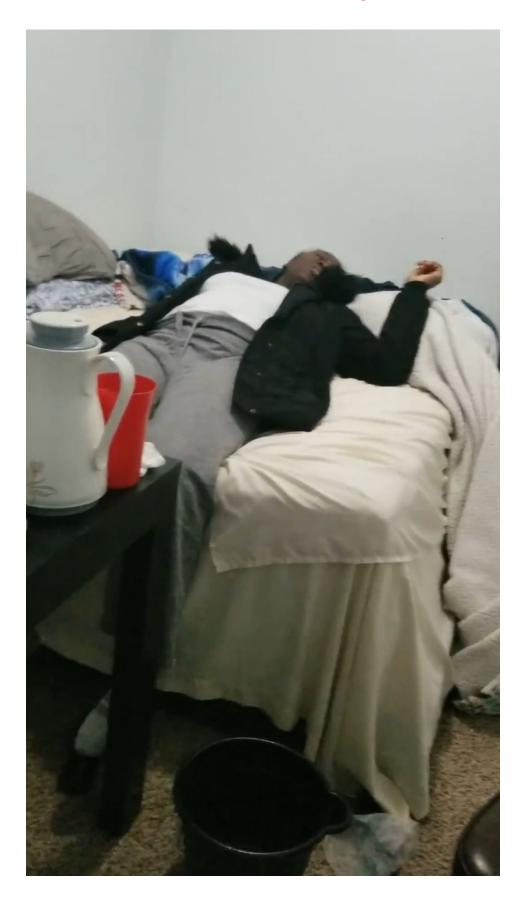


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Vaccine	Product Name/Manufacturer Lot Number	Date	Healthcare Professional or Clinic Site	
1st Dose COVID-19	COVID-19 Vaccine Wifg: Pfizer BioNTech	1 13 21 mm dd yy	M Health Fairview Southdale	-
2 <sup>nd</sup> Dose COVID-19	COVID-19 Vaccine	2,1,21 mm dd yy	M Health Fairview Southdale	
Other	Mfg: Pfizer BioNTech Lot: EL9262 Exp: 5/31/21	mm dd yy		
Other		mm dd yy		
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Vaccine	Date / Fecha  02 01 21  mm dd yy	
COVID-19 vaccine Vacuna contra el COVID-19 0800- Oc		
Other Otra	mm dd yy	
ring this vaccination record to every accination or medical visit. Check with your ealth care provider to make sure you are not dissing any doses of routinely recommended accines.  For more information about COVID-19 and COVID-19 vaccine, visit cdc.gov/pronavirus/2019-ncov/index.html.  For an report possible adverse reactions allowing COVID-19 vaccination to the accine Adverse Event Reporting System AERS) at vaers.hhs.gov.	Lleve este registro de vacunación a cada cita médica o de vacunación. Consulte con su proveedor de atención médica para asegurarse de que no le falte ninguna dosis de las vacunas recomendadas.  Para obtener más información sobre el COVID-19 y la vacuna contra el COVID-19, visite espanol.cdc.gov/coronavirus/2019-ncov/index.html.  Puede notificar las posibles reacciones adversas después de la vacunación contra el COVID-19 al Sistema de Notificación de Reacciones Adversas a las Vacunas (VAERS) en vaers.hhs.gov.	



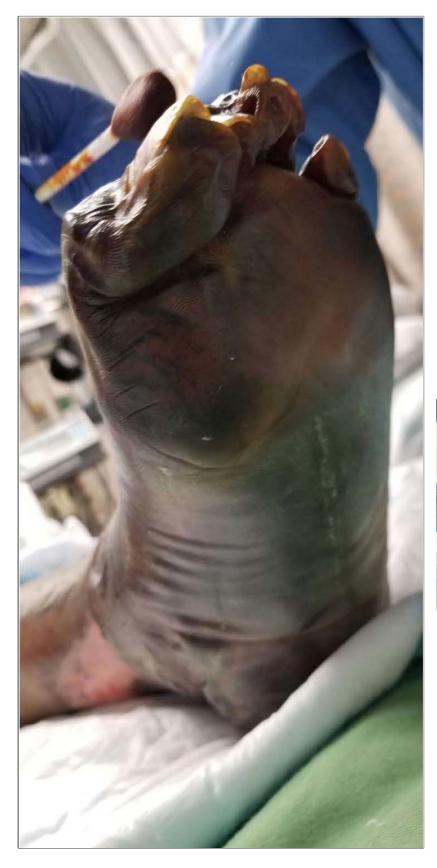






















Subject: Mrs. Jummai Nache and her Family













Subject: Mrs. Jummai Nache and her Family Conclusion



Mrs. Jummai Nache and her devoted husband Philip; learning how to walk stairs on prosthetic limbs. Amputation of her limbs **caused by mRNA injection** <u>mandated</u> **by her former employer**, the University of Minnesota (UMinn). A non-smoker, a non-drinker, a model of health **prior** to Pfizer needle; amputations in pictorial review above caused by mRNA-induced venous and arterial thromboembolism; not merely foreseeable but a known mRNA danger which had been discussed in-detail at the secret FDA meeting of 22 October 2020 (See cover letter Page 8-of-13).