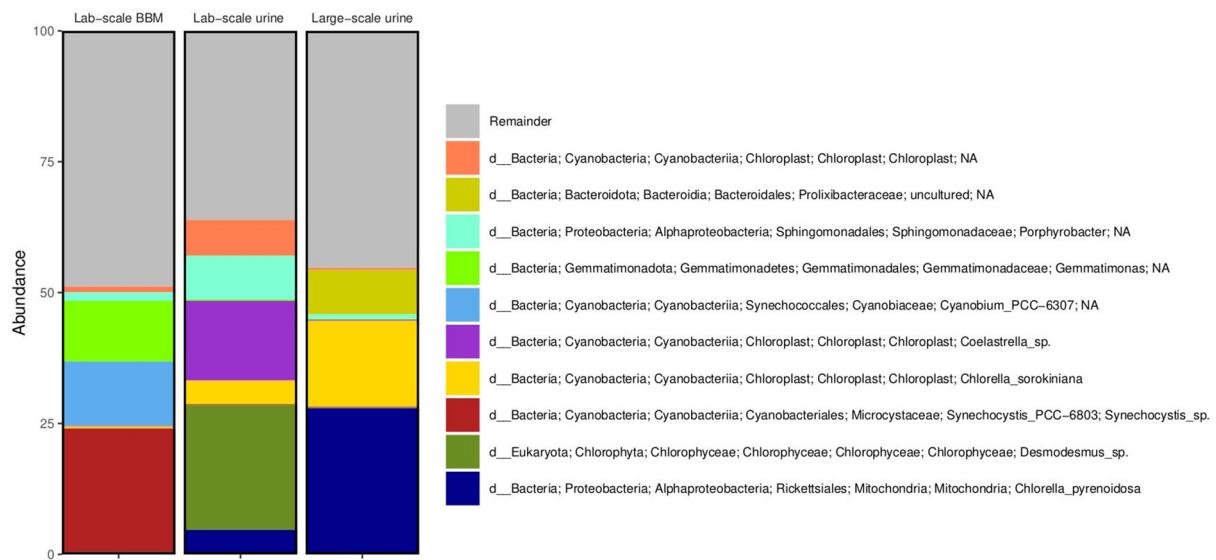


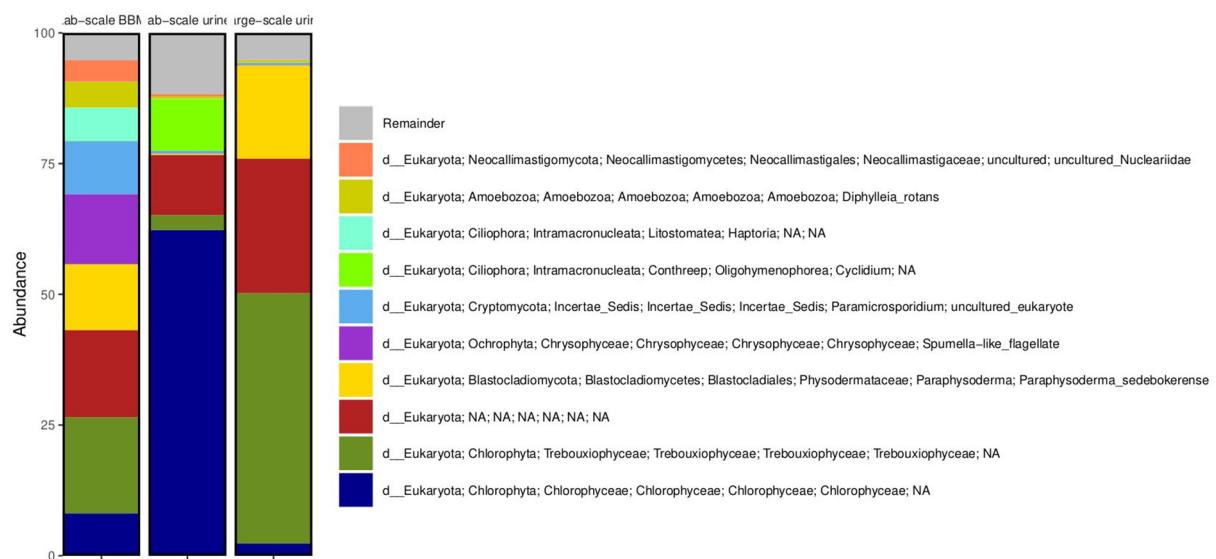
**Tech.phi2 Table S1.** Sequence information from QIIME 2 (2022.8 version) processing of NGS amplicon reads

	Samples	Number of reads	Number of sequences after denoising with Dada2	Number of observed ASVs
16S rRNA	1	53725	49269	415
	2	67913	61667	357
	10	63615	59403	394
18S rRNA	1	97940	95098	69
	2	96266	93876	74
	10	68905	67319	64
23S rRNA	1	113771	110267	82
	2	112697	109131	73
	10	96778	94580	51
tufA	1	31673	27026	97
	2	35074	29654	51
	10	11132	9465	64

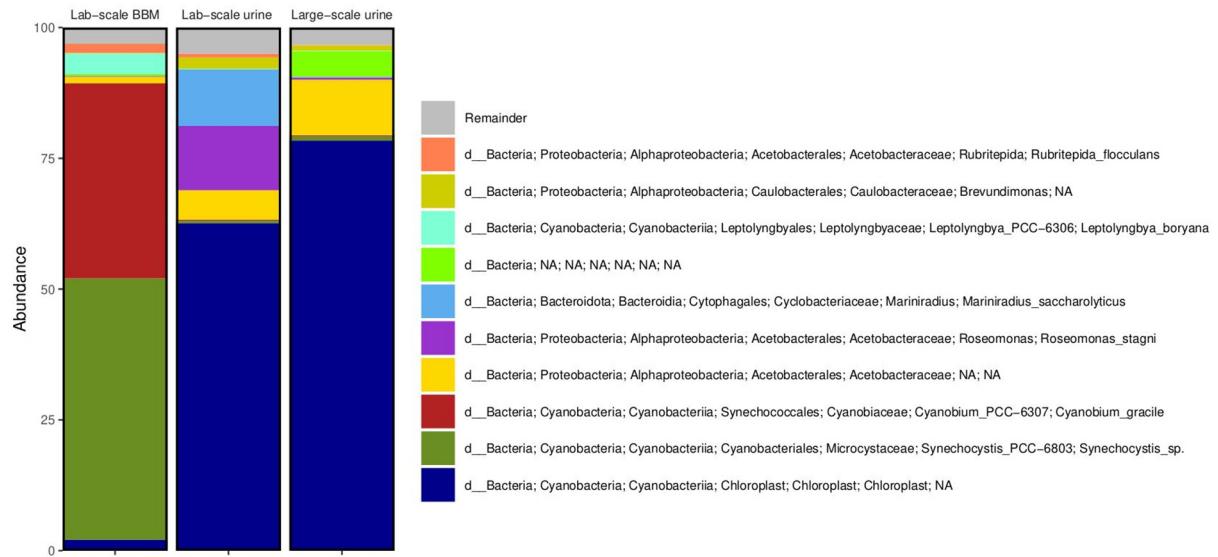
(a)



(b)



(c)

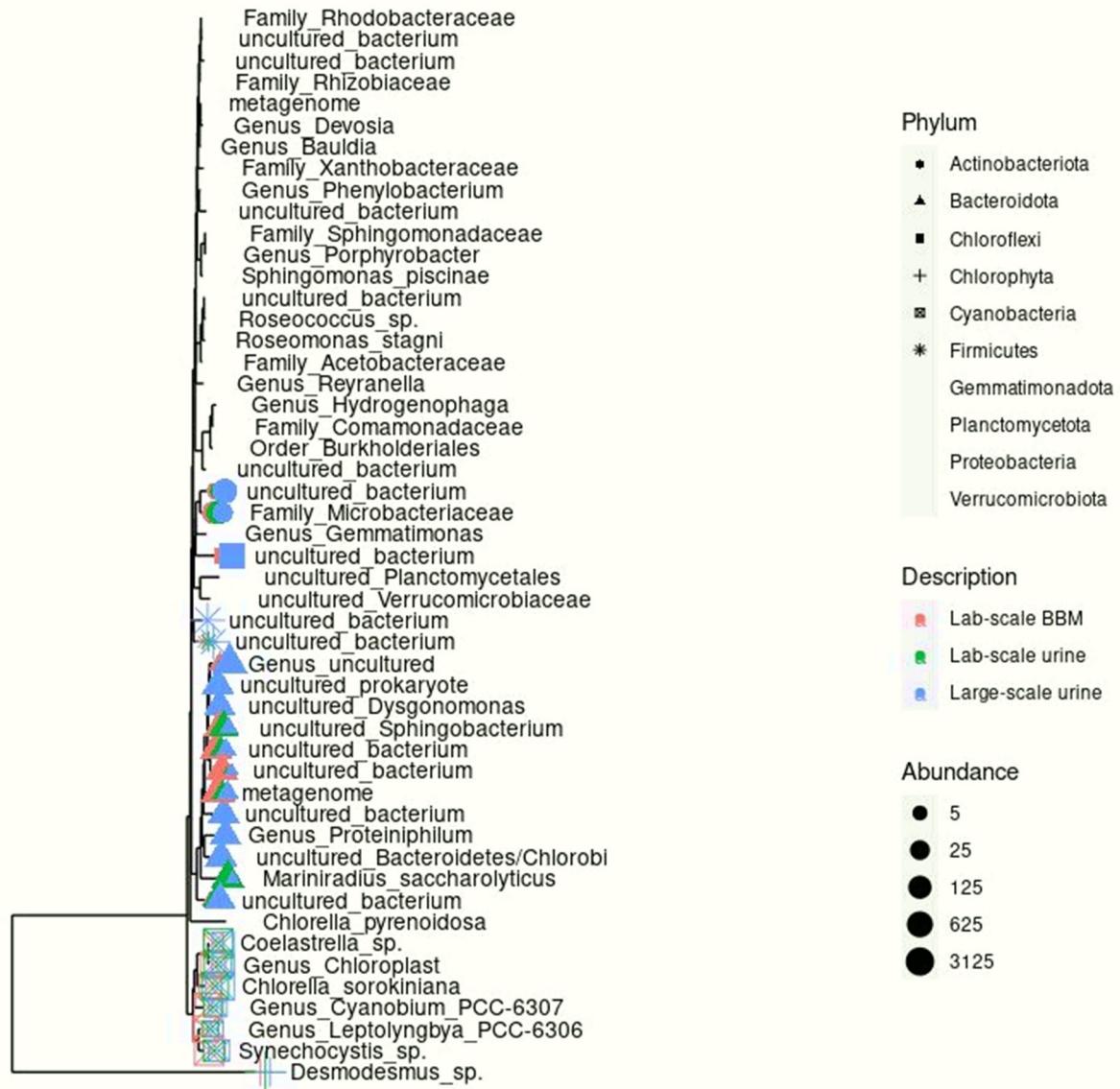


(d)

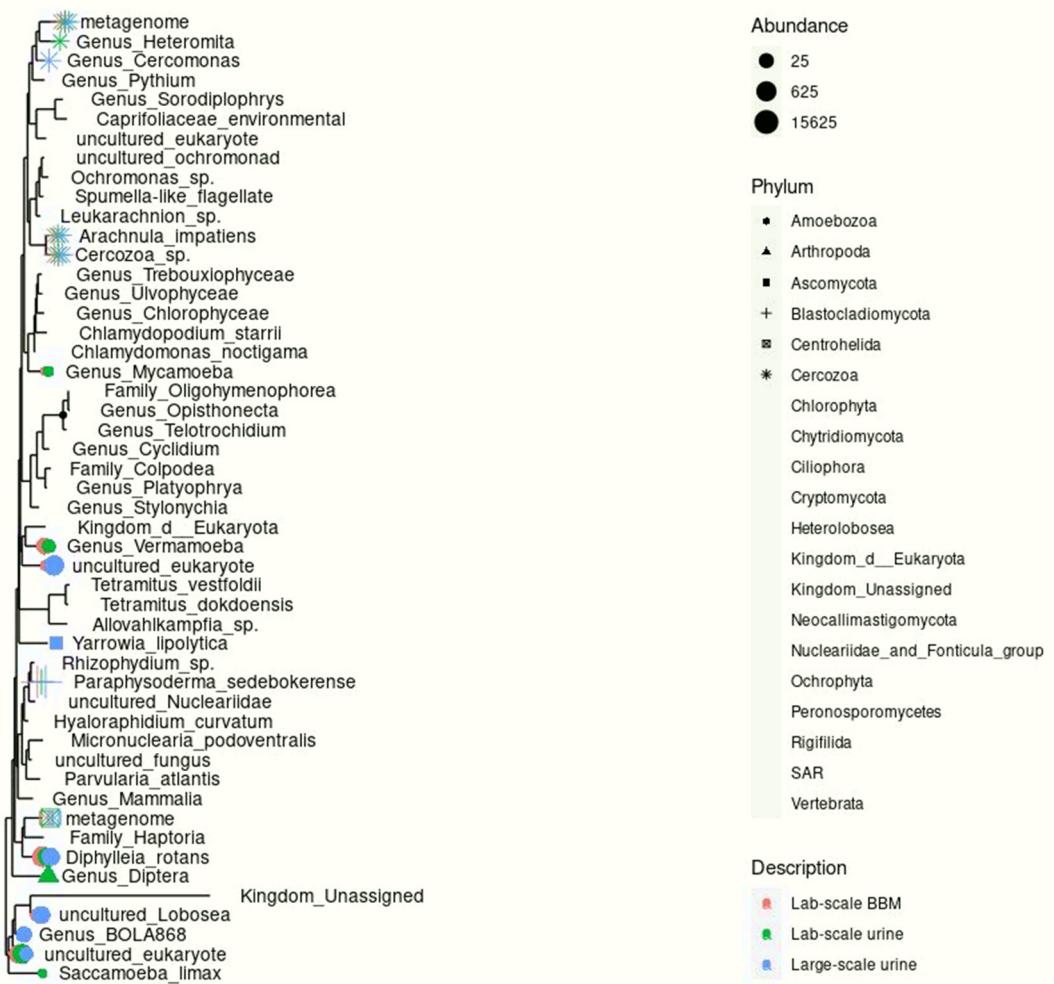


**Figure S1.** Bar plots showing variation in the relative abundances of taxonomies up to species level in lab-scale (urine and BBM) and large-scale urine microbial communities. Colors represent microbial taxonomy classified by Silva taxonomy (release\_138) with using (a) 16S rDNA marker regions, (b) 18S rDNA marker regions and (c) 23S rDNA marker regions, by (d) tufA database [Sauvage et al., 2016] [50].

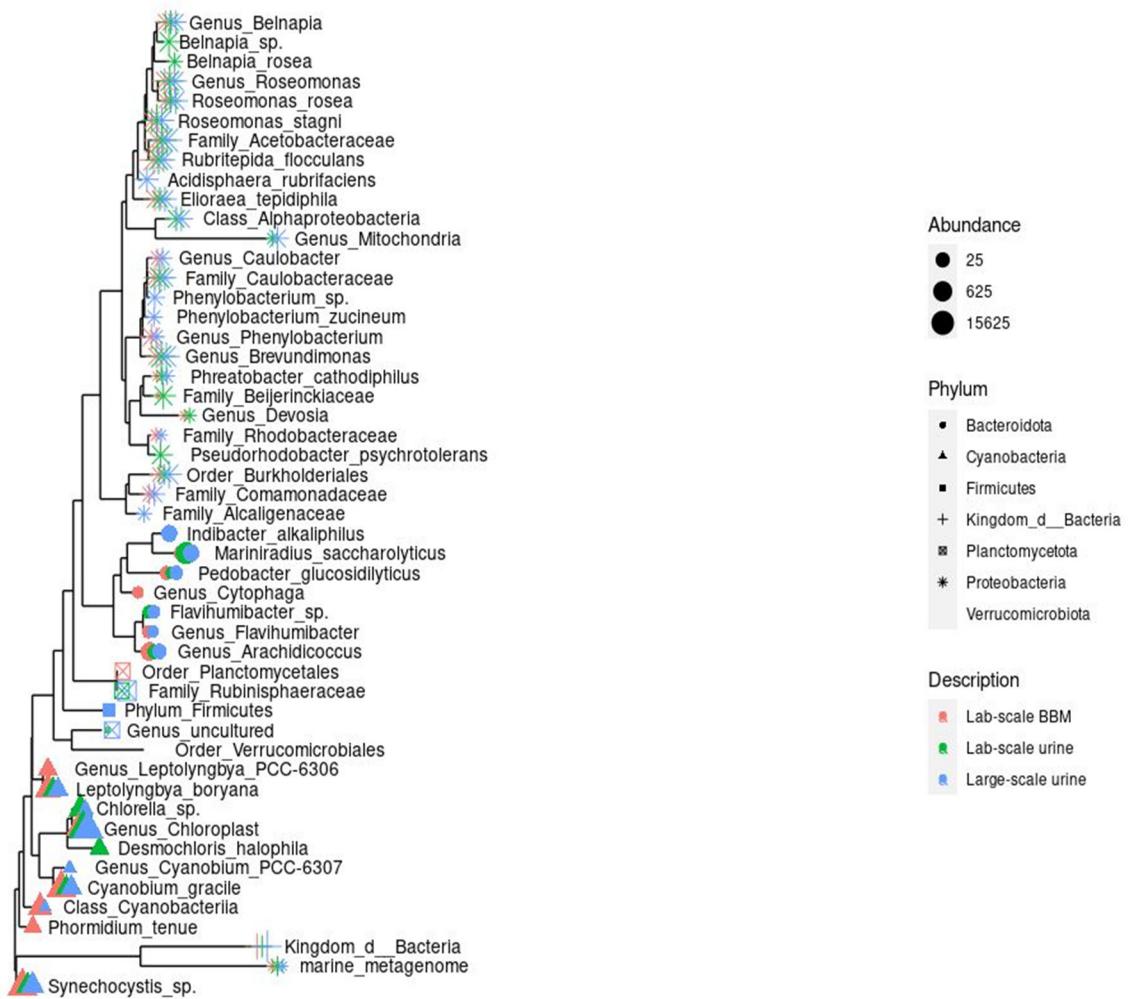
(a)



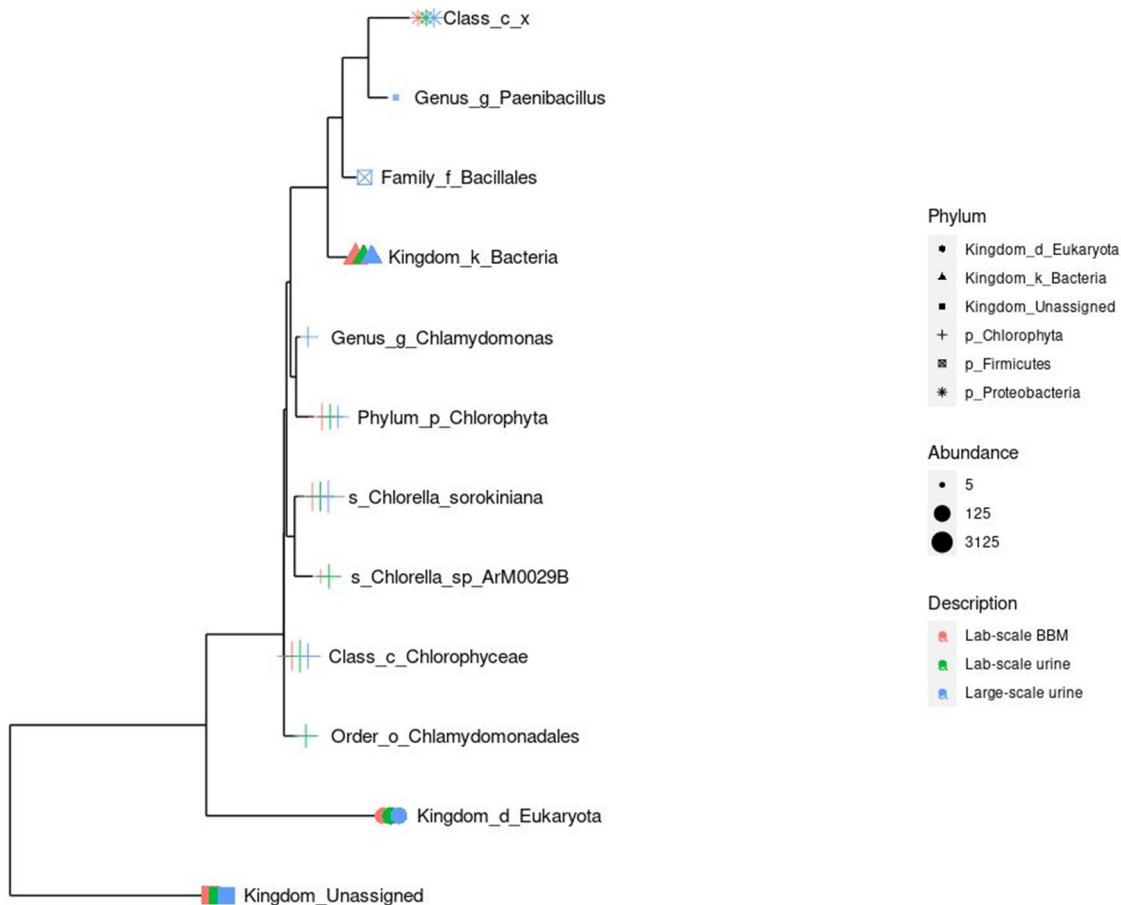
(b)



(c)



(d)



**Figure S2.** Phylogenetic trees showing the relationship of (a) 16S rRNA, (b) 18S RNA, (c) 23S rRNA and (d) tufA gene sequences. All the phylogenetic trees were constructed for top 50 taxa for better visualization.

50 Sauvage, T.; Schmidt, W.E.; Suda, S.; Fredericq, S. A metabarcoding framework for facilitated survey of endolithic phototrophs with tufa. *BMC Ecol.* **2016**, *16*, 1–21.