MiSeq", "Illumina MiniSeq", "Illumina HiSeq
2500","NextSeq 500", "NextSeq 550", "Illumina iSeq 100", "GridION")
variant caller: (from VCF or Galaxy or Master of Pores pipelines) → Variant in
Sample variantCaller
-BIOSAMPLE
biosample_id: (external ref) e.g "SRS6007144"
biosample_alt_id: (external ref) e.g "SAMN13872787"
individual_id
collection_date: e.g "2020-02-14"
biosample_type: (sample type/source) e.g "Bronchoalveolar lavage fluid" or "Cellular
passage" → sampleOrigin
procedure: → obtentionProcedure
✓ culture_cell: e.g: "Vero E6 cells (CRL 1586)" (NULL or none if not culture)
✓ culture_passage_history e.g "Original (not passaged)" (NULL or none if not culture)
info
→ out
biosample_ref_material: e.g "BEI Resources catalog NR-52281 (lot 70033135)
- HOST/ INDIVIDUAL (ON THIS 1 ENCOUNTER)
individual_id: (external ref)
host_taxon_id: e.g "9606" ("Homo sapiens") → where?
host_age: e.g "21" → Individual age
host_sex: "female", "male" (sex in default schema) → Individual sex
geo_origin: e.g "USA:WI:Madison" → Individual geographicOrigin
☐ disease: (relevant virus-related diseases) e.g "COVID 19 pneumonia" → Individual
diseasesDisease

	disease_stage: e.g "acute" → Individual diseasesStage
	comorbidities: (underlying chronic diseases, format as individualDiseases from default
	schema): e.g ICD10 for "diabetes mellitus type II" → Individual diseasesDisease and
	diseasesStage where they are set stage: chronic
\checkmark	disease_course: categorical "asymptomatic", "mild", "severe"
\checkmark	disease_outcome: e.g "resolution/discharge" , "fatal"
	info
	→ out
	— VIRUS (Should there be one generic for organism/entity which sequence data belongs to
	to include their relevant phenotypic features - in the form of whatever phenotypic feature
	ontology for any species, or one specific to Genus , or being removed altogether?)
	taxon_id: e.g "433733" → Variant Basic taxonId
\checkmark	taxon_name: e.g "Severe acute respiratory syndrome coronavirus 2"
\checkmark	strain_id:
\checkmark	strain_name: e.g "2019 nCoV/USA WI1/2020" → info?
\checkmark	annotation
	genomic annotation: file url
	✓ functional annotation: file url