Use this form to tell us important information about this document, then start the text on the following page. All information you give in this form will appear in the document.

SPA YML Schom	a 1 3 Poloaco Notos I	Draft B			Created (2011-01-14)	Updated (2011-05-1
SRA XML Schema 1.3 Release Notes Draft B					Created (2011-01-14)	Opualed (2011-05-1
Author Information	on					
Given Name(s)	Last Name	Suffix	Degrees	Affiliation	1	Email

Use one row for each author. List authors in order of appearance in the document. Add rows to add more authors.

### SRA XML Schema 1.3 Release Notes

Draft B - 17 May 2011

Status	Proposed
Active Date	2011-06-01
Inactive Date	
Scope	INSDC SRA

#### 1 Overview

This document summarizes the proposed changes for Release 1.3 of the Sequence Read Archive (SRA) schemas governing XML metadata. This schema will be used by the SRA archive instances and has been developed under the auspices of the International Nuceotide Sequence Database Collaboration (INSDC).

Release 1.3 is a change over Release 1.2, which was in introduced in October 2010. While the schemas are incompatible, all data have been migrated so that documents submitted or modified before release remain valid. The goal of this release is to update choices, introduce new features, and specify a usable Analysis object usable for BAM file submissions. These changes are being introduced with the objective of not invalidating any current valid XML documents.

Major new features in this release are:

- Addition of new instrument models
- Require certain fields that have already been migrated
- Allow for modification of already-loaded analysis objects

#### 1.1 Notice

The features and modalities described in the XML schema DO NOT constitute a statement of features and mechanisms available in the SRA. The schema changes frequently must precede actual implementation. New feature rollouts and functionality changes are made asynchronously with XML schema changes.

#### 1.2 Related Documents

The SRA schema for this release can be obtained from this site: http://www.ncbi.nlm.nih.gov/viewvc/v1/trunk/sra/doc/SRA\_1-3

### 1.3 Revision History

Drafts A- ... created 14 Jan 2011 and updated through .... Approved for release by INSDC partners .... Scheduled for release ....

# 2 Explanation of Changes

## 2.1 Changes to All Documents

## 2.1.1 Adjustment to import statements

All document importing SRA.common.xsd now point to a resolvable URL: http://www.ncbi.nlm.nih.gov/viewvc/v1/trunk/sra/doc/SRA/SRA.common.xsd?view=co

# 2.2 Changes to SRA.Common.xsd

#### 2.2.1 Add new instrument models

New instrument values have been added to Platform block:

- "Illumina HiSeq 1000" [Illumina],
- "Illumina MiSeg" [Illumina],
- AB SOLiD 5500xl SOLiD System

- AB SOLiD 5500 SOLiD System
- "PacBio RS" [Pacific Biosciences]
- "Complete Genomics" [Complete Genomics] (platform already exists)
- ION\_TORRENT (new platform) and instrument models:
  - o Ion Torrent PGM

#### 2.2.1 Remove deprecated instrument models

• Solexa 1G Genome Analyzer (use Illumina choices)

### 2.2.2 Add GapDescriptor

A new structure called the GapDescriptor is introduced that will encode the placement of spot subsequences (tags) against a reference or assembly substrate. This structure encodes mate pair gaps and tandem read gaps. It is possible to express gaps distances in three ways: as mean/standard deviation, as min-max range, and as histogram. Orientation of the tag pairs can be described as "innie", "outie", "normal", and strand-opposite "anti-normal", following the nomenclature of the Celera Assembler.

Introduction of the GapDescriptor element was motivated by the need to describe CompleteGenomics platform sequencing. It is also intended that the GapDescriptor replace the LIBRARY\_LAYOUT element in the LibraryType. The GapDescriptor can be specified at the level of Run in order to override any general settings at the level of experiment.

### 2.2 Changes to SRA Experiment

## 2.2.1 Require LIBRARY\_STRATEGY

The design parameter LIBRARY\_STRATEGY is now required.

# 2.2.2 New Library Source choice METATRANSCRIPTOMIC

This was requested by EBI.

# 2.2.3 Removed deprecated library strategy choice BARCODE

This change was requested by EBI. No records have this designation.

### 2.3 Changes to Study

### 2.3.1 The STUDY\_TYPE block is now optional

In preparation for migration to BioProjects, this block has been made optional, and will be deprecated.

#### 2.3.1 The RELATED STUDIES/STUDY block removed

In preparation for migration to BioProjects, this deprecated block has been removed.

#### 2.4 Changes to Sample

#### 2.4.1 TAXON ID now required

The TAXON\_ID field in the SAMPLE\_NAME block is now required. All records already have this.

### 2.5 Changes to Submission

### 2.5.1 Submission handle removed

Submission handle attribute has been removed.

### 2.5.1 PROTECT action is now a complex type

This is a technical improvement requested by a major submitter.

### 2.6 Changes to Run

#### 2.6.1 Replicated descriptors at Run level

- Replicated SAMPLE\_DESCRIPTOR at the level of Run. If specified at Run, it will override the setting at the level of Experiment.
- Replicated GAP\_DESCRIPTOR at the level of Run. If specified at Run, it will override the setting at the level of Experiment.
- Replicated SAMPLE\_DESCRIPTOR at the level of Run. If specified at Run, it will override the setting at the level of Experiment.

## 2.6.2 Require checksum and checksum method

The DATABLOCK/FILES/FILE/@checksum and @checksum\_method will now be required attributes.

# 2.7 Changes to Analysis

## 2.7.1 DATA\_BLOCK not required for modification

The DATA\_BLOCK is now required for add submissions, but no longer for modify submissions.

## 3 Deprecated Fields

SRA 1.3 contains the following fields, branches, and options that should no longer be used in current submissions.

		T	
SRA.common.xsd	SPOT_DECODE_METHOD		
SRA.common.xsd	NUMBER_OF_READS_PER_SPOT		
SRA.common.xsd '454 Titanium'		use '454 GS FLX Titanium'	
SRA.common.xsd	'GS 20'	use '454 GS 20'	
SRA.common.xsd	'GS FLX'	use 'GS FLX'	
SRA.common.xsd	'Solexa 1G Genome Analyzer'	use 'Illumina Genome Analyzer'	
SRA.common.xsd	CYCLE_SEQUENCE	use SEQUENCE_LENGTH	
SRA.common.xsd	CYCLE_COUNT	use SEQUENCE_LENGTH	
SRA.study.xsd	CENTER_NAME	use STUDY@center_name	
SRA.study.xsd	PROJECT_ID	use RELATED_STUDIES instead	
SRA.study.xsd	RELATED_STUDIES/STUDY	use RELATED_STUDIES/RELATED_STUDY	
		instead	
SRA.experiment.xsd	LIBRARY_STRATGEY/BARCODE	use another library strategy	
SRA.experiment.xsd	LIBRARY_SOURCE/NON	use METAGENOMIC or TRANSCRIPTOMIC	
	GENOMIC	instead	
SRA.experiment.xsd	PROCESSING/BASE_CALLS	use PIPELINE instead	
SRA.experiment.xsd	PROCESSING/QUALITY_SCORES	use PIPELINE instead	
SRA.experimentxsd	@expected_number_spots		
SRA.experimentxsd	@expected_number_reads		
SRA.run.xsd	'_seq.txt, _prb.txt, _sig2.txt, _qhg.txt'	use 'Illumina_native' instead	
SRA.run.xsd	@total_spots		
SRA.run.xsd	@total_reads		
SRA.run.xsd	@number_channels		
SRA.run.xsd	@format_code		
SRA.run.xsd	@instrument_model	use PLATFORM/INSTRUMENT_MODEL instead	
SRA.run.xsd	@run_file		
SRA.run.xsd	@total_data_blocks		
SRA.submission.xsd	HoldForPeriod		
SRA.submission.xsd	@submission_id	use alias instead	

SRA.submission.xsd	@handle	

## **4 Future Planned Revisions**

The next revision is anticipated to be contracting revision (one that potentially invalidates current documents). The main changes will be to remove deprecated fields. This will involve migration of data in anticipation of future schema changes.

## Figures, Tables and Boxes Appendix (do not delete)

Place numbered figures, tables and boxes (referred to from the main text) below.

"In-line" figures (e.g. equations) and tables should be placed within the main text in their desired final location.

Percentage have a single level of sections: the titles for those sections should be marked up in "Percentage".

Boxes can have a single level of sections; the titles for these sections should be marked up in "Box subhead" style.