

## 1. General Information

<b>Cohort ID</b>	5020_21
<b>Title (Study Name)</b>	The CoVVacBoost Study
<b>Principal investigator</b>	Assoc. Prof. Martin Stradner, Prof. Ivo Steinmetz
<b>Contact information</b>	<a href="mailto:pm-biobank@medunigraz.at">pm-biobank@medunigraz.at</a>
<b>Funding agency</b>	----

## 2. Description

The aim of this study is to evaluate the ability of a COVID-19 booster vaccine to induce a salivary antibody response and to investigate a possible correlation with the serum antibody response and the cellular response. A quantitative and functional comparison of the salivary anti-SARS-CoV-2 antibody response before and after 3<sup>rd</sup> booster vaccination will be investigated. Volunteers, who get their 3<sup>rd</sup> vaccination at the "Impfstraße" at the LKH-Univ. Klinikum Graz are asked to participate in this study. Blood and saliva samples will be obtained during 3 visits.

Visit 1 (before the 3<sup>rd</sup> booster vaccination): Serology, PBMCs, saliva

Visit 2 (3 – 8 weeks after the 3<sup>rd</sup> booster vaccination): Serology, PBMCs, saliva

Visit 3: 6 months after the 3<sup>rd</sup> booster vaccination: Serology, saliva

The following data are collected for all participants: type of COVID-19 vaccinations, medical history (including immunosuppressive therapies, chronic graft versus host disease, severe opportunistic infections, prior vaccination history (types, titers if available), current and past disease including COVID-19), current medication, BMI, smoking habits.

## 3. Details

<b>ICD 10/O codes / Healthy</b>	healthy	
<b>Key words</b>	vaccination, COVID-19, SARS-CoV-2, mRNA vaccine, booster, 3 <sup>rd</sup> dose, Moderna, Comirnaty, AstraZeneca	
<b>Collection / Cohort size</b> 12/2023	1.571 serum, 370 saliva and 553 PBMC aliquots from 140 patients	
<b>Informed Consent (IC)</b>	<input checked="" type="checkbox"/> Broad Biobank IC	
	<input checked="" type="checkbox"/> Specific Study IC	
<b>Status</b>	<input checked="" type="checkbox"/> In progress / compl. date: 01/2023	
	<input type="checkbox"/> Completed	
<b>Inclusion criteria</b>	<b>Age distribution</b>	18+
	<b>Sex distribution (f:m)</b>	~50:50
	<b>Others</b>	<ul style="list-style-type: none"> <li>• second dose AstraZeneca plus third dosis Moderna</li> <li>• second dose AstraZeneca plus third dosis Comirnaty</li> <li>• second dose Moderna plus third dosis Moderna</li> <li>• second dose Comirnaty plus third dosis Comirnaty</li> </ul>
<b>Earliest access</b>	As of now	

	<b>O-FIS Qualitätsmanagementsystem</b> Formblatt <b>Collection and Cohort Profile</b>	CL312 Seite 2 von 2
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<b>Quality-standards</b>	<input checked="" type="checkbox"/> ISO 9001:2015 (SOPs)
<b>Associated publications / references</b>	-----

#### 4. Material available (aliquot size) and storage conditions

<b>Material</b>	<input checked="" type="checkbox"/> Serum (580 µl)	<input checked="" type="checkbox"/> -80°C	<input type="checkbox"/> liq. N <sub>2</sub>
	<input checked="" type="checkbox"/> PBMC´s	<input checked="" type="checkbox"/> -80°C	<input checked="" type="checkbox"/> liq. N <sub>2</sub>
	<input checked="" type="checkbox"/> Saliva	<input checked="" type="checkbox"/> -80°C	<input type="checkbox"/> liq. N <sub>2</sub>

<b>Dokument erstellt</b> (tt/mm/yyyy): 22/12/2021	<b>Letzte inhaltliche Aktualisierung</b> (tt/mm/yyyy): 21/03/2024
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